Diversity, taxonomic composition, and functional aspects of fungal communities in living, senesced, and fallen leaves at five sites across North America

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ABSTRACT

Background. Fungal endophytes inhabit symptomless, living tissues of all major plant lineages to form one of earth’s most prevalent groups of symbionts. Many reproduce from senesced and/or decomposing leaves and can produce extracellular leaf-degrading enzymes, blurring the line between symbiotrophy and saprotrophy. To better understand the endophyte–saprotroph continuum we compared fungal communities and functional traits of focal strains isolated from living leaves to those isolated from leaves after senescence and decomposition, with a focus on foliage of woody plants in five biogeographic provinces ranging from tundra to subtropical scrub forest.

Methods. We cultured fungi from the interior of surface-sterilized leaves that were living at the time of sampling (i.e., endophytes), leaves that were dead and were retained in plant canopies (dead leaf fungi, DLF), and fallen leaves (leaf litter fungi, LLF) from 3–4 species of woody plants in each of five sites in North America. Our sampling encompassed 18 plant species representing two families of Pinophyta and five families of Angiospermae. Diversity and composition of fungal communities within and among leaf life stages, hosts, and sites were compared using ITS-partial LSU rDNA data. We evaluated substrate use and enzyme activity by a subset of fungi isolated only from living tissues vs. fungi isolated only from non-living leaves.

Results. Across the diverse biomes and plant taxa surveyed here, culturable fungi from living leaves were isolated less frequently and were less diverse than those isolated from non-living leaves. Fungal communities in living leaves also differed detectably in composition from communities in dead leaves and leaf litter within focal sites and host taxa, regardless of differential weighting of rare and abundant fungi. All focal isolates grew on cellulose, lignin, and pectin as sole carbon sources, but none displayed ligninolytic or pectinolytic activity in vitro. Cellulolytic activity differed among fungal classes. Within Dothideomycetes, activity differed significantly between fungi from living vs. non-living leaves, but such differences were not observed in Sordariomycetes.

Discussion. Although some fungi with endophytic life stages clearly persist for periods of time in leaves after senescence and incorporation into leaf litter, our sampling across diverse biomes and host lineages detected consistent differences between fungal assemblages in living vs. non-living leaves, reflecting incursion by fungi from the leaf exterior after leaf death and as leaves begin to decompose. However, fungi found...
only in living leaves do not differ consistently in cellulolytic activity from those fungi detected thus far only in dead leaves. Future analyses should consider Basidiomycota in addition to the Ascomycota fungi evaluated here, and should explore more dimensions of functional traits and persistence to further define the endophytism-to-saprotrophy continuum.

Subjects Ecology, Microbiology, Mycology
Keywords Endophytic fungi, Extracellular enzymes, Diversity, Ascomycota, Plant-fungal symbioses, Saprotroph

INTRODUCTION

Fungal endophytes inhabit symptomless, living photosynthetic tissues of all major plant lineages to form one of earth’s most prevalent groups of symbionts (e.g., Arnold et al., 2010; U'Ren et al., 2012; Zimmerman & Vitousek, 2012; Davey et al., 2013; Balint et al., 2015). Known from a wide range of biological provinces and agroecosystems, endophytes are a ubiquitous feature of plant biology (e.g., Arnold & Lutzoni, 2007; Saunders, Glenn & Kohn, 2010). Although classified together due to ecological similarities (i.e., colonization and transmission patterns, in planta biodiversity, and host interactions; see Rodriguez et al., 2009), endophytic fungi represent a diversity of evolutionary histories, life history strategies, and functional traits that are only beginning to be understood (reviewed by Porras-Alfaro & Bayman, 2011).

Endophytes that inhabit photosynthetic tissues of most plants are horizontally transmitted, form localized infections in aerial tissues, and represent highly diverse and often novel lineages (e.g., Arnold et al., 2009; Gazis et al., 2012; Chen et al., 2015; U'Ren et al., 2016) (Class 3 endophytes sensu Rodriguez et al., 2009; hereafter endophytes). Many reproduce from senesced and/or decomposing leaves (Fröhlich & Hyde, 1999; Promputtha et al., 2007; Promputtha et al., 2010; U'Ren et al., 2010; Chaverri & Gazis, 2011; He et al., 2012). Some also produce extracellular leaf-degrading enzymes (Carroll & Petrini, 1983; Korkama-Rajala, Müller & Pennanen, 2008; Osono & Hirose, 2011; Promputtha et al., 2010; Sun, Guo & Hyde, 2011). Thus many endophytes blur the line between symbiotrophy (during the endophytic phase) and saprotrophy (when they occur in association with dead tissue), creating a challenge for estimating trophic modes and the scale of fungal diversity based on species richness in particular functional groups.

The prevalence of saprotrophic life phases among endophytes and the dynamics of such fungi on the endophyte-to-saprotroph continuum are not yet clear. Current evidence suggests that different groups of fungi may persist for longer or shorter periods in senesced leaves, but the relevance of host lineages and abiotic factors is not yet understood in many cases. For example, Osono (2006) estimated that approximately two thirds of fungi with endophytic life stages can persist in and degrade leaf litter. U'Ren et al. (2016) found that 74% of Xylariaceae taxa were represented by isolates from asymptomatic plant tissues, as well as senesced/decomposing leaves, wood, bark, fruits, and/or flowers.
Some endophytes have ligninolytic and cellulytic activity, cause mass loss from dead plant tissues \textit{in vitro}, and persist as litter decomposers over multiple years (e.g., some \textit{Lophodermium} spp. and xylariaceous fungi; see Koide, Osono & Takeda, 2005; Osono, 2006; Korkama-Rajala, Müller & Pennanen, 2008; Osono & Hirose, 2009; Osono & Hirose, 2011; see also Lindahl et al., 2007; Yuan & Chen, 2014). The presence of such fungi can increase respiration rates and lignocellulolytic activity in litter, altering the litter substrate and the activity of subsequent decomposers (see Koide, Osono & Takeda, 2005; Šnajdr et al., 2011; He et al., 2012; Lin et al., 2015). In contrast, other fungi with endophytic life phases may occupy litter only transiently, quickly sporulating from senesced and decomposing leaves to infect living tissues (e.g., \textit{Rhabdocline parkeri} on \textit{Pseudotsuga menziesii} and \textit{Coccomyes nipponicum} on \textit{Camellia japonica}; Stone, 1987; Koide, Osono & Takeda, 2005). The impact of such fungi on litter degradation is less well known. Such patterns can be documented via culture-based studies, but also by culture-free methods. For example, fungal genotypes that were dominant in the phyllosphere of living leaves of \textit{Quercus petraea} disappeared by two to four months post abscission (Voříšková & Baldrian, 2013). Similarly, fungal communities in living leaves of European beech (\textit{Fagus sylvatica}) in autumn differed from fungi inhabiting leaf litter and bark in the following spring (Unterseher, Peršoh & Schnittler, 2013; but see Peršoh et al., 2013).

Together, these studies speak to continuity between fungi that occur within living and non-living leaves (see also Promputtha et al., 2007; Chaverri & Gazis, 2011; Sun, Guo & Hyde, 2011; Peršoh et al., 2013; Voříšková & Baldrian, 2013; Unterseher, Peršoh & Schnittler, 2013). However, it is not clear whether such continuity is consistent in the distinctive plant and fungal communities present in different biomes, nor whether patterns of persistence vary among phylogenetically diverse plants. It is also unclear whether endophytes in a given plant species might occur in non-living leaves of co-occurring plant species: if undetected, such occurrences could alter our understanding of the endophyte–saprotroph continuum. Finally, little is known about functional traits of these fungi, raising the question: do strains that occur only living leaves differ in substrate use and enzyme activity from those that consistently occur in both living and non-living leaves?

The goal of this study was to explore the occurrence of endophytes in leaves after senescence and incorporation into the leaf litter, with a focus on woody plants in five biogeographic provinces ranging from tundra to subtropical scrub forest. Our surveys included >7,000 tissue pieces from living leaves, senesced leaves in plant canopies, and recently fallen leaf litter from 3–4 plant species in each site, yielding >2,000 fungal cultures. Here, we address (1) the degree to which fungal communities within leaves differ as a function of leaf type (living, senesced, or in leaf litter) at a given point in time; (2) whether such patterns are consistent among diverse biogeographic/bioclimatic zones and host lineages; and (3) how functional differences in carbon substrate utilization and enzyme activity reflect the occurrence of particular fungi in living vs. non-living leaves.

**MATERIALS AND METHODS**

We collected living leaves, senesced leaves in plant canopies, and leaf litter from five sites representing distinct environmental, biological, and biogeographic regions of
North America: the Madrean Sky Island Archipelago of southeastern Arizona (AZC); the Appalachian Mountains of western North Carolina (NCH); sub-tropical scrub forest in Florida (FLA); Beringian tundra and boreal forest in the Seward Peninsula ecoregion of western Alaska (AKN); and inland, subalpine tundra in the Interior Highlands of east-central Alaska (AKE) (see U’Ren et al., 2012 for site details). Within each site, we selected three to four species of woody plants that were representative of the community of the region (Table 1; Hultén, 1965; Radford, Ahles & Bell, 1968; Barton, 1994; http://www.archbold-station.org/html/datapub/species/lists/plantlist.html). Overall we examined 18 host species representing two families of Pinophyta and five families of Angiospermae (Table 1).

In each site we collected fresh tissues from one individual of each species in each of three replicate microsites located ca. 30 m apart along a 100 m transect, for a total of 9–12 host individuals per site. Within each microsite, focal plants occurred within close proximity to one another (<1–10 m apart), which allowed us to decouple spatial heterogeneity from host associations (U’Ren et al., 2010). The only exception was Picea in AKN (see Table 1; U’Ren et al., 2012).

From each individual we collected three small branches containing both healthy leaves and dead leaves attached to branches, as well as leaves in below-crown leaf litter in an intermediate state of decomposition (i.e., intact leaves with obvious changes in color and texture). Leaves of deciduous plants were selected to represent the same year of growth.

**Isolation of fungi**

Plant material was transported in plastic bags to the laboratory and processed within 24 h (except AKN; 48 h) following U’Ren et al. (2010). Each sample was washed thoroughly in running tap water for 30 sec. Although putatively saprotrophic fungi can be isolated from spores or hyphae on leaf surfaces (e.g., Promputtha et al., 2002; Promputtha et al., 2007), all leaves were sterilized to exclude fungi that were incidental on leaf surfaces and to maximize comparability with endophytic fungi (which are isolated from surface-sterilized leaves).

Leaves were cut into 2 mm$^2$ segments and surface-sterilized by agitating sequentially in 95% ethanol for 10 sec, 10% bleach (0.5 % NaOCl) for 2 min, and 70% ethanol for 2 min (Arnold et al., 2007). Segments were surface-dried under sterile conditions before being placed on 2% malt extract agar (MEA) in Petri dishes (16 tissue segments/dish) or 1.5 ml microcentrifuge tubes (1 segment/tube). Plates or tubes were sealed with Parafilm and incubated under ambient light/dark condition at room temperature (ca. 21.5 °C) for up to one year. Emergent fungi were isolated into pure culture, vouchered in sterile water, and deposited at the Robert L. Gilbertson Mycological Herbarium at the University of Arizona (ARIZ) (Table S1). Overall, 7,725 tissue segments were placed into culture (Table 2). All samples relevant to this work were handled in accordance with standard operating procedures for USDA permit P526P-1400151.

