# Metabarcoding of insect-associated fungal communities: a comparison of internal transcribed spacer (ITS) and large-subunit (LSU) rRNA markers

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## Abstract

## Complex communities of fungi are regularly characterised by deep sequencing with standard barcode markers, especially the Internal Transcribed Spacer (ITS) and the Large-Subunit (LSU) gene of the rRNA locus. Full taxonomic characterisation of fungal communities is necessary for establishing ecological associations or for early detection of pathogens and invasive species, but reliance on a single short sequence fragment may be problematic due to various technological and analytical obstacles. Here we conducted a side-by-side comparison of fungal communities associated with bark beetles (Scolytinae), the likely vectors of several tree pathogens, using ITS and LSU. Both markers revealed similar patterns of overall species richness and response to key variables (beetle species, forest type). Identification against the respective reference databases using various classifiers revealed similar higher-level taxonomic composition, but decreasing resolution towards lower levels, especially at the species level. Thus, Operational Taxonomic Units (OTUs) could not be linked via taxonomic classifiers across ITS and LSU fragments. However, a phylogenetic analysis (focused on the epidemiologically important Sordariomycetes) placed OTUs relative to reference sequences spanning both loci and demonstrated the largely similar phylogenetic distribution of ITS and LSU-derived OTUs. The analysis of congruence in both markers also suggested the biologically most defensible threshold values for OTU delimitation (98% for ITS, 99% for LSU). Studies of complex fungal communities using the canonical ITS metabarcode marker require corroboration across additional loci, and benefit from phylogenetic analyses and their greater precision compared to conventional taxonomic classifiers, even in the face of incomplete and partially identified reference databases.

## Introduction

Fungal communities associated with insects have been widely studied to disentangle the nature of their interactions (Ganter 2006; Li et al. 2016; Malacrinò et al. 2017; Raman et al. 2012; Six 2020). For these studies to succeed, an accurate and reliable fungal identification is essential. However, these identifications are challenging due to the cryptic nature and incomplete taxonomy of the fungi, as only 3-8% of fungal species have been described so far (Hibbett et al. 2016; Wedaralalage et al. 2020). Conventional studies of fungal communities have been conducted by isolating and culturing each fungus recovered from the substrate (Batra 1963), which generated biases towards culturable species. High-throughput sequencing has provided an alternative methodology, especially through metabarcoding, i.e. the sequencing of short amplicons from mixed communities (Yu et al. 2012). Metabarcoding is now widely applied in characterising species composition and biodiversity of fungal communities associated with insects. In the specific case of fungal communities associated with bark beetles, metabarcoding usually detects dozens of species of fungi on a single insect specimen (Bálint et al. 2014; Hulcr et al. 2020; Johnson et al. 2018; Malacrinò et al. 2017; Miller et al. 2016; Miller et al. 2019).

There is broad agreement that the internal transcribed spacer (ITS) of the nuclear rRNA gene cluster should be the standard DNA barcode in fungi (Schoch et al. 2012). Its use in metabarcoding is now equally well established, and extensive reference databases and universal primer combinations are in wide use (Porras-Alfaro et al., 2014; Tedersoo et al., 2015). However, various challenges remain for accurate characterisation of communities. PCR amplification biases may skew species recovery (Bellemain et al. 2010; Dreaden et al. 2014; Harrington et al. 2011; Li et al. 2020), in particular given that primer design is constrained in studies of associations with insects, which requires to avoid co-amplification of the corresponding genomic regions of the hosts. In addition, the recovered short sequence fragments have limited power for placement in the phylogenetic system (Porras-Alfaro et al. 2014; Vrålstad 2011), exacerbated by the taxonomic insufficiencies of the reference databases (Agerbo Rasmussen et al. 2020; Miller et al. 2016; Porras-Alfaro et al. 2014; Tedersoo et al. 2015b). In response to these concerns many fungal phylogenetic and barcoding studies have used a combination of ITS and partial large and small subunit (LSU and SSU) rRNA genes, as well as other markers such as RPB2 and TEF1α (Lutzoni et al. 2004; Stielow et al. 2015; Zhang et al. 2006). The latter two markers lack representation in databases, while curated reference databases and analysis tools like SILVA and RDP (Ribosomal Database Project) are built specifically for SSU and LSU genes (Wang et al. 2007; Quast et al. 2012).

In practise, both ITS and the LSU/SSU markers exhibit particularities whose benefits and drawbacks depend on the aim and scope of a study (Porras-Alfaro et al., 2014). While the LSU/SSU genes are less variable than the ITS intergenic region, which favours alignment and tree-based analyses, their low molecular rate reduces the taxonomic resolution at the species-level. In turn, ITS provides better species resolution, but the higher mutation rate leads to intragenomic variation of the tandem repeat units, and high variability often translates into poor alignments, making it difficult to apply phylogenetic methods (Porras-Alfaro et al. 2014; Vrålstad 2011). However, this intraspecific and intragenomic variation is still poorly documented, and may affect not only the ITS (Lücking et al. 2020). Furthermore, the suitability of the primers used to amplify fungal communities is often tested by using mock communities (Bakker 2018; Egan et al. 2018; Frau et al. 2019), but only a few studies have addressed the relative power of ITS and LSU/SSU markers in complex natural fungal communities (Jusino et al. 2019; Li et al. 2020; Parada et al. 2016).