**DNA extraction, PCR, and sequencing**

Total genomic DNA was extracted directly from each isolate following Arnold & Lutzoni (2007). The nuclear ribosomal internal transcribed spacers and 5.8s gene
Table 1  Isolation frequency, richness and diversity of cultivable fungi from the interior of surface-sterilized leaves, including living leaves (i.e., endophytic fungi), fungi from dead leaves in canopies of woody plants (DLF), and fungi from leaf litter (LLF), from 18 plant species in five sites across North America. Site abbreviations match those in U'Ren et al. (2012): (AZC, Chiricahua Mountains, Arizona; NCH, Highlands Biological Station, North Carolina; FLA, Archbold Biological Station, Florida; AKE, Eagle Summit, Alaska; AKN, Nome, Alaska).

| Site | Host family | Host species | Fungal type | Leaf segments | Isolates recovered | Isolation frequency/ microsite ± SD | Isolates sequenced (%) | Basidiomycota sequences | Ascomycota sequences (Putative species) | Ascomycota ACE (95% CI) | Fisher’s alpha |
|------|-------------|--------------|-------------|---------------|-------------------|-------------------------------------|------------------------|---------------------|-------------------------------|---------------------|---------------|
| AZC  | Cupressaceae | Juniperus deppeana | endophyte | 144 | 22 | 0.15 ± 0.12 | 19 (86%) | 0 | 19 (10) | 27.2 (15.9, 16.3) | 8.5 |
|      |             |              | DLF        | 141 | 87 | 0.62 ± 0.18 | 82 (94%) | 0 | 82 (24) | 50.5 (36.2, 81.5) | 11.4 |
|      |             |              | LLF        | 144 | 43 | 0.30 ± 0.09 | 37 (86%) | 0 | 37 (18) | 22.8 (19.1, 38.0) | 13.8 |
| AZC  | Pinaceae    | Pinus arizonica var. arizonica | endophyte | 144 | 9 | 0.06 ± 0.02 | 9 (100%) | 0 | 9 (2) | 3.8 (2.2, 17.0) | 0.8 |
|      |             |              | DLF        | 144 | 36 | 0.25 ± 0.40 | 31 (86%) | 0 | 31 (3) | N/A | 0.8 |
|      |             |              | LLF        | 143 | 57 | 0.40 ± 0.35 | 55 (96%) | 0 | 55 (8) | 23.2 (13.5, 49.9) | 2.6 |
| AZC  | Pinaceae    | Pseudotsuga menziesii | endophyte | 144 | 21 | 0.15 ± 0.22 | 16 (76%) | 0 | 16 (3) | 4.0 (3.2, 7.1) | 1.1 |
|      |             |              | DLF        | 142 | 29 | 0.20 ± 0.21 | 19 (66%) | 0 | 19 (3) | NA | 1.0 |
|      |             |              | LLF        | 144 | 29 | 0.20 ± 0.03 | 25 (86%) | 0 | 25 (9) | 20.7 (12.7, 46.0) | 5.0 |
| AZC  | Fagaceae    | Quercus rugosa | endophyte | 130 | 0 | 0 | N/A | N/A | N/A | N/A | N/A |
|      |             |              | DLF        | 144 | 34 | 0.24 ± 0.39 | 32 (97%) | 0 | 32 (7) | N/A | 2.8 |
|      |             |              | LLF        | 143 | 29 | 0.20 ± 0.18 | 27 (90%) | 0 | 27 (7) | 9.5 (7.3, 25.3) | 3.1 |
| NCH  | Pinaceae    | Pinus strobus | endophyte | 114 | 14 | 0.14 ± 0.11 | 13 (93%) | 0 | 13 (6) | 30.6 (15.1, 72.5) | 4.3 |
|      |             |              | DLF        | 144 | 63 | 0.44 ± 0.31 | 61 (97%) | 1 | 60 (9) | 30.0 (10.6, 277.5) | 2.9 |
|      |             |              | LLF        | 144 | 62 | 0.43 ± 0.28 | 49 (79%) | 0 | 49 (16) | 25.9 (18.5, 55.2) | 8.3 |
| NCH  | Pinaceae    | Tsuga canadensis | endophyte | 144 | 54 | 0.38 ± 0.26 | 49 (91%) | 0 | 49 (13) | 19 (16.5, 23.4) | 5.8 |
|      |             |              | DLF        | 144 | 53 | 0.37 ± 0.39 | 42 (79%) | 0 | 42 (24) | 162.2 (108.9, 248.8) | 23.3 |
|      |             |              | LLF        | 144 | 123 | 0.85 ± 0.08 | 95 (77%) | 2 | 93 (26) | 48.0 (35.8, 75.4) | 12.0 |

(continued on next page)
| Site | Host family | Host species | Fungal type | Leaf segments | Isolates recovered | Isolation frequency/ microsite ± SD | Isolates sequenced (%) | Basidiomycota sequences | Ascomycota sequences (Putative species) | Ascomycota ACE (95% CI) | Fisher’s alpha |
|------|-------------|--------------|-------------|---------------|-------------------|---------------------------------|-------------------------|------------------------|---------------------------------|------------------------|---------------|
| NCH  | Ericaceae   | Kalmia latifolia L. | endophyte | 144 | 29 | 0.20 ± 0.31 | 29 (100%) | 0 | 29 (6) | 9.8 (7.6, 15.0) | 2.3 |
|      |             |              | DLF 144 | 71 | 0.49 ± 0.34 | 65 (92%) | 0 | 65 (14) | 43.4 (26.6, 82.4) | 5.5 |
|      |             |              | LLF 144 | 114 | 0.79 ± 0.15 | 91 (80%) | 0 | 91 (18) | 66.1 (41.3, 117.5) | 6.7 |
| NCH  | Fagaceae    | Quercus montana | endophyte | 144 | 60 | 0.42 ± 0.33 | 57 (95%) | 0 | 57 (6) | 25.2 (11.5, 73.1) | 1.7 |
|      |             |              | DLF 144 | 58 | 0.40 ± 0.42 | 47 (81%) | 0 | 47 (12) | 170.2 (102.8, 287.5) | 5.2 |
|      |             |              | LLF 144 | 46 | 0.32 ± 0.17 | 27 (59%) | 0 | 27 (13) | 38.9 (23.6, 76.4) | 9.9 |
| FLA  | Pinaceae    | Pinus elliottii | endophyte | 144 | 138 | 0.96 ± 0.04 | 130 (94%) | 57 | 73 (32) | 60.6 (42.5, 109.9) | 21.7 |
|      |             |              | DLF 144 | 102 | 0.71 ± 0.29 | 84 (82%) | 3 | 81 (32) | 126.6 (82.0, 210.9) | 19.5 |
|      |             |              | LLF 144 | 100 | 0.69 ± 0.23 | 85 (85%) | 0 | 85 (28) | 53.0 (36.6, 100.8) | 14.6 |
| FLA  | Pinaceae    | Pinus clausa | endophyte | 144 | 52 | 0.36 ± 0.07 | 47 (90%) | 29 | 18 (11) | 111.1 (59.3, 218.8) | 12.0 |
|      |             |              | DLF 144 | 88 | 0.61 ± 0.33 | 75 (85%) | 2 | 73 (30) | 88.9 (61.5, 140.2) | 19.0 |
|      |             |              | LLF 144 | 104 | 0.72 ± 0.10 | 88 (85%) | 3 | 85 (23) | 38.0 (27.6, 71.9) | 10.4 |
| FLA  | Arecaceae   | Serenoa repens | endophyte | 144 | 65 | 0.45 ± 0.40 | 58 (89%) | 17 | 41 (10) | 17.4 (12.1, 35.9) | 4.2 |
|      |             |              | DLF 144 | 81 | 0.56 ± 0.21 | 56 (69%) | 1 | 55 (33) | 154.3 (105.4, 236.4) | 34.8 |
|      |             |              | LLF 144 | 73 | 0.51 ± 0.17 | 58 (79%) | 2 | 56 (39) | 70.5 (62.3, 81.5) | 57.0 |
| FLA  | Fagaceae    | Quercus inopina | endophyte | 144 | 80 | 0.56 ± 0.25 | 51 (64%) | 1 | 50 (19) | 89.0 (55.7, 152.6) | 11.2 |
|      |             |              | DLF 144 | 95 | 0.66 ± 0.50 | 67 (71%) | 0 | 67 (28) | 119.4 (81.4, 184.5) | 18.1 |
|      |             |              | LLF 144 | 100 | 0.69 ± 0.35 | 72 (72%) | 0 | 72 (41) | 153.6 (107.1, 232.9) | 39.5 |

(continued on next page)
| Site | Host family | Host species | Fungal type | Leaf segments\(^b\) | Isolates recovered | Isolation frequency/ microsite ± SD | Isolates sequenced (%) | Basidiomycota sequences | Ascomycota sequences (Putative species) | Ascomycota ACE (95% CI)\(^c\) | Fisher’s alpha |
|------|-------------|--------------|-------------|---------------------|--------------------|--------------------------------------|-------------------------|--------------------------|------------------------------------------|-----------------------------|---------------|
| AKE  | Pinaceae    | *Picea glauca* | endophyte   | 144 8               | 0.06 ± 0.03        | 7 (88%)                              | 1                       | 6 (6)                    | N/A                       | N/A                         |               |
|      |             |              |             | DLF 144 43         | 0.30 ± 0.16        | 36 (84%)                             | 0                       | 36 (17)                  | 48.8 (31.3, 87.7)           | 12.6                        |               |
|      |             |              |             | LLF 144 77         | 0.53 ± 0.32        | 64 (83%)                             | 1                       | 63 (29)                  | 51.1 (36.9, 91.1)           | 20.8                        |               |
| AKE  | Betulaceae  | *Betula nana* | endophyte   | 144 2               | 0.01 ± 0.01        | 2 (100%)                             | 0                       | 2 (2)                    | N/A                       | N/A                         |               |
|      |             |              |             | DLF 144 10         | 0.07 ± 0.05        | 10 (100%)                            | 0                       | 10 (6)                   | 65.7 (28.8, 162.5)          | 6.3                         |               |
|      |             |              |             | LLF 144 37         | 0.26 ± 0.25        | 36 (97%)                             | 0                       | 36 (18)                  | 48.0 (30.9, 88.0)           | 14.3                        |               |
| AKE  | Salicaceae  | *Salix pulchra* | endophyte | 144 3               | 0.02 ± 0.02        | 2 (67%)                              | 0                       | 2 (2)                    | N/A                       | N/A                         |               |
|      |             |              |             | DLF 144 15         | 0.10 ± 0.04        | 12 (80%)                             | 0                       | 12 (9)                   | 26.8 (12.3, 104.3)          | 16.4                        |               |
|      |             |              |             | LLF 144 19         | 0.13 ± 0.10        | 13 (68%)                             | 0                       | 13 (9)                   | 14.6 (12.9, 17.1)           | 12.9                        |               |
| AKN  | Pinaceae    | *Picea glauca* | endophyte   | 144 68              | 0.47 ± 0.37        | 65 (96%)                             | 0                       | 65 (8)                   | 9.2 (8.1, 18.8)             | 2.4                         |               |
|      |             |              |             | DLF 144 75         | 0.52 ± 0.15        | 60 (80%)                             | 0                       | 60 (20)                  | 42.0 (29.6, 70.1)           | 10.5                        |               |
|      |             |              |             | LLF 144 35         | 0.24 ± 0.23        | 32 (91%)                             | 5                       | 27 (11)                  | 17.9 (12.4, 45.0)           | 6.9                         |               |
| AKN  | Salicaceae  | *Salix pulchra* | endophyte | 144 0               | 0                  | N/A                                  | N/A                     | N/A                      | N/A                       | N/A                         |               |
|      |             |              |             | DLF 144 0           | 0                  | NA                                   | N/A                     | N/A                      | N/A                       | N/A                         |               |
|      |             |              |             | LLF 144 1           | 0.01 ± 0.01        | N/A                                  | N/A                     | N/A                      | N/A                       | N/A                         |               |
| AKN  | Betulaceae  | *Betula nana* | endophyte   | 144 0               | 0                  | N/A                                  | N/A                     | N/A                      | N/A                       | N/A                         |               |
|      |             |              |             | DLF 144 2           | 0.01 ± 0.02        | 1 (50%)                              | 0                       | 1 (1)                    | N/A                       | N/A                         |               |
|      |             |              |             | LLF 144 2           | 0.01 ± 0.01        | 1 (50%)                              | 0                       | 1 (1)                    | N/A                       | N/A                         |               |
|      |             |              |             |                     |                     |                                      |                         |                          |                           |                             |               |
| Total|             |              |             | 7,725 2618          | 0.34 ± 0.26        | 2189 (84%)                           | 125                     | 2064 (306)               | 437.8 (395.7, 499.9)        | 99.3                        |               |

Notes.

\(^a\) Hosts were collected approximately 60 km east of Nome, in Council, AK (the nearest site that contains trees).

\(^b\) We initially sampled 144 leaf segments from each host species in each site (i.e., 48 per microsite for three microsites), but a small number were lost to contamination, overgrowth, and desiccation.

\(^c\) The diversity of Ascomycota sequences was estimated with ACE (abundance-based coverage estimator).
## Table 2  Summary of isolation frequency, richness and diversity of endophytic fungi, fungi from dead leaves in tree canopies (DLF), and fungi from leaf litter (LLF) in five North American sites.

| Site | Host species (yielding cultures) | Isolates | Isolation frequency ± SD | Isolates sequenced (%) | Basidiomycota sequences | Ascomycota sequences | Ascomycota putative species (95% CI) | Richness estimates (Bootstrap, ACE) | Fisher’s alpha (FA) | Mean FA/host species ± SD |
|------|----------------------------------|----------|--------------------------|------------------------|-------------------------|----------------------|--------------------------------------|----------------------------------|------------------|--------------------------|
|      |                                  |          |                          |                        |                         |                      |                                      |                                  |                  |                          |
|      | Endophyte                        | AZC 4 (3)| 52                       | 0.09 ± 0.07<sup>ab</sup> | 44 (85%)               | 0                     | 44                    | 15 (10.9, 19.1)                     | 17.2, 17.9 | 8                        | 3.5 ± 4.4       |
|      |                                  | NCH 4 (4)| 157                      | 0.28 ± 0.14<sup>ab</sup> | 148 (94%)              | 0                     | 148                   | 27 (25.1, 28.9)                     | 29.6, 29.3 | 9.7                      | 3.5 ± 1.9       |
|      |                                  | FLA 4 (4)| 335                      | 0.58 ± 0.26<sup>a</sup>  | 286 (85%)              | 104                   | 182                   | 60 (51.7, 68.3)                     | 71.7, 95.4 | 31.2                     | 12.3 ± 7.2      |
|      |                                  | AKE 3 (3)| 13                       | 0.03 ± 0.02<sup>b</sup>  | 11 (85%)               | 1                     | 10                    | 10 (4.5, 15.6)                      | 13.5, 55.0 | N/A                      | N/A             |
|      |                                  | AKN 3 (1)| 68                       | 0.16 ± 0.27<sup>ab</sup> | 65 (96%)               | 0                     | 65                    | 8 (5.5, 10.5)                       | 8.9, 10.0  | 2.4                      | 2.4             |
|      |                                  | Total 18 (15)| 625          | 0.24 ± 0.26<sup>ab</sup> | 554 (89%)              | 105                   | 449                   | 108 (99.6, 116.1)                    | 122.6, 128.5 | 45.1                     | 6.3 ± 6.2<sup>b</sup> |
|      | DLF                              | AZC 4 (4)| 186                      | 0.33 ± 0.19<sup>ab</sup> | 164 (88%)              | 0                     | 164                   | 33 (28.3, 37.7)                     | 37.8, 42.9 | 12.4                     | 4.0 ± 5.0<sup>b</sup> |
|      |                                  | NCH 4 (4)| 245                      | 0.43 ± 0.05<sup>ab</sup> | 215 (88%)              | 1                     | 214                   | 50 (42.1, 57.9)                     | 57.6, 67.1 | 20.5                     | 9.22 ± 9.4<sup>b</sup> |
|      |                                  | FLA 4 (4)| 366                      | 0.64 ± 0.06<sup>a</sup>  | 282 (77%)              | 6                     | 276                   | 81 (69.1, 92.9)                     | 95.9, 122.3 | 38.6                     | 22.9 ± 8.0<sup>a</sup>  |
|      |                                  | AKE 3 (2)| 68                       | 0.16 ± 0.12<sup>b</sup>  | 58 (85%)               | 0                     | 58                    | 25 (18.7, 31.3)                     | 30.1, 38.6 | 16.7                     | 11.8 ± 5.1<sup>b</sup> |
|      |                                  | AKN 3 (3)| 77                       | 0.18 ± 0.30<sup>b</sup>  | 61 (79%)               | 0                     | 61                    | 21(16.3, 25.7)                      | 25.1, 33.0 | 11.3                     | 10.5            |
|      |                                  | Total 18 (17)| 942          | 0.36 ± 0.23<sup>a</sup>  | 780 (83%)              | 7                     | 773                   | 182 (168.7, 193.3)                   | 211, 240.9 | 75.1                     | 11.9 ± 9.5<sup>a</sup> |
|      | LLF                              | AZC 4 (4)| 158                      | 0.28 ± 0.09<sup>ab</sup> | 144 (91%)              | 0                     | 144                   | 35 (29.7, 40.2)                     | 40.8, 46.4 | 14.7                     | 6.1 ± 5.2<sup>b</sup>  |
|      |                                  | NCH 4 (4)| 345                      | 0.60 ± 0.26<sup>a</sup>  | 262 (76%)              | 2                     | 260                   | 58 (49.8, 66.3)                     | 65, 67.0   | 23.2                     | 9.2 ± 2.3<sup>b</sup> |
|      |                                  | FLA 4 (4)| 377                      | 0.66 ± 0.10<sup>a</sup>  | 303 (80%)              | 5                     | 298                   | 93 (84.1, 101.9)                    | 109.3, 125.0 | 46.4                     | 30.4 ± 21.9<sup>b</sup> |
|      |                                  | AKE 3 (3)| 133                      | 0.31 ± 0.20<sup>ab</sup> | 113 (85%)              | 1                     | 112                   | 41 (33.3, 48.7)                     | 49.6, 70.7 | 23.3                     | 16 ± 4.2<sup>b</sup> |
|      |                                  | AKN 3 (3)| 38                       | 0.09 ± 0.15<sup>b</sup>  | 33 (90%)               | 5                     | 28                    | 12 (9.7, 14.3)                      | 14.1, 15.0 | 8                        | 6.9             |
|      |                                  | Total 18 (18)| 1051         | 0.41 ± 0.26<sup>a</sup>  | 853 (81%)              | 13                    | 842                   | 195 (179.1, 210.9)                   | 226.9, 266.3 | 79.6                     | 14.9 ± 14.2<sup>b</sup> |

**Notes.**

* Different letters represent significant differences ($P < 0.05$) in isolation frequency among sites for each leaf type (lowercase; endophytic: ANOVA $F_{4,10} = 7.31, P = 0.0051$; DLF: ANOVA $F_{4,12} = 3.30, P = 0.0482$; LLF: ANOVA $F_{4,13} = 7.41, P = 0.0025$) based on post-hoc Tukey’s HSD comparisons and among leaf types (uppercase: linear contrasts following ANOVA on residuals after test for site effects; $F_{1,47} = 4.11, P = 0.0484$).

** Different letters represent significant differences ($P < 0.05$) in diversity per-host-species among leaf types (uppercase; ANOVA on residuals following test for site effects, $F_{4,39} = 11.56, P < 0.0001$) and among sites for DLF and LLF (lowercase; DLF: ANOVA on log FA, $F_{3,19} = 5.35, P = 0.0162$; LLF: ANOVA on log FA, $F_{3,11} = 5.27, P = 0.0170$), but endophytic fungi only approached significance (ANOVA on log FA, $F_{2,8} = 3.87, P = 0.0667$).
ITS rDNA; ca. 600 bp) and an adjacent portion of the nuclear ribosomal large subunit (LSU rDNA; ca. 500 bp) were amplified by PCR as a single fragment using primers ITS1F or ITS5 and LR3 (White et al., 1990; Gardes & Bruns, 1993; Vilgalys & Hester, 1990; ITS rDNA-partial LSU rDNA) following U'Ren et al. (2010). ITS rDNA was amplified using primers ITS1 and ITS4 (White et al., 1990) for 19 isolates that failed to amplify using the primers listed above. PCR conditions are described in U'Ren et al. (2010).

PCR products were evaluated by staining with SYBR Green I (Molecular Probes, Invitrogen; Carlsbad, CA, USA) after electrophoresis on a 1% agarose gel. All positive amplicons yielded single bands. PCR products were cleaned, quantified, normalized, and sequenced directly with the Applied Biosystems BigDye® Terminator v3.1 cycle sequencing kit and the original PCR primers at the University of Arizona Genetics Core. Bidirectional sequencing was performed on an Applied Biosystems 3730xl DNA Analyzer (Foster City, CA, USA).

The software applications phred and phrap (Ewing & Green, 1998; Ewing et al., 1998) were used to call bases and assemble contigs with automation provided by the ChromaSeq package in Mesquite v. 1.06 (http://mesquiteproject.org). Base calls were verified by inspection of chromatograms in Sequencher v. 4.5 (Gene Codes, Ann Arbor, MI, USA). Overall, 1,037 new sequences were analyzed in conjunction with 1,030 sequences previously published as part of a study of endophytic and endolichenic fungal communities (U'Ren et al., 2010; U'Ren et al., 2012; U'Ren et al., 2016). Each sequence was queried against the non-redundant UNITE+INSDC database (v. 7.97.31.01, released 2016-01-31) using the Ribosomal Database Project (RDP) naïve Bayesian classifier (Wang et al., 2007) with a confidence threshold cutoff of 80% to estimate taxonomic placement. Basidiomycota comprised <6% of the overall data set and were excluded from all analyses due to their rarity, potentially reflecting use of a culture medium demonstrated to recover a high diversity of ascomycetous endophytes (Fröhlich & Hyde, 1999; Arnold, 2002). All sequences have been deposited in GenBank (Table S2).