A limited number of studies comparing different regions of the nuclear rRNA cluster have revealed similar ecological patterns and broad conclusions that are not profoundly affected by marker choice (George et al. 2019; Nilsson et al. 2019a; Tedersoo et al. 2015a). However, these studies generally have applied a coarse-grain approach of higher-level taxonomic analysis, rather than the species level. As the databases become more complete and the quality of HTS reads is improving, increasingly precise matches of database and query allow a greater detail of community analysis, ultimately at the species (Operational Taxonomic Unit, OTU) or haplotype (Amplified Sequence Variant, ASV) level. In the few cases where this approach has been taken to date, the choice of markers seems to be critical. For example, the ITS marker may not detect key pathogen species that are recovered using LSU (Hulcr et al. 2020; Skelton et al. 2019). Furthermore, ideas of what constitutes a species and what are the most appropriate ways to circumscribe them based on barcode sequences are always shifting (Hibbett et al. 2016; Hyde et al. 2013; Wedaralalage et al. 2020; Kõljalg et al. 2013; Lücking et al. 2020).

This leaves the question about which marker is preferrable, and given the presumed advantages of each, to what degree does the marker choice matter for the conclusions from metabarcoding studies? Here we address these questions for the case of fungal communities associated with bark beetles (Coleoptera: Scolytinae). These insects breed in living or dead trees and form close associations with fungi important for access to nutrients from wood that cannot be utilised directly by the beetles themselves (Batra, 1963). Fungal communities associated with these beetles are highly diverse and form symbioses of various strength and specificity, which in many cases involve the active transport of fungal associates in specifically adapted pockets of the beetles’ exoskeleton, the mycangia (Six, 2020). A beetle-fungus complex can cause enormous damage to forest ecosystems, e.g. resulting in the demise of chestnuts in North America, elms across the Northern Hemisphere, or the recent large-scale decline of conifer forests in Central Europe and North America, which usually involve fungi from the ascomycote orders Ophiostomatales, Microascales and Hypocreales (Class Sordariomycetes) (Ploetz et al. 2013). Metabarcoding provides a tool for detailed studies of these complex communities, but the results may be influenced by the choice of barcode markers and other problems of using short sequences from PCR mixtures to characterise complex species communities. Confidence in the metabarcoding approach can be improved by analysing ITS and LSU marker sets from a given sample in parallel, which should produce the same outcome. However, given different evolutionary rates, the limited power of phylogenetic placement, and different resolution of species differentiation may result in mismatched conclusions from either marker. We used individuals from four bark beetle species obtained from three forest types to characterise the associated fungal communities, conducting a comparison of the two markers with regard to: (1) broad ecological trends of fungal associations taking a whole-community approach, and (2) species identifications against existing ITS and LSU fungal reference databases, using various taxonomic classifiers and explicit phylogenetic methods. The side-by-side comparison addresses the power of either markers to infer critical parameters of fungal community metabarcoding, such as the number and taxonomic identity of OTUs, their ecological associations, and the inferences on whole-community diversity and turnover.

## Materials and Methods

#### Samples used and laboratory procedures

Sequence data were generated from 20 specimens each of four beetle species: *Xylosandrus germanus*, *Xyleborinus saxesenii*, *Gnathotrichus materiarius*, and *Tomicus piniperda* (**Table 1**). Only the latter is a xylophagous ‘bark beetle’ in the strict sense, while the three others are considered mycelia feeding (xylomycetophagous) ‘ambrosia beetles’ that apparently rely on active transport of fungi indicated by the presence mycangia (see Six, 2020b). Specimens were collected by Forest Research during 2013-2015 in the New Forest National Park (50º50'52.08" N 1º35'33.51" W), Hampshire, UK, using Lindgren multiple-funnel traps (Lindgren 1983) (Phero Tech). These traps were located in oak, spruce and pine forests and were baited with lures (100% ethanol, plus α-pinene) (Inward 2019). Propylene glycol (65%) was used as the preservation fluid at the bottom of the traps. Specimens were morphologically identified and selected at random to obtain the same number of specimens per beetle species and forest type.

In the laboratory, the specimens were rinsed with pure water to remove loosely adhering fungal tissue, and thoroughly macerated individually to ensure that all fungi associated with the specimens were released. DNA was extracted using the Qiagen DNeasy Blood and Tissue spin column extraction kit (Qiagen, Valencia, CA, USA). Individual DNA extracts were first tested for correct species identification using the COI barcode marker, which was amplified for a 418 bp fragment and sequenced on Illumina HiSeq following methods of (Arribas et al. 2016). In all cases the most abundant read, as determined with the NAPselect script (Creedy et al. 2019), had an exact match to existing reference sequences of the respective species, confirming the morphological identification.

The DNA extracts were then used for fungal metabarcoding of ITS with primers ITS86F (5′-GTGAATCATCGAATCTTTGAA-3′) / ITS4 (5′-TCCTCCGCTTATTGATATGC-3′)(Op De Beeck et al. 2014)and LSU using primers LR0R (5′-ACCCGCTGAACTTAAGC-3′) (Vilgalys and Hester 1990) / JH-LSU-369rc (5′-CTTCCCTTTCAACAATTTCAC-3′) (Li et al. 2016) targeting the D1-D2 region at the 5’ end of the LSU gene immediately downstream of the ITS2 region. Both markers were amplified from each beetle DNA extraction in separate reactions. Unique six-nucleotide indices added to each primer pair were used to distinguish the libraries. PCRs were pooled from three replicates conducted under slightly different annealing temperatures (54°C, 55°C and 56°C) to accommodate differences in optimal amplification conditions of the fungal species (Schmidt et al. 2013) and blank PCR reactions were used as negative control. Successful PCR amplicons were purified using the AMPure XP magnetic beads (Beckman Coulter) and amplicons were indexed using a secondary PCR with Nextera XT DNA Library Preparation Kit (Illumina Inc.) and sequenced on an Illumina® HiSeq® 2500 platform to generate 2 x 300 bp paired-end reads.

#### Bioinformatics

Raw reads were demultiplexed, primer trimmed and singleton reads removed with Cutadapt (Martin 2011). Read quality was evaluated using FASTQC (Andrews 2014). The raw reads generated for these analyses are available in the BioSample Submission Portal as Bio-Project PRJNA727174 and Sequence Read Archive (SRA) accession numbers PRJNA727174 (barcoding) and (metabarcoding).

Forward and reverse reads were merged and quality filtered (Phred score ≥30) using PEAR v. 0.9.8 (Zhang et al. 2014). After merging, the average read length was 252 bp for ITS and 357 bp for LSU. Subsequent steps were carried out using VSEARCH v. [2.15.0](https://github.com/torognes/vsearch/releases/tag/v2.15.0) (Rognes et al., 2016). A further quality test was effectuated using, with the --fastx\_filter command and -fastq\_maxee 1.0. After dereplication (--derep\_fulllength), assemblies were denoised (--cluster\_unoise --minsize 4 --unoise\_alpha 2) and length filtered for a range of 100 to 500 bp (--fastx\_filter) and all singletons removed. Chimera filtering was performed (--uchime3\_denovo) and reads were then clustered into Operational Taxonomic Units (OTUs) at various similarity thresholds (97%, 98%, 99%) using VSEARCH. The average length of the OTUs was 270 bp for ITS and 347 bp for LSU (Supplementary Figure S1). Reads were then mapped to the OTU clusters, outputting an OTU table of read abundances suitable for the ecological analysis of the samples.

#### OTU identification and classification

Fungal OTUs were classified following three widely used methods for species identification. The Ribosomal Database Project (RDP) Bayesian Classifier (Wang et al. 2007) was used for fungal identification employing the Warcup fungal ITS (v2, release March 2018) and UNITE (version February 2020) training sets (Deshpande et al. 2016; Edgar 2018). In addition, OTUs were processed through the PROTAXFUNGI pipeline (Abarenkov et al. 2018), implemented in the PlutoF (Abarenkov et al. 2010) platform and based on the UNITE fungal database (February 2020). PROTAXFUNGI hierarchically assigns the OTU identities from the root node of the taxonomy through to the species level (Nilsson et al. 2019b). It has not been implemented for LSU, and thus was applied to the ITS data only. A third classifier, IDTAXA, employs machine learning to reduce over-classification errors to obtain a higher accuracy (Murali et al. 2018). Taxonomic assignment was carried out separately on class, order, genus, and species level. The percentage accuracy of identification from these methods was averaged as a measure of confidence of each identification (average of three values for ITS and two for LSU data). A minimum threshold of 70% probability for at least one of the classifiers was set, below which the OTUs were considered as “unclassified”, together with other sequences that were identified with high confidence against database entries labelled as “unclassified”, “unidentified” or “*incertae sedis*”. When identifications disagreed among the classifiers, the one with the highest probability was selected. Taxonomic composition of samples was presented as the number of OTUs assigned to a given taxonomic level in a barplot created with *ggplot2* in Rstudio (team 2015).

In a more detailed study of OTU assignments in the ecologically important class Sordariomycetes, OTUs generated from clustering at 97%, 98%, and 99% similarity thresholds were classified using the Warcup training set and the RDP fungal data set for ITS2 and LSU, respectively (Deshpande et al. 2016). All OTUs with a confidence of assignment >80% to class Sordariomycetes were retained. Order-level assignments (the Sordariomycetes are split into 28 orders) with a confidence >50% were taxonomized, while all others were kept as “unclassified Sordariomycetes”. To assess the effects of differing clustering thresholds on downstream taxonomic assignment, OTUs at each clustering threshold were also closed-reference clustered (i.e. only retaining sequences with hits in the reference set) against the composite LSU/ITS2 reference sequences used to construct the tree (Edgar 2010; Rognes et al. 2016).

#### Alignment and tree building in Sordariomycetes

Reference sequences for the class Sordariomycetes were downloaded from GenBank, consisting of species that were complete for at least 2/3 of the rRNA cluster composed of SSU, LSU, and ITS2 sequence (full list of accessions in Suppl. Table 1). 80% of species in this reference set had accessions covering all three regions. ITS2 reference sequences were processed through ITSx to eliminate redundancy in the concatenated alignment (Bengtsson-Palme et al. 2013).

The reference sequences and OTUs were aligned using MUSCLE (Edgar 2004) under default settings and the aligned matrices were concatenated. The concatenated three-gene alignment was then inspected in Mesquite and Geneious Prime (v 2020.0.4) and problematic accession sequences were removed. The alignment was partitioned for each marker region, and the best model for each partition was selected according to BIC values. Model testing, tree building, and ultrafast bootstrap approximation (n = 1000) were performed in IQ-Tree2 (Chernomor et al. 2016; Minh et al. 2020).

#### Phylogenetic diversity metrics

Phylogenetic distribution of ITS and LSU copies was assessed by metrics of clustering and overdispersion originally developed for community ecology (Webb et al. 2008). In the ideal case of capturing the same species with both markers, copies of ITS2 and LSU corresponding to the same species should be in close vicinity on the tree, i.e. the copies of each marker should be ‘overdispersed’ (more dispersed than a random phylogenetic structure). Deviations from this pattern can be assessed with the metrics calculating the Mean Pairwise Distances (MPD) and Mean Nearest Taxon Distances (MNTD) of each set (ITS2 and LSU). We report standardized values as the net relatedness index (NRI) and nearest taxon index (NTI) relative to null models of randomly distributed communities. Positive NRI and NTI scores indicate phylogenetic clustering, negative values indicate phylogenetic overdispersions, while random phylogenetic structure results in values not significantly different from zero (Webb et al. 2008). Calculations were performed with the *R* packages *picante*, *ape*, and *phylomeasures* (Webb et al. 2008; Paradis and Schliep 2019; Tsirogiannis and Sandel 2016).