Species richness and diversity
Operational taxonomic units (OTU) were defined using ITS-partial LSU rDNA sequences. To designate OTU a distance matrix based on pairwise Needleman-Wunsch alignments was generated with the needledist module in ESPRIT (Sun et al., 2009) under the default parameters. Gaps of any length were treated as a single evolutionary event and terminal gaps were not penalized (Sun et al., 2009). We used mothur (Schloss et al., 2009) to cluster sequences into OTU using the complete-linkage method (i.e., furthest neighbor) and to calculate richness and OTU overlap among communities. Sequences were assembled into OTU based on sequence similarities ranging from 95% to 100% following U'Ren et al. (2012). Results are presented at the genotype level (i.e., 100% sequence similarity) and at the putative species level (i.e., 95% sequence similarity; see U'Ren et al., 2009). Previous assessment of four endophyte-rich genera in the Sordariomycetes and Dothideomycetes demonstrated that ca. 5% ITS rDNA divergence (i.e., 95% sequence similarity) conservatively estimates sister species boundaries when compared against published phylogenies (U'Ren et al., 2009; see also Liggenstoffer et al., 2010). OTU richness...
increased when more stringent sequence similarity levels were used (e.g., 348 OTU were estimated at 97% sequence similarity vs. 306 OTU at 95% sequence similarity), generally reflecting an increase in singletons that could not be used for comparative analyses. Even so, estimates of overlap among different leaf types were congruent based on 95–100% sequence similarity groups.

OTU accumulation curves (Fig. 1; Fig. S1), rarefaction analyses, estimates of total richness (bootstrap and abundance-based coverage estimator (ACE), recommended by Hortal, Borges & Gaspar, 2006), and diversity were inferred in EstimateS v. 8.0 (http://viceroy.eeb.uconn.edu/EstimateS) using 50 randomizations of sample order without replacement. Fisher’s alpha, a parameter of the log series model that is theoretically independent of sample size (Fisher, Corbet & Williams, 1943; Taylor, 1978; Magurran, 2004), was chosen to measure fungal diversity. Fisher’s alpha is robust for comparing diversity among communities that are sampled unequally (given variation in isolation frequency) and are characterized by a log-series distribution of OTU abundance (Magurran, 2004).

We analyzed isolation frequency and diversity as a function of site, host family, and leaf type (i.e., living, senesced, or leaf litter) using multiple regression with these explanatory factors as main effects. Isolation frequency and diversity were logit and log transformed, respectively, to achieve normality. The fit of each model was assessed using a lack-of-fit F test. Pairwise comparisons between fungal communities from different leaf types were computed using least-squares means contrasts. Analyses were done in JMP v. 9.0.0 (SAS Institute, Cary, NC).

**Similarity of fungal assemblages**

We used several approaches to examine the relationship of site, host, and leaf type to the composition of fungal communities. First, we used analysis of similarity (ANOSIM) coupled with visualization by non-metric multidimensional scaling (NMDS) to assess whether fungal communities differed significantly among sites (Fig. S2). Because fungal communities did differ among sites, NMDS ordinations also were computed within each site to examine the effect of host species and leaf type (Fig. S2). ANOSIM and NMDS were conducted in PAST v. 1.88 (Hammer, Harper & Ryan, 2001) following U’Ren et al. (2012).

Second, these analyses were complemented by hierarchical clustering to clarify the relationships among fungal communities in different sites and leaf types. The dendrogram was generated using the unweighted pair-group average (UPGMA) algorithm. Hierarchical clustering was conducted in PAST v. 1.88 (Hammer, Harper & Ryan, 2001) using 10,000 bootstrap replicates.

Third, permutational multivariate analysis of variance (PERMANOVA) was used to examine variation in community composition as a function of (1) leaf type and host species, while constraining permutations with each site; and (2) leaf type, while constraining permutations with each site and each host species. PERMANOVA was implemented with the adonis and permute functions in the R (R Core Team) package vegan (Oksanen et al., 2015). ANOSIM, NMDS, hierarchical clustering, and PERMANOVA were conducted using all non-singleton OTU (i.e., those occurring more than once in the entire
Figure 1  Species accumulation curves (Mao Tau), 95% confidence intervals, and bootstrap estimates of richness based on ITS-partial LSU rDNA. (A) 449 isolates of endophytic fungi; (B) 773 isolates of fungi from senesced leaves in plant canopies (dead leaf fungi, DLF); and (C) 842 isolates of fungi from leaf litter (leaf litter fungi, LLF) in five sites.
dataset) to minimize the potential under-estimation of similarity due to undersampling (Manter & Bakker, 2015). The Morisita-Horn metric of similarity was used to minimize sensitivity to under-sampling, as it reflects the distribution of abundant taxa (Beck, Holloway & Schwanghart, 2013). Host individuals/leaf types from which <4 isolates were sequenced were excluded from all analyses (see Table 1).

Finally, because sample sizes generally were low for each host species-leaf type combination, and because distributions of rare taxa can be informative, we examined community structure for all OTU grouped by site and leaf type using OTUshuff (Manter & Bakker, 2015). To estimate the influence of rare and abundant taxa on community similarity, OTUshuff uses a variably weighted version of the Odum index (wOdum). At $\alpha = 1$, $D_{wOdum}$ is equal to $D_{Odum}$; when $\alpha < 1$, the influence of abundant taxa will be down-weighted; and when $\alpha > 1$, the influence of abundant abundance taxa will be up-weighted. We varied the alpha weighting factor from 0 to 4 to test the null hypothesis that samples from different leaf types within a site represent the same community (Table S1). Analyses were conducted using a Monte Carlo simulation with 1,000 iterations. Low isolation frequency of endophytes prevented comparisons of endophytes and DLF/LLF in one site (AKE).

**Functional characterization of fungal isolates**

We compared *in vitro* growth and cellulolytic, pectinolytic, and ligninolytic activity for representative isolates of eight OTU found only in living leaves and nine OTU found only in non-living leaves. Each OTU was represented by a single isolate. Leaf type designation for each OTU (i.e., living leaves vs. non-living leaves) was based on host and substrate information from a larger collection of endophytic and endolichenic fungi (see U'Ren et al., 2012), as well as BLASTn comparisons to the NCBI nr database. When possible, OTU representing each leaf status (living, non-living) were chosen from the same class (i.e., Sordariomycetes and Dothideomycetes). However, paired comparisons were not possible for Pezizomycetes or Leotiomycetes because no OTU in those classes were found only as endophytes.

Pure cultures were started on 2% MEA plates and were allowed to grow for up to 7 d. At that point 5 mm agar plugs containing actively growing mycelium were placed on water agar plates (WA; pH 5.6, 1.5% agar, containing 50 ml 20X nitrate salts, 1 ml 1000X trace elements, and 1 ml vitamin solution per liter; see Tucker & Orbach, 2007) amended with (1) 5 g carboxymethylcellulose (cellulose substitute; Sigma-Aldrich, St. Louis, Missouri, USA); (2) 10 g citrus pectin (grade 1 from citrus fruits, Sigma-Aldrich); or (3) 0.5 g indulin (lignin substitute; Mead-Westvaco, Richmond, Virginia, USA) as the sole carbon source. Three replicates were prepared for each isolate per medium. A culture with verified enzymatic activity was used as the positive control.

Cultures were grown for 5–28 days in a temperature controlled room at 26 °C under constant light conditions. Once cultures were ca. 3–4 cm in diameter, plates were stained with aqueous solutions of 0.2 % (wt/vol) Congo red, equal parts of 1.0 % (wt/vol) FeCl$_3$ and K$_3$[Fe(CN)]$_6$, or 0.05% (wt/vol) ruthenium red to detect cellulolytic, pectinolytic, and ligninolytic activity, respectively (modified from Gazis et al., 2012; Cotty et al., 1990).
Following staining, plates were rinsed with either 1M sodium chloride (for cellulose plates) or deionized water (pectin and lignin plates) and the colony diameter and clearing at the periphery of mycelial growth (i.e., extra-hyphal clearing) of each isolate was measured.

Average growth rates (colony diameter/days of growth) were compared on each carbon source as a function of fungal taxonomy at the class level and leaf type using non-parametric tests. The proportion of isolates that demonstrated diagnostic clearing was compared as a function of leaf type using two-tailed Fisher’s exact tests. Where present, the amount of clearing at the periphery of mycelial growth was scaled by growth rate and compared using Wilcoxon rank sum tests.

Mantel tests were used to examine the relationship of fungal genetic distance to patterns of cellulolytic activity and growth rates on different carbon sources. ITS-partial LSU rDNA sequences for Sordariomycetes and Dothideomycetes isolates were aligned using MAFFT v6.821b (Katoh & Toh, 2008) with the L-INS-I setting for high accuracy. Ambiguously aligned nucleotides were masked using the Gblocks server v.0.91b (Castresana, 2000) with parameters to allow for less stringent selection (i.e., to allow for smaller final blocks; to allow gap positions within the final blocks; and to allow less strict flanking positions) resulting in retention of 83% of the original 1,223 positions. A distance matrix was calculated in PAUP* v4.0 (Swofford, 2003) using uncorrected distances (p). Pairwise distance matrices for (1) cellulolytic activity; (2) average growth rate on cellulose; (3) average growth rate on pectin; and (4) average growth rate on lignin were calculated in JMP v. 9.0.0. Mantel tests were performed in R with the ade4 package (Dray & Dufour, 2007) using 9,999 permutations.

RESULTS

We obtained 2,618 fungal isolates from the interior of 7,725 leaf segments representing living leaves (i.e., endophytes), dead leaves in plant canopies (dead leaf fungi, DLF), and leaf litter (leaf litter fungi, LLF; Table 1) of 18 host species. ITS-partial LSU rDNA data obtained from 2,064 isolates of Ascomycota comprised 555 unique genotypes (100% sequence similarity) and 306 putative species (i.e., OTU at 95% sequence similarity) (Table 1).

Our sampling effort did not capture the estimated richness of cultivable leaf-associated fungi for the entire study (Fig. 1; Table 2). However, sampling in eight of 15 site-leaf type combinations encompassed the local richness of fungi based on bootstrap estimates (Table 2), and average differences between bootstrap and ACE estimates and the upper 95% confidence intervals around observed richness were only 3.0 and 13.7 OTU, respectively (Table 2; see also Fig. S1). Therefore, we used these data to compare communities of cultivable fungi as described below.