#### Assessment of species richness and community composition

Community ecological analyses were carried out on samples rarefied to 1000 reads, which was sufficient for generating largely complete OTU sets as judged by species accumulation curves (Supplementary Figure 2), thus retaining 73 samples from the four species. Species accumulation curves were built with the *specaccum* function of the *vegan* package (Oksanen and et al. 2018). An OTU table and species classification was generated for fungal communities separately from ITS2 and LSU sequencing, after singletons and doubletons reads were removed. Fungal OTU richness among samples was assessed with a Generalised Linear Model (GLM), built using the *lme4* package (Bates et al. 2015) of *R*, with fungal OTU richness as a response variable and beetle species and forest type as dependent variables. The Negative Binomial model was chosen, as it is suitable for over dispersed data. A post hoc pairwise comparison (Tukey HSD test at the 95% significance level) was carried out to compare the means among the different factors.

Sorensen and Jaccard indexes were used to calculate beta-diversity between sample pairs based on presence-absence data (richness), and the Bray-Curtis index was used for turnover based on the relative abundance of each OTU. The variation was visualised using multidimensional scaling MDS (*betapart* *R* package; (Baselga and Orme 2012), and the significance of the distribution was assessed using permutational multivariate analysis of variance (PERMANOVA) under a model for forest type, beetle species and their interaction, calculated with the *adonis* function of the *vegan* package (permutations=999) (Oksanen and et al. 2018). A permutational multivariate analysis of dispersion (PERMUTEST) was performed to evaluate the assumption of homogeneity of variance in the data by using the *betadisper* function (Anderson and Walsh 2013). The non-euclidean distances between the group centroids were also calculated to assess the group dispersion.

To evaluate the stringency of association of fungal OTUs with tree species and beetle hosts for each assembly, a multilevel pattern analysis was carried out by calculating Pearson’s phi coefficient of association (“p.g”) (Chytrý et al. 2002) between sample pairs, correcting this index to account for the differences in specimen numbers among the compared groups (function *multipatt* of the *indicspecies R* package; (De Cáceres et al. 2011). The identified OTUs for which the association values were significant were displayed as a heatmap (*aheatmap* function, NMF *R* package; (Gaujoux and Seoighe 2010).

## Results

Sequencing of 80 libraries produced 2,436,075 quality-filtered and merged reads for ITS2 and 1,742,119 reads for LSU, which resulted in 1157 OTUs from ITS2 and 548 OTUs from LSU after bioinformatics filtering and clustering (1546 and 632 OTUs if singleton and doubleton reads were retained and without applying rarefaction on each library). Identifications of OTUs at ≥70% confidence level obtained with IDtaxa, PROTAXfungi and RDP were higher for ITS2 than for LSU at all hierarchical levels from class to order, family, genus and species level (Figure 1). Yet, OTUs identified by one or multiple identifiers never exceeded 61.5% for ITS and 41.5% for LSU of the total OTUs. Identifications dropped consistently from class to species level, and with each hierarchical level an increasing proportion of identifications was provided by a single classifier only, indicating the growing uncertainty of taxonomic assignments. A classification at species-level was generally not possible for LSU because of the limitations of the databases that generally provide a taxonomy string to genus level only, and thus nearly 100% of the OTUs remained unidentified at this level. Nearly 50% of sequences were identified to species level with ITS but in virtually all cases only a single classifier produced these results (Figure 1).

The 73 beetle specimens remaining after rarefaction harboured a total of 1180 OTUs for ITS and 553 OTUs for LSU. These OTUs could be assigned to 24 classes, 66 orders, 129 families and 369 genera. Identification at class level revealed the presence of 23 classes for ITS2 and 17 classes for LSU. The dominant classes were Dothideomycetes for ITS2 and Sordariomycetes for LSU (Figure 2A, B, Table S1). ITS2 produced twice as many identified OTUs, and in the classes Leotiomycetes and Tremellomycetes more than five times as many, compared to LSU, due to the greater total number of OTUs and the higher proportion of them being fully identified. ITS2 also detected seven fungal classes not retrieved with LSU (Archaeorhizomycetes, Chytridiomycetes, Mucoromycetes, Orbiliomycetes, Spizellomycetes, Tritirachiomycetes and Ustilaginomycetes), while LSU recovered only one class not obtained with the ITS2 (Atractiellomycetes). Only for the Sordariomycetes and Agaricomycetes the proportion of OTUs detected with LSU was higher than with the ITS2 marker.

#### Effects of the amplified sub-region on the ecological analysis

Fungal communities obtained with either marker were compared with regard to total richness and differentiation across beetle species and forest type. For both markers, species accumulation curves displayed a similar shape, despite the roughly twice higher OTU number in ITS2, with a slow increase and not reaching a plateau, although LSU generally showed a more pronounced ‘shoulder’ indicating a fraction of OTUs that is encountered commonly in multiple samples. Across the different forest types, species accumulation in oak forest clearly lagged behind pine and spruce forests in both markers (Supplementary Figure S2).

Richness in a single-beetle extract ranged from 9 to 140 fungal OTUs (mean of 56) in ITS2 and from 11 to 109 fungal OTUs (mean 48) in LSU (Figure 2C, D). Despite some scatter among individual beetles, the number of OTUs per sample differed in a characteristic way between beetle species and forest types, and these differences were closely correlated in ITS2 and LSU, indicating that both markers detected a similar set of fungal species. This correlation was also evident at specimen level in the two outliers in each of the libraries corresponding to the same beetle individual. The variation in species richness explained by forest type and beetle species was broadly similar in ITS2 and LSU derived fungal communities (Table 2), but the LSU data attributed a greater proportion of the variation to the forest type alone (27.47% compared to 18.75% from ITS2), rather than to the interaction of forest type and beetle species. Both ITS2 and LSU explained around 7% of the variation to be due to beetle species.MDS plots on the OTU composition revealed a very similar pattern of community separation of the three forest types in ITS2 and LSU (Figure 3).

Tests of associations of OTUs to particular beetle species and forest types with the *indval* function revealed a large number of significantly positive and negative associations (Figure 4). Despite representing fewer OTUs in total, the number of OTUs with significant levels of association was slightly higher in LSU compared to ITS2 (60 versus 50 OTUs). Positive associations of OTUs were prevalent in pine and spruce forests, but much less in oak. Positive associations were particularly strong with *T. piniperda*, and to some extent *G. materiarius*, whereas positive associations with *Xs. germanus* and *Xb. saxesenii* were limited to a small number of oak associated OTUs. Most other associations in these species were negative; e.g. the pine/spruce associated OTUs were absent, despite the fact that both beetle species were also sampled from spruce. General patterns of OTU associations and non-associations were similar for the two xyleborine species, and they were quite similar with those in Oak. In contrast, association patterns in *T. piniperda* and *G. materiarius* were similar to Pine and Spruce (Figure 4). The similarity in these association patterns differed only slightly if assessed with the ITS2 and LSU-based OTUs (Figure 4), despite the fact that the OTUs themselves could not be linked up between both markers, as they mostly were not identified or the identifications did not overlap between the two marker sets.

#### OTU identifications across markers

Because it remained unclear if the observed similarities in community composition were based on the same set of OTUs, a phylogenetic approach was used to associate ITS2-based and LSU-based OTUs with each other, focusing on the class Sordariomycetes as a model system. OTUs were clustered at minimum similarity thresholds of 97, 98 and 99%, which resulted in between 120-150 OTUs for ITS and 80-120 OTUs for LSU classified as Sordariomycetes using the RDP classifier at confidence >80%(Table 3). OTU sequences from either marker were used for phylogenetic analysis together with publicly available full-length sequences covering all or most of the rRNA cluster, including the LSU and ITS2 regions, besides the SSU gene present in most accessions. These sequences served as a scaffold for the major orders of Sordariomycetes (full list of accessions in Suppl. Table S1), to which the metabarcode data were added, using clustering thresholds of 99% similarity for LSU and 98% for ITS2. These conditions were used because they generated a similar number of OTUs for each marker (Table 3), and thus potentially represent a similar set of species.

Maximum-likelihood trees for the three-gene reference alignment and OTUs from metabarcoding resolved basal relationships similar to those found in the literature (Hongsanan et al. 2017; Zhang et al. 2006) (Suppl. Figure S3). OTUs obtained from ITS and LSU were widely distributed across this tree, and their placement generally matched the order-level assignment obtained with the RDP classifier, which was dominated by the orders Hypocreales, Diaporthales, Ophiostomatales and Xylariales (Figure 5) (Suppl. Figure S4). Across most orders in the trees, ITS2 and LSU sequences were recovered in close proximity and the distribution of both types of sequences across orders was similar, showing that overall community diversity at the order level could equally be inferred using either ITS2 or LSU (Suppl. Figure S2). Order-level subsets of trees for these orders showed the placement of ITS2 and LSU sequence fragments relative to the reference set (Figures 6A-C). If both sequences are derived from the same genomic template in the metabarcoding amplification they were expected to be represented by one OTU representative sequence for each marker, and these sequences to fall in close proximity on the tree (taking the same phylogenetic position) (Figure 6D). However, due to their non-overlapping sequence, they can only be placed relative to the nearest full-length reference sequence. We found 15 instances where one ITS and one LSU barcode were in close proximity together with a reference sequence, potentially representing the same species. In an additional six instances, one or both of the barcodes formed a cluster on zero-length branches when matched to full length rRNA reference sequences, i.e. representing an exact match to an existing database entry.

Closed-reference clustering against the reference dataset to each order within Sordariomycetes by the RDP classifier revealed species-level matches for both ITS2 and LSU sequences (Figure 7A). Notably, four species had matches to both markers, i.e. the same species were amplified. In addition, one ITS2 sequence produced a hit not reciprocated in LSU. Vice versa, LSU sequences produced hits to a minimum of eight additional species not seen in ITS, which was increased to 11 and 17 species under the higher 98 and 99% threshold values, respectively, as the trees became increasingly populated with the additional taxa from splitting of larger OTUs (Figure 7B). Under these lower threshold values closely related sequences apparently were less affected by ‘lumping’, which obscured the true diversity in the sequencing mixture.