Isolation frequency and diversity

Isolation frequency and diversity differed as a function of site, host family, and leaf type (Table 3). Across all sites, fungi were isolated less frequently from living leaves than from dead leaves in plant canopies or leaf litter (Table 3). Endophytes were isolated from 24% ± 26% (mean ± SD) of leaf segments per host species (range: 0–96%; Table 2). DLF and LLF were isolated from 36% ± 23% (range, 0–71%) and 41% ± 26% (range, 1–85%) of
leaf segments per host species, respectively (Table 1). A greater number of hosts failed to yield cultures from living leaves (i.e., *Salix pulchra* and *Betula nana* at AKN and *Quercus rugosa* at AZC; Table 2) compared to a single host for non-living leaves (*Betula nana* at AKN; only LLF were obtained; Table 1).

Endophytes also were less diverse than DLF and LLF (Table 3). Cumulative diversity of fungi from non-living leaves over the entire study was 1.7 to 1.8 times that of fungi from living leaves (Table 2).

**Fungal community analyses**

Fungal communities differed among sites (Fig. 2; Fig. S2A). Within sites, communities differed among host species and leaf type (Fig. 2; Table 4; Fig. S2B–S2D).

Within each site, we rejected the null hypothesis that endophytes and DLF/LLF were from the same community at all levels of alpha ($P < 0.05$; Table S1). Preferentially weighting rare species decreased similarity between endophytes and DLF, and between DLF and LLF, relative to analyses in which abundant species were weighted preferentially. In contrast, variably weighting rare or abundant OTU had no discernible effect on similarity estimates of endophytes and LLF, which were consistently low (Fig. 3; Table S1).

Overall, 29% of non-singleton OTU (Fig. S3) and 52% of common OTU (those represented by >5 sequenced isolates) were found in both living and non-living leaves (Fig. 4). After removing infrequent taxa, OTU from only one leaf status were rare: only 3%, 5%, or 8% of OTU were unique to living leaves, dead leaves in plant canopies, or leaf litter, respectively (Fig. 4; see also Fig. S4). However, OTU that were highly abundant in leaf litter were seldom found as endophytes (Fig. 4), and the abundance of OTU from leaf litter was not correlated with abundance of the same OTU in living leaves (Table S3).

**Substrate utilization**

We compared *in vitro* growth for representative isolates of eight OTU found only in living leaves and nine OTU found only in non-living leaves. All isolates, regardless of leaf type of origin, grew on cellulose, lignin, and pectin as sole carbon sources.
Figure 2  Order-level taxonomy and hierarchical clustering analysis of fungal communities in living (E), senesced (D), and fallen (L) leaves among sites. (A) Proportion of isolates representing orders of Pezizomycotina as a function of site/leaf type. Sequences from AKE (Eagle Summit, Alaska) and AKN (Nome, Alaska) were pooled for this analysis (AK). (B) Unweighted pair-group average (UPGMA) dendrogram showing hierarchical clustering of endophytes (Endo), DLF, and LLF from each site (AZC, Chiricahua Mountains, Arizona; NCH, Highlands Biological Station, North Carolina; FLA, Archbold Biological Station, Florida; AK, see above). Bootstrap values are based on 10,000 bootstrap replicates. Host individual/leaf type with <4 sequences and singleton OTU were excluded from hierarchical clustering analysis.
Table 4  Results of PERMANOVA analysis of the Morisita-Horn dissimilarities for fungal OTU community structure (DF, degrees of freedom; SS, sum of squares; MS, mean sum of square; F, pseudo-F by permutation).

| Blocks: site (1000 permutations) | DF | SS   | MS   | F     | R²  | P     |
|---------------------------------|----|------|------|-------|-----|-------|
| Host species                    | 11 | 17.978 | 1.634 | 5.546 | 0.431 | 0.001 |
| Leaf type                       | 2  | 1.372 | 0.686 | 2.328 | 0.033 | 0.001 |
| Residuals                       | 78 | 22.398 | 0.295 | 0.537 |      |       |
| Total                           | 89 | 41.748 |   | 1.000 |      |       |

| Blocks: site; plots: host species (200 permutations) | DF | SS   | MS   | F     | R²  | P     |
|-----------------------------------------------------|----|------|------|-------|-----|-------|
| Leaf type                                           | 2  | 1.446 | 0.723 | 1.560 | 0.035 | 0.005 |
| Residuals                                           | 87 | 40.303 | 0.463 |       | 0.965 |       |
| Total                                               | 89 | 41.748 |   | 1.000 |      |       |

We focused on OTU in two classes to evaluate the relationship of substrate utilization to leaf type of origin. Dothideomycetes from living leaves grew more rapidly than those isolated originally from non-living leaves on all three carbon sources (Fig. 5C). However, growth of Sordariomycetes on cellulose, lignin, or pectin did not differ for fungi from living vs. non-living tissues (Fig. 5C).

We found no significant correlation between genetic distance and differences in growth on cellulose or lignin, but differences in fungal growth on pectin were correlated with genetic distance between isolates (Table S3).

**Enzyme activity**

Extra-hyphal clearing (indicative of enzyme activity) was observed for isolates only on media containing cellulose as the sole carbon source. Only the positive control displayed in vitro ligninolytic or pectinolytic activity.

The number of OTU with detectable cellulolytic activity vs. no cellulolytic activity did not differ as a function of leaf type of origin (Table S3). For OTU with detectable clearing, the degree of cellulolytic activity differed significantly among fungi from different classes, with Leotiomycetes from non-living leaves displaying the greatest enzymatic activity (Table 5; Fig. 5A). Dothideomycetes from non-living leaves had greater cellulolytic activity compared to those isolated only from living leaves (Fig. 5B). Sordariomycetes from different leaf types did not differ significantly in cellulolytic activity (Fig. 5B). We found no significant correlation between genetic distance and differences in cellulolytic activity (Table S3).

**DISCUSSION**

The occurrence of some endophytic fungi in dead leaves and/or leaf litter and the ability of some endophytes to produce extracellular leaf-degrading enzymes in a manner consistent with saprotrophs (Carroll & Petrini, 1983; Fröhlich & Hyde, 1999; Promputtha et al., 2007; Korkama-Rajala, Müller & Pennanen, 2008; Promputtha et al., 2010; U'Ren et al., 2010; Chaverri & Gazis, 2011; Osono & Hirose, 2011; Sun, Guo & Hyde, 2011; He et al., 2012) has
Figure 3  Pairwise community similarity among different leaf types as a function of site. Similarity is measured as $1-D_{\text{wOdum}}$, as calculated in OTUshuff using 1,000 Monte Carlo iterations. Similarity was assessed at different alpha weighting factors (i.e., when $\alpha = 1$, $D_{\text{wOdum}}$ is equal to $D_{\text{Odum}}$; when $\alpha < 1$, the influence of abundant taxa will be down-weighted; and when $\alpha > 1$, the influence of low abundance taxa will be down-weighted). Low sample size of endophytes precluded comparisons of endophyte and DLF/LLF at AKE.
Figure 4  Heat map showing the distribution of 104 putative species (based on 95% rDNA sequence similarity; OTU with <5 isolates were excluded) of endophytic, dead leaf fungi (DLF), and leaf litter fungi (LLF) from plants in five North American sites. Species abundance on each leaf type is shown as the percentage of the total number of isolates of that species. Taxonomy was estimated for each species using the Ribosomal Database Project naïve Bayesian classifier (continued on next page...)
led to suggestions that many endophytes are incidental symbionts that exist in leaves as a prelude to completing their life cycles as primary decomposers. Here, we explore the endophyte–saprotroph continuum, with the goal of understanding the degree to which endophytes may occur in non-living tissues in diverse plants and terrestrial biomes and whether ecological patterns reflect fungal carbon substrate utilization and enzyme activity.

**Fungal communities differ in living and non-living leaves**

As a whole, culturable fungi inhabiting living leaves were isolated less frequently and were less diverse than those in non-living leaves. This may reflect the plant actively restricting fungal colonization within leaves, limited growth by fungi within living leaves for intrinsic reasons, or culturing biases (see below). Culturable fungal communities in living leaves also differed in composition from communities in senesced leaves and leaf litter, regardless of differential weighting of rare and abundant taxa.

These observations are congruent with colonization of senesced and abscised leaves by additional epiphytic- and/or litter fungi (see Lindahl et al., 2007). As leaves senesce, epiphytic fungi can colonize the interior of leaves rapidly, leading to a subsequent decline in the relative abundance of endophytic taxa (Cabral, 1985; Stone, 1987; Osono, 2002; but see Peršoh et al., 2013). When we examine common OTU as a function of their abundance in leaves of each type, 35% have patterns of leaf-association that match that of ambient, non-endophytic fungi colonizing leaf interiors after leaf death (Fig. 4). Additional fungi appear to colonize fallen leaves from the underlying litter layer or from other sources; these represented a relatively small number (8% of OTU), potentially reflecting preferential growth of Ascomycota (vs. Basidiomycota and other fungal taxa) on the culture medium used here.

Although culturable ascomycete communities as a whole differed significantly as a function of leaf type, 29% of non-singleton OTU (Fig. S3) and 52% of common OTU (OTU represented by >5 sequences) from living leaves also inhabited non-living leaves. These OTU typically were present in living vs. non-living leaves at different abundances (Fig. 4). This is consistent with previous studies that noted the transitory nature of many endophytes in leaf litter (Stone, 1987; Hudson, 1968; Osono & Takeda, 2001), as well as the quick turnover in litter of all phyllosphere fungi (including both epiphytes and endophytes). For example, Voříšková & Baldrian (2013) found that the majority of phyllosphere OTU were absent from leaves two to four months after abscission. In contrast, a greater proportion of Xylariaceae appear to inhabit living leaves, as well as decomposing leaves, wood, and fruit (U'Ren et al., 2016). Factors that mediate these shifts merit further exploration, including the exhaustion of readily available sugars, the ability of some fungi to decompose structural polymers, and processes relevant to competitive exclusion or antibiosis (see Osono, 2006; Yuan et al., 2011; Yuan & Chen, 2014).
Figure 5  Normalized in vivo cellulolytic activity (mean activity/colony diameter) and growth (mean colony diameter/day) of 17 representative fungal OTU isolated from living (i.e., endophyte) and non-living leaves (DLF/LLF). (A) Mean cellulolytic activity as a function of class-level taxonomy and (B) class-level taxonomy and leaf type. (C) Mean growth rate as a function of both class-level taxonomy (S, Sordariomycetes; D, Dothideomycetes) and carbon source (continued on next page...).
Figure 5 (...continued)

(cellulose, lignin, or pectin). Different letters represent significant differences in enzyme activity as a function of fungal class after *post-hoc* nonparametric multiple comparisons (A). Asterisks (**) indicate significant differences in cellulytic activity or growth between fungal OTU found only in non-living leaves vs. fungal OTU isolated from only living leaves (*P* < 0.05; Table S3) (B and C). No significant differences were observed between OTU of Sordariomycetes from different leaf types (Table S3) (B and C).

Differences in fungal communities as a function of leaf type are consistent among sites and host lineages

Although the same major lineages of Ascomycota were present in all sites, fungal communities differed among sites in terms of the relative abundance of those lineages, as well as at finer taxonomic levels. In some cases, these differences may reflect unequal recovery of fungi from different plant lineages: in AKN and AZC, for example, fungi were more commonly isolated from coniferous hosts, and isolation frequency from angiosperms was low compared to other sites. Yet, communities still differed between sites when lineages were sampled equally (e.g., NCH and FLA), consistent with our previous results on endophytic and endolichenic fungi from the same locations (U’Ren *et al.*, 2012) and with other studies illustrating the influence of climate and biogeographic factors on endophyte community composition (e.g., Davis & Shaw, 2008; Zimmerman & Vitousek, 2012; Langenfeld *et al.*, 2013).

Within each site we found that fungal community composition differed markedly among host species. These results are consistent with the view that plant defenses may mediate infection and colonization by particular endophytes (see Schulz & Boyle, 2005), with persistent signatures of host chemistry or other traits contributing to the structure of fungal assemblages in non-living leaves. Previous work has shown that litter traits (i.e., nutrient quality or other traits relevant to host species) may have stronger effects on fungal community composition than other environmental factors (Aneja *et al.*, 2006; Šnajdr *et al.*, 2011; Bray, Kitajima & Mack, 2012; Urbanová, Šnajdr & Baldrian, 2015). More generally, the consistent differences observed between fungal communities in living vs. non-living leaves in different hosts and sites suggest that factors influencing the ecology and evolution of plant-fungal associations may be relatively consistent at large scales.

Evaluating the strength of our conclusions

Differences in fungal communities observed here could result from artifacts of insufficient sampling, unintentional comparisons among leaves of different ages (e.g., leaf litter representing previous years of growth vs. living leaves representing the current year), and/or biasing our work toward fungi that are readily cultured on standard nutrient media or specific isolation techniques (see Unterseher & Schnittler, 2009 for comparison of fragment plating vs. dilution-to-extinction culturing). The last issue might be especially problematic for fungi that do not normally occur in symbiosis and/or have the capacity to grow on non-living substrates vs. those with a symbiotrophic lifestyle. However, several lines of evidence suggest our conclusions are robust. First, species accumulation curves and bootstrap estimates of species richness for all sites and site/leaf-type combinations indicate our sampling was at or near statistical completion for culturable fungi. Second,
| Isolate Name | 95% ITS-partial LSU OTU | Substrate type | Isolates in OTU\(^a\) Sites\(^a\) Plant host species (families)\(^b\) | Taxonomy\(^b\) | Mean ± SD growth rate on cellulose medium (mm/day) | Mean ± SD growth rate (mm/day) on lignin medium | Mean ± SD growth rate (mm/day) on pectin medium | Mean ± SD cellulolytic activity/colony diameter (mm) |
|--------------|------------------------|---------------|---------------------------------------------|-----------------|-----------------------------------------------|-----------------------------------------------|-----------------------------------------------|-----------------------------------------------|
| NC0075       | 403                    | Living        | 26                                          | 1 2 (2) | Dothideomycetes (*Phyllosticta*)              | 10.7 ± 1.1                                    | 7.2 ± 1.0                                     | 25.0 ± 3.1                                    | 22.0 ± 6.6                                     |
| NC0101       | 404                    | Living        | 6                                           | 1 2 (2) | Dothideomycetes (*Phyllosticta*)              | 11.4 ± 2.0                                    | 7.8 ± 0.4                                     | 22.1 ± 3.8                                    | 40.0 ± 16.8                                    |
| FL0319       | 261                    | Living        | 7                                           | 1 1 (1) | Dothideomycetes (*Ochrocladosporium*)         | 17.1 ± 0.3                                    | 10.4 ± 0.3                                    | 14.5 ± 0.4                                    | 45.3 ± 1.5                                     |
| FL1985       | 366                    | Non-living    | 13                                          | 1 2 (2) | Dothideomycetes (*Penidiella*)                | 7.3 ± 0.3                                     | 6.2 ± 0.5                                     | 9.6 ± 0.6                                     | 49.0 ± 6.0                                     |
| FL0303       | 259                    | Living        | 12                                          | 1 1 (1) | Dothideomycetes (*Mycosphaerella*)            | 26.7 ± 0.4                                    | 26.9 ± 1.6                                    | 28.3 ± 0.8                                    | 15.3 ± 4.0                                     |
| FL1704       | 353                    | Non-living    | 9                                           | 1 2 (1) | Dothideomycetes (*Teratosphaeria marasasii*)  | 10.2 ± 5.8                                    | 6.6 ± 0.4                                     | 8.5 ± 0.1                                     | 82.0 ± 42.5                                    |
| AK1907       | 118                    | Non-living    | 16                                          | 3 4 (1) | Leotiomycetes (*Hyellodendriella*)            | 11.3 ± 0.2                                    | 11.7 ± 0.9                                    | 12.6 ± 0.3                                    | 46.0 ± 2.0                                     |
| FL2076       | 360                    | Non-living    | 9                                           | 1 2 (2) | Leotiomycetes (*Rhytisma*)                    | 6.2 ± 1.4                                     | 6.8 ± 0.5                                     | 6.2 ± 1.6                                     | 145.7 ± 38.8                                   |
| FL2145       | 355                    | Non-living    | 7                                           | 1 1 (1) | Leotiomycetes (*Rhytisma*)                    | 7.0 ± 0.8                                     | 4.1 ± 1.3                                     | 23.1 ± 1.2                                    | 167.7 ± 28.9                                   |
| AZ0245       | 163                    | Non-living    | 7                                           | 1 2 (2) | Pezizomycetes (*Strobiliscypha*)              | 13.2 ± 0.5                                    | 5.9 ± 0.4                                     | 31.3 ± 3.3                                    | 40.3 ± 4.6                                     |
| FL0231       | 253                    | Living        | 7                                           | 2 5 (2) | Sordariomycetes (*Lecythophora*)              | 20.5 ± 0.6                                    | 25.9 ± 1.1                                    | 22.8 ± 1.0                                    | 0.0 ± 0.0                                      |
| NC0063       | 406                    | Living        | 7                                           | 1 4 (3) | Sordariomycetes (*Eutypa*)                   | 25.2 ± 1.3                                    | 21.7 ± 0.6                                    | 21.0 ± 0.3                                    | 28.3 ± 4.5                                     |
| NC0012       | 399                    | Living        | 12                                          | 1 1 (1) | Sordariomycetes (*Tsukuba seocharakensis*)    | 9.0 ± 0.2                                     | 20.3 ± 1.2                                    | 105.0 ± 4.4                                   | 116.7 ± 4.9                                    |
| FL2151       | 352                    | Non-living    | 31                                          | 1 4 (3) | Sordariomycetes (*Magnaporthe*)               | 24.4 ± 0.3                                    | 20.2 ± 0.2                                    | 21.2 ± 0.5                                    | 0.0 ± 0.0                                      |
| NC1320       | 378                    | Non-living    | 6                                           | 2 2 (2) | Sordariomycetes (*Fusarium*)                  | 81.0 ± 4.4                                    | 55.2 ± 2.9                                    | 91.0 ± 1.7                                    | 3.0 ± 0.0                                      |
| FL1642       | 170                    | Non-living    | 13                                          | 3 3 (1) | Sordariomycetes (*Polyscytalum*)              | 17.8 ± 1.3                                    | 6.9 ± 1.1                                     | 21.9 ± 2.4                                    | 39.0 ± 5.6                                     |
| NC0068       | 263                    | Living        | 13                                          | 2 2 (2) | Sordariomycetes (*Collectotrichum*)           | 86.3 ± 0.6                                    | 66.7 ± 0.8                                    | 91.3 ± 0.6                                    | 0.0 ± 0.0                                      |

**Notes.**

- **a** Totals include fungi isolated from lichens and plants as part of a larger study on endophytic and endolichenic fungi (U'Ren et al., 2012).
- **b** Taxonomic information is based on queries of UNITE database using the RDP Bayesian classifier (see methods). However, when matches to the UNITE database lacked genus and species information (e.g., the species hypothesis for FL1985 is Capnodiaceae sp.), taxonomic information is estimated based on a BLASTn query of the NCBI nr database (indicated with an *).
although leaves may have emerged at different times and thus been subject to different pools of inoculum (see Arnold & Herre, 2003), patterns are consistent across hosts from different microsites, geographic locations, and host species, including both deciduous and evergreen angiosperms as well as conifers. Third, although missing species found using culture-independent methods have the potential to alter our estimates of diversity and the degree of overlap among leaf-types (Arnold et al., 2007; Gallery, Dalling & Arnold, 2007; Jumpponen & Jones, 2009; U’Ren et al., 2014; Balint et al., 2015), the consistent patterns we observed across hosts and sites suggest that our conclusions are valid for prevalent members of these fungal communities. In addition, our results are congruent with similar studies that used culture-free methods (Voříšková & Baldrian, 2013).

The abundance and identity of endophytes in leaf litter typically differs as a function of leaf age and season (Hirose & Osono, 2006; Osono, 2006; Peršoh et al., 2013). Had we sampled freshly fallen leaves, communities may have been more similar among leaf-types; conversely, greater differences may have been observed had we examined more thoroughly decomposed leaves (see Voříšková & Baldrian, 2013). Accordingly, our estimates of overlap among endophytic and saprotrophic communities are restricted to culturable members from leaf litter in a moderate state of decay sampled during the summer. However, the relatively consistent patterns observed from tundra to subtropical communities suggest strong signal in the data.

Importantly, the relative abundance of isolates in living or dead tissues is not necessarily indicative of an isolate’s ability to decompose plant tissues. This prompted our evaluation of substrate use and enzyme activity (below). Future studies that assess endophyte and litter communities over multiple time points (including multiple time points for living leaves and during the first weeks of decomposition) in combination with functional assays and metatranscriptomic methods, will provide much-needed insight into the persistence and functional roles of endophytes as decomposers.

**Functional differences**
The limited persistence of endophytes in leaf litter over time led us to conduct a preliminary investigation of *in vitro* patterns of substrate use between OTU found only in living tissues vs. only in non-living tissues in our surveys (including the larger sampling from U’Ren et al., 2012). Across four classes of Pezizomycotina, cellulolytic activity differed primarily as a function of class-level taxonomy. However, cellulolytic activity was not correlated with finer-scale genetic relationships within Sordariomycetes and Dothideomycetes. Although some isolates demonstrated high levels of cellulolytic activity (e.g., Leotiomycetes representing the Rhytismataceae and Helotiaceae), our study differs from previous ones in that none of our isolates had detectable ligninolytic or pectinolytic activity (see Oono et al., 2014; Fouda et al., 2015). We detected differences in cellulolytic activity and growth rate on pectin-, lignin-, and cellulose media between Dothideomycetes fungi from living vs. non-living leaves. However, such differences were not observed in the Sordariomycetes. Overall, more precise measurements investigating a broader array of enzymes (e.g., β-mannanase, xylanase, laccase, etc.), in conjunction with genomic and transcriptomic studies of multiple
isolates per OTU, are necessary to identify functional differences among fungi associated with leaves at different life stages.