Where closely related reference sequences were missing, ITS2 and LSU sequences may be matched based on their phylogenetic proximity, but the ITS2 and OTU sequences obtained from a single genome may not appear as sister taxa because the gene sequences are non-overlapping and thus lack characters that could group them. We tested the degree to which ITS2 and LSU sequences interleave on the tree, by assessing phylogenetic clustering and dispersion with the NRI and NTI (Table 3). For ITS2, most values were negative, indicating overdispersion relative to the LSU sequences, except for the 99% similarity value, which produced positive NRI (clustering) possibly from selective oversplitting of OTUs that is not matched in the less variable LSU sequences. For LSU there was a progression from positive (clustering) at 97% similarity to negative (indicating overdispersion) at 99% similarity, which coincided with a near doubling in the number of OTUs (against only a small increase in the ITS2 data) (Table 3). This indicated that OTUs newly formed by splitting were not clustered, unlike the ITS2-derived OTUs, but instead were widely distributed on the tree and interleaved with the ITS2 sequences. A ‘mixed’ threshold value of 98% for ITS2 and 99% for LSU presented slightly negative NRI/NTI values for both markers (Table 3).

The detailed observations were confirmed by the global classification of OTUs at order level, which showed an increase in the proportion of identified OTUs with increasing threshold value for LSU, but not ITS2 (Figure 8). Both markers produced broadly similar proportions of the four dominant orders, Xyliariales, Ophiostomatales, Diaporthales and Hypocreales, but differed to various degrees in the assignment of the ‘small’ orders. It was also evident that OTU numbers in Ophiostomatales were comparatively lower in ITS2, as also suggested from the phylogenetic tree (Figure 5). This may be explained by the fact that the ITS2 forward primer binding site in this group differs from the consensus (Suppl. Figure S4)**.** The RDP classifications and tree-based assignments were largely in agreement regarding the relative proportion of order-level taxa, although generally the trees assigned a greater proportion of OTUs, reaching nearly 95%, and the result was more variable under the various similarity threshold values. This was particularly evident in ITS2, which saw hardly any change in the RDP classification under changing threshold values.

## Discussion

Metabarcoding has revolutionised the study of fungal communities, revealing the huge proportion of hitherto unobserved species, including the unexpectedly diverse communities associated with bark beetles (Tedersoo et al. 2019). However, these inferences are based on short sequences and lack the biological underpinning of conventional assessment of communities using fungal cultures. Independent corroboration of species limits is needed, and principally can be achieved by using multiple markers that confirm the same groupings independently (e.g. (DeSalle et al. 2005). This was attempted here, in part motivated by the claims that the most widely used ITS fungal barcode marker may not be suitable for efficient amplification of key groups of fungi (Hulcr et al. 2020). The test of phylogenetic congruence is complicated because the amplicons come from complex mixtures of species, which does not allow to establish genetic linkage (phasing) across the two markers, despite the close proximity of the ITS2 and LSU in the genome. Instead, an indirect approach had to be used that identifies the amplicons relative to full-length reference sequences separately for ITS2 and LSU, which was not straightforward given the highly incomplete and taxonomically incompatible databases available for both markers.

We addressed the equivalency of ITS and LSU for the characterisation of fungal communities, by (1) assessing the ecological associations at the higher taxonomic level for the entire fungal set, and (2) establishing explicit correspondences of species-level taxa in metabarcoding from both markers (for the class Sordariomycetes only). As we showed with both approaches, OTU identification is challenging and depends on the available reference databases, as well as the specific strategy for linking the metabarcode sequences into the taxonomic system. Taxonomic classifiers are now widely used and are becoming increasingly sophisticated. However, we note that placement is possible mostly to higher taxonomic levels, in line with comparisons of taxonomic classification software showing that the RDP classifier performed best in taxonomic classification to order (Richardson et al. 2017). Only a fairly small proportion of reads can be placed to genus or species level, in particular for the LSU marker (Figure 1). Of those OTUs that could be placed, the exact taxonomic membership could be established only with one of the three classifiers. These difficulties in identifying species greatly compromised the comparison of community composition obtained with either marker, which was impossible to link with this approach, as virtually none of the species level entities encountered were labelled with the same Linnaean binomial (Figure 3). Thus, we did not obtain a corroboration of community composition from using both markers via taxonomic identification of reads against their respective reference databases.

In contrast, simple counts of OTU numbers (a proxy of species richness) produced a good correlation between both markers in several key parameters describing the community composition. First, plots of the number of OTUs obtained from each treatment (beetle species, forest type) from the ITS2 and LSU data closely mirror each other (Figure 1). Equally, the proportion of explained OTU diversity by beetle species, forest type and beetle x forest interactions was closely similar between both markers, even if the absolute number of OTUs was generally much lower in LSU (Table 2). For both markers, communities from different forest types occupy remarkably similar portions of the multivariate space (Figure 4). In addition, the higher-level composition of fungal communities associated with each beetle species is very similar whether generated with ITS2 or LSU (Figure 2), which also holds for the composition of classes within the order Sordariomycetes (Figure 8). The broad patterns of individual OTU associations in the *indval* analysis also show similar affinities with the beetle species and tree type (Figure 4), even if the correspondences of species identity between ITS2 and LSU datasets could not be determined. All of these findings point to a high level of corroboration between both markers and provide justification for the widely used approach of fungal community analysis based on higher level classification and read abundances. The latter is particularly remarkable given the frequently raised concern about PCR bias in relative read abundance (Bálint et al. 2016; Krehenwinkel et al. 2017). Thus, even a single metabarcode marker can safely represent the broad ecological trends determining fungal communities.