Conclusions
Approximately half of the common OTU found here occurred in both living and non-living leaves. However, our sampling across diverse biomes and host lineages detected consistent differences in living vs. non-living leaves in terms of diversity and fungal community composition. Fungi that were common in decomposing leaves collected from leaf litter were not typically common in living leaves of the same plants. Thus some fungi with endophytic life stages clearly persist for periods of time in leaves after senescence and incorporation into leaf litter, with shifts after leaf death and in the early stages of decomposition likely reflecting incursion by additional fungi from the leaf exterior. Overall, our data—while preliminary in some regards—suggest that focal isolates found in dead and fallen leaves did not differ consistently from those in living leaves in terms of their capacity to use pectin, lignin, and cellulose as their sole carbon sources, nor in the degree of cellulolytic activity displayed in vitro. We suggest that culturable fungi with endophytic life stages may not persist in decomposing leaves once cellulose and hemicellulose are depleted. Future analyses should optimize methods to culture both Basidiomycota and Ascomycota endophytes, and should explore more dimensions of functional traits and persistence to further define the endophytism-to-saprotrophy continuum.

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The authors declare there are no competing interests.

Author Contributions
• Jana M. U’Ren conceived and designed the experiments, performed the experiments,
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paper.
• A. Elizabeth Arnold conceived and designed the experiments, contributed
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DNA Deposition
The following information was supplied regarding the deposition of DNA sequences:
GenBank accession numbers are provided for all sequences in Table S2.

Data Availability
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REFERENCES
Aneja MK, Sharma S, Fleischmann F, Stich S, Heller W, Bahnweg G, Munch JC,
Schloter M. 2006. Microbial colonization of beech and spruce litter— influence of
decomposition site and plant litter species on the diversity of microbial community.
Microbial Ecology 52:127 DOI 10.1007/s00248-006-9006-3.
Arnold AE. 2002. Fungal endophytes in Neotropical trees: abundance, diversity and
ecological interactions. Doctoral dissertation, University of Arizona, Tucson, 337p.
Arnold AE, Henk DA, Eells RL, Lutzoni F, Vilgalys R. 2007. Diversity and phylogenetic affinities of foliar fungal endophytes in loblolly pine inferred by culturing and environmental PCR. *Mycologia* 99:185–206 DOI 10.3852/mycologia.99.2.185.

Arnold AE, Herre EA. 2003. Canopy cover and leaf age affect colonization by tropical fungal endophytes: ecological pattern and process in *Theobroma cacao* (Malvaceae). *Mycologia* 95:388–398 DOI 10.2307/3761880.

Arnold AE, Lamit LJ, Bidartondo M, Gehring C, Callahan HS. 2010. Interwoven branches of the plant and fungal trees of life. *New Phytologist* 185:874–878 DOI 10.1111/j.1469-8137.2010.03185.x.

Arnold AE, Lutzoni F. 2007. Diversity and host range of foliar fungal endophytes: are tropical leaves biodiversity hotspots? *Ecology* 88:541–549 DOI 10.1890/05-1459.

Arnold AE, Miadlikowska J, Higgins KL, Sarvate SD, Gugger P, Way A, Hofstetter V, Kauff F, Lutzoni F. 2009. A phylogenetic estimation of trophic transition networks for ascomycetous fungi: are lichens cradles of symbiotrophic fungal diversification? *Systematic Biology* 58:283–297 DOI 10.1093/sysbio/syp001.

Balint M, Bartha L, O’Hara RB, Olson MS, Otte J, Pfenninger M, Robertson AL, Tiffin P, Schmitt I. 2015. Relocation, high-latitude warming and host genetic identity shape the foliar fungal microbiome of poplars. *Molecular Ecology* 24:235–248 DOI 10.1111/mec.13018.

Barton AM. 1994. Gradient analysis of relationships among fire, environment, and vegetation in a southwestern USA mountain range. *Bulletin of the Torrey Botanical Club* 121:251–265 DOI 10.2307/2997180.

Beck J, Holloway JD, Schwanghart W. 2013. Undersampling and the measurement of beta diversity. *Methods in Ecology and Evolution* 4:370–382 DOI 10.1111/2041-210x.12023.

Bray SR, Kitajima K, Mack MC. 2012. Temporal dynamics of microbial communities on decomposing leaf litter of 10 plant species in relation to decomposition rate. *Soil Biology & Biochemistry* 49:30–37 DOI 10.1016/j.soilbio.2012.02.009.

Cabraal D. 1985. Phyllosphere of *Eucalyptus viminalis*: dynamics of fungal populations. *Transactions of the British Mycological Society* 85:501–511 DOI 10.1016/S0007-1536(85)80047-4.

Carroll GC, Petrini O. 1983. Patterns of substrate utilization by some fungal endophytes from coniferous foliage. *Mycologia* 75:53–63 DOI 10.2307/3792923.

Castresana J. 2000. Selection of conserved blocks from multiple alignments for their use in phylogenetic analysis. *Molecular Biology and Evolution* 17:540–552 DOI 10.1093/oxfordjournals.molbev.a026334.

Chaverri P, Gazis RO. 2011. Linking *ex planta* fungi with their endophytic stages: *Perisporiopsis*, a common leaf litter and soil fungus, is a frequent endophyte of *Hevea* spp. and other plants. *Fungal Ecology* 4:94–102 DOI 10.1016/j.funeco.2010.09.003.

Chen K-H, Miadlikowska J, Molnár K, Arnold AE, U’Ren JM, Gaya E, Gueidan C, Lutzoni F. 2015. Phylogenetic analyses of eurotiomycetous endophytes reveal their close affinities to Chaetothyriales, Eurotiales, and a new order—Phaeomoniellales. *Molecular Phylogenetics and Evolution* 85:117–130 DOI 10.1016/j.ympev.2015.01.008.
Cotty PJ, Cleveland TE, Brown RL, Melton JE. 1990. Variation in polygalacturonase production among Aspergillus flavus isolates. Applied and Environmental Microbiology 56:3885–3887.

Davey ML, Heegaard E, Halvorsen R, Kauserud H, Ohlson M. 2013. Amplicon-pyrosequencing-based detection of compositional shifts in bryophyte-associated fungal communities along an elevation gradient. Molecular Ecology 22:368–383 DOI 10.1111/mec.12122.

Davis EC, Shaw AJ. 2008. Biogeographic and phylogenetic patterns in diversity of liverwort-associated endophytes. American Journal of Botany 95:914–924 DOI 10.3732/ajb.2006463.

Dray S, Dufour AB. 2007. The ade4 package: implementing the duality diagram for ecologists. Journal of Statistical Software 22:1–20.

Ewing B, Green P. 1998. Base-calling of automated sequencer traces using Phred. II. Error probabilities. Genome Research 8:186–194.

Ewing B, Hillier L, Wendl M, Green P. 1998. Base-calling of automated sequencer traces using Phred. I. Accuracy assessment. Genome Research 8:175–185.

Fisher RA, Corbet AS, Williams CB. 1943. The relation between the number of species and the number of individuals in a random sample of an animal population. Journal of Animal Ecology 12:42–58 DOI 10.2307/1411.

Fouda AH, El-Din Hassan S, Eid AM, El-Din Ewais E. 2015. Biotechnological applications of fungal endophytes associated with medicinal plant Asclepias sinaica (Bioss.). Annuals of Agricultural Sciences 60:95–104 DOI 10.1016/j.j.aaos.2015.04.001.

Fröhlich J, Hyde KD. 1999. Biodiversity of palm fungi in the tropics: are global fungal diversity estimates realistic? Biodiversity and Conservation 8:977–1004 DOI 10.1023/A:1008895913857.

Gallery R, Dalling JW, Arnold AE. 2007. Diversity, host affinity, and distribution of seed-infecting fungi: a case study with neotropical Cecropia. Ecology 88:582–588 DOI 10.1890/05-1207.

Gardes M, Bruns TD. 1993. ITS primers with enhanced specificity of basidiomycetes: application to the identification of mycorrhizae and rusts. Molecular Ecology 2:113–118 DOI 10.1111/j.1365-294X.1993.tb00005.x.

Gazis R, Miadlikowska J, Lutzoni F, Arnold AE, Chaverri P. 2012. Culture-based study of endophytes associated with rubber trees in Peru reveals a new class of Pezizomycotina: xylonomycetes. Molecular Phylogenetics and Evolution 65:294–304 DOI 10.1016/j.ympev.2012.06.019.

Hammer Ø, Harper DAT, Ryan PD. 2001. PAST: paleontological statistics software package for education and data analysis. Palaeontologia Electronica 4:1–9.

He X, Han G, Lin Y, Tian X, Xiang C, Tian Q, Wang F, He Z. 2012. Diversity and decomposition potential of endophytes in leaves of a Cinnamomum camphora plantation in China. Ecological Research 27:273–284 DOI 10.1007/s11284-011-0898-0.

Hirose D, Osono T. 2006. Development and seasonal variations of Lophodermium populations on Pinus thunbergii needle litter. Mycoscience 47:242–247 DOI 10.1007/S10267-006-0299-3.
Hortal J, Borges PAV, Gaspar C. 2006. Evaluating the performance of species richness estimators: sensitivity to sample grain size. *Journal of Animal Ecology* **75**:274–287 DOI 10.1111/j.1365-2656.2006.01048.x.

Hudson HJ. 1968. The ecology of fungi on plant remains above the soil. *New Phytologist* **67**:837–874 DOI 10.1111/j.1469-8137.1968.tb06399.x.

Hultén E. 1965. *Flora of Alaska and neighboring territories; a manual of the vascular plants*. Stanford: Stanford University Press.

Jumpponen A, Jones KL. 2009. Massively parallel 454 sequencing indicates hyperdiverse fungal communities in temperate *Quercus macrocarpa* phyllosphere. *New Phytologist* **184**:438–448 DOI 10.1111/j.1469-8137.2009.02990.x.

Katoh K, Toh H. 2008. Recent developments in the MAFFT multiple sequence alignment program. *Briefings in Bioinformatics* **9**:286–298 DOI 10.1093/bib/bbn013.

Koide K, Osono T, Takeda H. 2005. Fungal succession and decomposition of *Camellia japonica* leaf litter. *Ecological Research* **20**:599–609 DOI 10.1007/s11284-005-0077-2.

Korkama-Rajala T, Müller MM, Pennanen T. 2008. Decomposition and fungi of needle litter from slow- and fast-growing Norway spruce. *Microbial Ecology* **56**:76–89 DOI 10.1007/s00248-007-9326-y.

Langenfeld A, Prado S, Nay B, Cruaud C, Lacoste S, Bury E, Hachette F, Hosoya T, Dupont J. 2013. Geographic locality greatly influences fungal endophyte communities in *Cephalotaxus harringtonia*. *Fungal Biology* **117**:124–136 DOI 10.1016/j.funbio.2012.12.005.

Liggenstoffer A, Youssef N, Couger M, Elshahed M. 2010. Phylogenetic diversity and community structure of anaerobic gut fungi (phylum Neocallimastigomycota) in ruminant and non-ruminant herbivores. *ISME Journal* **4**:1225–1235 DOI 10.1038/ismej.2010.49.

Lin Y, He X, Ma T, Han G, Xiang C. 2015. Priority colonization of *Cinnamomum camphora* litter by endophytes affects decomposition rate, fungal community and microbial activities under field conditions. *Pedobiologia* **58**:177–185 DOI 10.1016/j.pedobi.2015.09.001.