Yet, the difficulty of linking these metabarcoding sequences amongst markers leaves great uncertainty about the biological relevance of the community data, which still may represent entirely different species within the major taxonomic groups recovered by either marker, as already suggested for the Ophiostomatales (Hulcr et al., 2020). Thus, ultimately the metabarcoding approach may fall short of linking any particular fungal species to a beetle, unlike the conventional approaches of culturing specific isolates that were used to reveal the symbioses of particular fungal and beetle species of direct relevance to the ecological processes. Phylogenetic analysis of individual sequences can improve the precision of placement, beyond the assignment to a broad taxonomic group, and using the phylogenetic position of particular ITS2 and LSU sequences relative to each other can potentially link the corresponding reads representing a given species from either marker, thus reconciling the distinct ‘views’ of the true community composition as revealed by each marker (Figure 6D).

As shown by our extensive analysis of the Sordariomycetes, phylogenetic analyses provide clear insight into the taxonomic makeup of fungal communities and the most appropriate treatment of fungal metabarcodes for species delimitation. First, we found that the classifiers are broadly in agreement with the phylogenetic analysis. However, average RDP classifier confidence scores varied from order to order, with LSU assignments less confident overall, ranging from an average of 41% in Sordariomycetes, to values generally between 80-100% in the ITS sequences (see Suppl. Table S2). Some OTUs were not identified beyond the class level by the RDP classifier, despite clear placement in the tree. Notably, the order Myrmecridiales was missing entirely from the classifier results, despite the presence of several OTUs placed clearly within the order and OTUs matching *Myrmecridium schulzeri* found in the closed-reference clustering at all three thresholds (Figures 7A, B). This outcome highlights the potential shortcomings of using classifiers that are dependent on reference databases and probabilistic assessments of short *k-mers*, which may be particularly problematic due to the limited sequence length of metabarcoding reads (Bacci et al. 2015; Porras-Alfaro et al. 2014; Wang et al. 2007; Xue et al. 2019).

Second, the phylogenetic analysis attempted to determine if both markers reveal the same species-level entities. Under ideal circumstances, each species is represented by exactly one sequence each of LSU and ITS, and these two sequences from both markers find themselves in the same position of the tree. Thus, the 1:1 represented sequences of ITS and LSU should be interleaved on the tree, which predicts a uniform phylogenetic distribution of both markers, contrary to a scenario of overdispersion or clustering of either type (Figure 6D). However, deviations from the uniform distribution are due to two factors; the non-overlapping sequence data that only can place ITS2 and LSU sequences relative to full-length reference sequences rather than to each other, and the problem of similarity thresholds that produce different numbers of potential species representatives by either lumping (too few species) or oversplitting (too many species) of OTUs in one or both of these loci. We here used the NRI/NTI framework to assess the uniformity of distributions of ITS and LSU sequences relative to each other on the phylogenetic tree.

Overall, the increase of the similarity threshold had a greater impact on the LSU than ITS2, almost doubling in numbers of recognised OTUs versus a small increase only (Table 3), and parity of OTUs in both markers was greatest at a ‘mixed’ threshold of 99% for LSU and 98% for ITS2. Slightly negative NRI/NTI values under these OTU thresholds (Table 3) approximate a pattern of overdispersed ‘communities’ of LSU and ITS2 sequences, as expected for the 1:1 correspondence of each marker, although the exact relationships were more complex. The 99% threshold for LSU is also supported by the greater matches in the closed-reference clustering (Figure 7). The similarities in OTU counts and relative NRI/NTI values thus were considered the best estimate of the OTU diversity in each marker. Placement in proximity on the tree therefore would suggest that the respective ITS2 and LSU sequences in fact are derived from closely related fungal species, if not from the same genomic template. Frequently in our analysis this was corroborated by the fact that these closely related ITS2 and LSU sequences were obtained from the same specimen sample (not shown).

There are uncertainties associated with this inference, as it is well known that there is great variation in the rate of sequence evolution. Across many fungal classes, intraspecific ITS variability varies considerably, highlighting the challenges and inevitable shortcomings resulting from the selection of a uniform OTU clustering threshold (Nilsson et al. 2008). For example, while a 97% clustering threshold is generally accepted (Kõljalg et al. 2013; Tedersoo et al. 2015a), in one study intraspecific ITS variability in *Xylaria hypoxylon* was found to be 24.2%. Thus, the abundance of ITS2 OTUs in Xylariales relative to LSU OTUs (Figure 6) may be the result of oversplitting of highly variable ITS2 OTUs (Nilsson et al. 2008). Yet, whether these OTUs correspond to biological species or some other hierarchical level, the approach generally is capable of linking close relatives across different markers. Reference sequences spanning both markers are required, but as shown for the Sordariomycetes, even an incomplete set can provide the scaffold for placing non-overlapping sequences, and in several instances the idealized placement of OTUs and reference sequences was found, in some cases across all clustering thresholds (Figure 5), while in many cases some uncertainties remain where reference sequences are distant. Matching of ITS2 and LSU sequences was even possible in the Ophistomatales, despite the deviation in the ITS2 primer binding site in this group (Suppl. Figure S4), as the bias against the amplification presumably is overcome by permissive PCR conditions, and similar effects can be expected in other groups where such variation may exist, although we did not find any evidence in the taxonomically broad reference sequences used here.

## Conclusion

Two streams of evidence suggest the equivalency of the two most widely used metabarcoding markers, ITS2 and LSU, in assessing fungal communities associated with insects: community-level diversity metrics showed great consistency when inferred with either marker, and the taxonomic and phylogenetic distribution of taxa recovered were highly similar. Metabarcoding of these communities only began recently, revealing a huge diversity of mostly anonymous (“dark”) fungal isolates that have been used to draw conclusions on biodiversity patterns, but the taxonomic assignment and estimates of species richness were lacking independent corroboration. The use of two independent metabarcode markers changes this situation dramatically, as each inference is independent, and although the markers are genetically linked, i.e. likely represent the same evolutionary history, the experimental approach is treating these samples as separate inferences of the fungal communities. This provides strong support for the metabarcoding approach of fungal diversity directly from the insect specimens, which can reveal critical information about the distribution, spread and epidemic conditions of fungal tree pathogens by sampling the insect specimens.

## Acknowledgments

## This work was funded by a PhD studentship of the John Spedan Lewis Foundation to AC-E. We thank Dr Ruth Bone for grant management and scientific advice. We acknowledge technical help from the Core Research Laboratory at the NHM.

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## Figure Legends

Figure 1. The proportion of fungi classified with IDtaxa, RDP and PROTAXfungi from class to species level. “ALL” refers to the proportion of OTUs for which the three classifiers agreed in their classification.

Figure 2. Classification of OTUs and number of OTUs associated with each beetle. Top panel: The proportion of OTUs identified as members of a fungal Class determined by the LSU (left panel) and ITS (right panel) regions, separate for each of the three forest types and beetle species, as given at the bottom of the figure. Proportions are based on the total OTU numbers from 20 specimens of each beetle/forest type category; for the Spruce Forest, only nine and four *X. germanus* and *X. saxesenii* specimens were retained after rarefaction. Bottom panel: The number of fungal OTUs per beetle specimen, separate for each beetle species and forest type, for LSU (left panel) and ITS (right panel).

Figure 3. MDS ordination of fungal communities obtained from each beetle specimens collected in Oak, Pine and Spruce forests, based on the LSU (left panel) and ITS (right panel) data. Stress values fell within acceptable ranges (<0.2).

Figure 4. Similarity of associations of individual fungal OTUs with beetle species and forest type. The heatmap shows the strength of association (positive or negative) for OTUs with significant values in the IndVal analysis. The dendrograms on either side show the degree of similarity of fungal communities of significantly associated OTUs, based on the ITS2 (left panel) and LSU (right panel) data. Note the similar clustering obtained with both markers. Fungal OTUs (at bottom) were classified to the genus or species level where possible; they are shown in a random order and cannot be linked taxonomically between both markers.

Figure 5. Phylogenetic tree of Sordariomycetes. The circular tree shows the orders represented by the colour of branches, and the distribution of ITS (blue) and LSU (green) relative to the full-length ITS-LSU reference set (yellow) on the outer circle. Note the limited presence of ITS sequences in the Ophiostomatales (in top right quadrant). Red dots mark the three orders Xylariales, Ophiostomatales and Diaporthales shown in greater detail in Figure 6. The orange dot marks the Myrmecridiales whose presence in the sample was only detected with the tree-based method, but not with the classifiers.

Figure 6. Order-level trees with mixed OTU clustering thresholds (99% LSU, 98% ITS2). Full tree in supplementary materials. *Leotia lubrica* was used as the outgroup (not pictured). Brackets indicate reference taxa linked to an ITS2 and/or LSU OTU, with colours indicating potential splitting/lumping (blue, splitting; green, lumping; orange, 1:1) by one of the markers as illustrated in Figure 6B. The interaction of non-overlapping ITS2 and LSU sequences in the combined community tree is only possible through their placement relative to the full-length reference sequences that include both markers (labelled “Reference”). The combined OTU inference shows how the different clustering artefacts may present themselves on the tree.

Figure 7. Results from the closed reference clustering of OTUs at each clustering threshold against composite LSU/ITS2 reference sequences. Figure 7A. LSU matches in green, ITS2 matches in blue, linked matches (for which both an ITS2 and LSU OTU were matched to a reference sequence of the same species) in yellow. Underlined taxa indicate new matches at each clustering threshold. Figure 7B. Phylogenetic tree of LSU OTUs under increasingly stringent clustering thresholds, with arrows marking newly added taxa as threshold values are increased.

Figure 8. Classification of OTUs from metabarcoding with LSU (left panel) and ITS (right panel) markers based on the RDP classifier and the phylogenetic tree, under increasing threshold values.

# Tables

Table 1. Beetle species and number of the specimens per forest type included in the study along with information related to their origin and main fungal structures.

Table

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Table 2. Overview of the GLM analysis results, showing the significant *p* values, the percentage of explained variance and the F parameter for every factor included in the model.

Graphical user interface, application

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Table 3: Net relatedness index (NRI), nearest taxon index (NTI), and number of OTUs recovered for LSU and ITS2 across all clustering thresholds. “Mixed” refers to a clustering threshold of 99% for LSU and 98% for ITS2. Reference sequences were included when building the trees used, though pruned (leaving only OTUs in the tree) for the above calculations.

Table

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# Figures

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Figure 1. The proportion of fungi classified with IDtaxa, RDP and PROTAXfungi from class to species level. “ALL” refers to the proportion of OTUs for which the three classifiers agreed in their classification.

Chart, timeline, box and whisker chart

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Figure 2. Classification of OTUs and number of OTUs associated with each beetle. Top panel: The proportion of OTUs identified as members of a fungal Class determined by the LSU (left panel) and ITS (right panel) regions, separate for each of the three forest types and beetle species, as given at the bottom of the figure. Proportions are based on the total OTU numbers from 20 specimens of each beetle/forest type category; for the Spruce Forest, only nine and four *X. germanus* and *X. saxesenii* specimens were retained after rarefaction. Bottom panel: The number of fungal OTUs per beetle specimen, separate for each beetle species and forest type, for LSU (left panel) and ITS (right panel).

Chart, scatter chart, bubble chart

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Diagram

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Chart, bar chart

Description automatically generated

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