Lindahl BD, Ihrmark K, Boberg J, Trumbore SE, Högberg P, Stenlid J, Finlay RD. 2007. Spatial separation of litter decomposition and mycorrhizal nitrogen uptake in a boreal forest. *New Phytologist* **173**:611–620 DOI 10.1111/j.1469-8137.2006.01936.x.

Magurran AE. 2004. *Measuring biological diversity*. Oxford: Blackwell Publishing Ltd.

Manter DK, Bakker MG. 2015. Estimating beta diversity for under-sampled communities using the variably weighted Odum dissimilarity index and OTUshuff. *Bioinformatics* **31**:3451–3459 DOI 10.1093/bioinformatics/btv394.

Oksanen J, Guillaume Blanchet F, Kindt R, Legendre P, Minchin PR, O’Hara RB, Simpson GL, Solymos P, Henry M, Stevens H, Wagner H. 2015. vegan: Community Ecology Package. R package version 2.3-2. Available at https://CRAN.R-project.org/package=vegan.

Oono R, Lutzoni F, Arnold AE, Kaye L, U’Ren JM, May G, Carbone I. 2014. Genetic variation in horizontally transmitted fungal endophytes of pine needles reveals
population structure in cryptic species. *American Journal of Botany* 101:1362–1374 DOI 10.3732/ajb.1400141.

Osono T. 2002. Phyllosphere fungi on leaf litter of *Fagus crenata*: occurrence, colonization, and succession. *Canadian Journal of Botany* 80:460–469 DOI 10.1139/b02-028.

Osono T. 2006. Role of phyllosphere fungi of forest trees in the development of decomposer fungal communities and decomposition processes of leaf litter. *Canadian Journal of Microbiology* 52:701–716 DOI 10.1139/w06-023.

Osono T, Hirose D. 2009. Effects of prior decomposition of *Camellia japonica* leaf litter by an endophytic fungus on the subsequent decomposition by fungal colonizers. *Mycoscience* 50:52–55 DOI 10.1007/S10267-008-0442-4.

Osono T, Hirose D. 2011. Colonization and lignin decomposition of pine needle litter by *Lophodermium pinastri*. *Forest Pathology* 41:156–162 DOI 10.1111/j.1439-0329.2010.00648.x.

Osono T, Takeda H. 2001. Organic chemical and nutrient dynamics in decomposing beech leaf litter in relation to fungal ingrowth and succession during 3-year decomposition processes in a cool temperate forest in Japan. *Ecological Research* 16:649–670 DOI 10.1046/j.1440-1703.2001.00426.x.

Peršoh D, Segert J, Zigan A, Rambold G. 2013. Fungal community composition shifts along a leaf degradation gradient in a European beech forest. *Plant Soil* 362:175–186 DOI 10.1007/s11104-012-1271-y.

Porras-Alfaro A, Bayman P. 2011. Hidden fungi, emergent properties: endophytes and microbiomes. *Annual Review of Phytopathology* 49:291–315 DOI 10.1146/annurev-phyto-080508-081831.

Promputtha I, Hyde KD, McKenzie EHC, Peberdy JF, Lumyong S. 2010. Can leaf degrading enzymes provide evidence that endophytic fungi becoming saprobes? *Fungal Diversity* 41:89–99 DOI 10.1007/s13225-010-0024-6.

Promputtha I, Lumyong S, Dhanasekaran V, McKenzie EHC, Hyde KD, Jeewon R. 2007. A phylogenetic evaluation of whether endophytes become saprotrophs at host senescence. *Microbial Ecology* 53:579–590 DOI 10.1007/s00248-006-9117-x.

Promputtha I, Lumyong S, Lumyong P, McKenzie EHC, Hyde KD. 2002. Fungal succession on senescent leaves of *Manglietia garrettii* in Doi Suthep-Pui National Park, northern Thailand. *Fungal Diversity* 10:89–100.

Radford AE, Ahles HE, Bell CR. 1968. *Manual of the vascular flora of the Carolinas*. Chapel Hill North Carolina: University of North Carolina Press.

Rodriguez RJ, White JF, Arnold AE, Redman RS. 2009. Fungal endophytes: diversity and functional roles. *New Phytologist* 182:314–330 DOI 10.1111/j.1469-8137.2009.02773.x.

Saunders M, Glenn AE, Kohn LM. 2010. Exploring the evolutionary ecology of fungal endophytes in agricultural systems: using functional traits to reveal mechanisms in community processes. *Evolutionary Applications* 3:525–537 DOI 10.1111/j.1752-4571.2010.00141.x.

Schloss PD, Wescott SL, Ryabin T, Hall JR, Hartmann M, Hollister EB, Lesniewski RA, Oakley BB, Parks DH, Robinson CJ, Sahl JW, Stres B, Thallinger GG,
Van Horn DJ, Weber CF. 2009. Introducing mothur: open-source, platform-independent, community-supported software for describing and comparing microbial communities. *Applied and Environmental Microbiology* 75:7537–7541 DOI 10.1128/AEM.01541-09.

Schulz B, Boyle C. 2005. The endophytic continuum. *Mycological Research* 109:661–686 DOI 10.1017/S095375620500273X.

Šnajdr J, Cajthaml T, Valášková V, Merhautová V, Petránková M, Spetz P, Leppänen K, Baldrian P. 2011. Transformation of *Quercus petraea* litter: successive changes in litter chemistry are reflected in deferential enzyme activity and changes in the microbial community composition. *FEMS Microbiology Ecology* 75:291–303 DOI 10.1111/j.1574-6941.2010.00999.x.

Stone JK. 1987. Initiation and development of latent infections by *Rhabdocline parkeri* on Douglas-fir. *Canadian Journal of Botany* 65:2614–2621 DOI 10.1139/b87-352.

Sun X, Guo L-D, Hyde KD. 2011. Community composition of endophytic fungi in *Acer truncatum* and their role in decomposition. *Fungal Diversity* 47:85–95 DOI 10.1007/s13225-010-0086-5.

Sun Y, Cai Y, Yu F, Farrrel ML, McKendree W, Farmerie W. 2009. ESPRIT: estimating species richness using large collections of 16S rRNA pyrosequences. *Nucleic Acids Research* 37(10):e76 DOI 10.1093/nar/gkp285.

Swofford DL. 2003. *PAUP*. Phylogenetic Analysis Using Parsimony (*and Other Methods). Version 4. Sunderland, Massachusetts: Sinauer Associates.

Taylor LR. 1978. Bates, Williams, Hutchinson—a variety of diversities. In: Mound LA, Warloff N, eds. *Diversity of insect faunas*. Oxford: Blackwell Publishing Ltd.

Tucker SL, Orbach MJ. 2007. *Agrobacterium*-mediated transformation to create an insertion library in *Magna porthe grisea*. In: Ronald PC, ed. *Methods in molecular biology 354: plant-pathogen interactions*, vol 354. Totowa: Humana Press, p 57–68.

Unterseher M, Persoh D, Schnittler M. 2013. Leaf-inhabiting endophytic fungi of European Beech (*Fagus sylvatica* L.) co-occur in leaf litter but are rare on decaying wood of the same host. *Fungal Diversity* 60:43–54 DOI 10.1007/s13225-013-0222-0.

Unterseher M, Schnittler M. 2009. Dilution-to-extinction cultivation of leaf-inhabiting endophytic fungi in beech (*Fagus sylvatica* L.)—different cultivation techniques influence fungal biodiversity assessment. *Mycological Research* 113:645–654.

Urbanová M, Šnajdr J, Baldrian P. 2015. Composition of fungal and bacterial communities in forest litter and soil is largely determined by dominant trees. *Soil Biology and Biochemistry* 84:53–64 DOI 10.1016/j.soilbio.2015.02.011.

U’Ren JM, Dalling JW, Gallery RE, Maddison DR, Davis EC, Gibson CM, Arnold AE. 2009. Diversity and evolutionary origins of fungi associated with seeds of a neotropical pioneer tree: a case study for analyzing fungal environmental samples. *Mycological Research* 113:432–449 DOI 10.1016/j.mycres.2008.11.015.

U’Ren JM, Lutzoni F, Miadlikowska J, Arnold AE. 2010. Community analysis reveals close affinities between endophytic and endolichenic fungi in mosses and lichens. *Microbial Ecology* 60:340–353 DOI 10.1007/s00248-010-9698-2.
U'Ren JM, Lutzoni F, Miadlikowska J, Laetsch A, Arnold AE. 2012. Host- and geographic structure of endophytic and endolichenic fungi at a continental scale. *American Journal of Botany* **99**:898–914 DOI 10.3732/ajb.1100459.

U'Ren JM, Miadlikowska J, Zimmerman NB, Stajich JE, Lutzoni F, Arnold AE. 2016. Contributions of North American endophytes to the phylogeny, ecology, and taxonomy of Xylariaceae (Sordariomycetes, Ascomycota). *Molecular Phylogenetics and Evolution* **98**:210–232 DOI 10.1016/j.ympev.2016.02.010.

U'Ren JM, Riddle J, Monacell JT, Carbone I, Miadlikowska J, Arnold AE. 2014. Tissue storage and primer selection influence pyrosequencing-based inferences of diversity and community composition of endolichenic and endophytic fungi. *Molecular Ecology Resources* **14**:1032–1048 DOI 10.1111/1755-0998.12252.

Vilgalys R, Hester M. 1990. Rapid genetic identification and mapping of enzymatically amplified ribosomal DNA from several *Cryptococcus* species. *Journal of Bacteriology* **172**:4238–4246 DOI 10.1128/jb.172.8.4238-4246.1990.

Voříšková J, Baldrian P. 2013. Fungal community on decomposing leaf litter undergoes rapid successional changes. *ISME Journal* **7**:477–486 DOI 10.1038/ismej.2012.116.

Wang Q, Garrity GM, Tiedje JM, Cole JR. 2007. Naïve bayesian classifier for rapid assignment of rRNA sequences into the new bacterial taxonomy. *Applied and Environmental Microbiology* **73**:5261–5267 DOI 10.1128/AEM.00062-07.

White TJ, Bruns T, Lee S, Taylor J. 1990. Amplification and direct sequencing of fungal ribosomal RNA genes for phylogenetics. In: Innis MA, Gelfand DH, Sninsky JJ, White TJ, eds. *PCR Protocols: a guide to methods and applications*. San Diego: Academic Press.

Yuan Z, Chen L. 2014. The role of endophytic fungal individuals and communities in the decomposition of *Pinus massoniana* needle litter. *PLoS ONE* **9**(8):e105911 DOI 10.1371/journal.pone.0105911.

Yuan Z-L, Rao L-B, Chen Y-C, Zhang C-L, Wu Y-G. 2011. From pattern to process: species and functional diversity in fungal endophytes of *Abies beshanzuensis*. *Fungal Biology* **115**:197–213 DOI 10.1016/j.funbio.2010.11.002.

Zimmerman NB, Vitousek PM. 2012. Fungal endophyte communities reflect environmental structuring across a Hawaiian landscape. *Proceedings of the National Academy of Sciences of the United States of America* **109**:13022–13027 DOI 10.1073/pnas.1209872109.