New species and records of *Trichoderma* isolated as mycoparasites and endophytes from cultivated and wild coffee in Africa

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A survey for species of the genus *Trichoderma* occurring as endophytes of *Coffea*, and as mycoparasites of coffee rusts (*Hemileia*), was undertaken in Africa; concentrating on Cameroon and Ethiopia. Ninety-four isolates of *Trichoderma* were obtained during this study: 76 as endophytes of healthy leaves, stems and berries and, 18 directly from colonized rust pustules. A phylogenetic analysis of all isolates used a combination of three genes: translation elongation factor-1α (*tef1*), *rpb2* and *cal* for selected isolates. GCPSR criteria were used for the recognition of species; supported by morphological and cultural characters. The results reveal a previously unrecorded diversity of *Trichoderma* species endophytic in both wild and cultivated *Coffea*, and mycoparasitic on *Hemileia* rusts. Sixteen species were delimited, including four novel taxa which are described herein: *T. botryosum*, *T. caeruloviride*, *T. lentissimum* and *T. pseudopyramidale*. Two of these new species, *T. botryosum* and *T. pseudopyramidale*, constituted over 60% of the total isolations, predominantly from wild *C. arabica* in Ethiopian cloud forest. In sharp contrast, not a single isolate of *Trichoderma* was obtained using the same isolation protocol during a survey of coffee in four Brazilian states, suggesting the existence of a ‘*Trichoderma* void’ in the endophyte mycobiota of coffee outside of Africa. The potential use of these African *Trichoderma* isolates in classical biological control, either as endophytic bodyguards—to protect coffee plants from *Hemileia vastatrix*, the fungus causing coffee leaf rust (CLR)—or to reduce its impact through mycoparasitism, is discussed, with reference to the on-going CLR crisis in Central America.

Species of the ascomycete genus *Trichoderma* (*Hypocreales: Hypocreaceae*) are widely distributed in different environments and have a variety of biological activities¹. In the last two decades various studies have investigated the diversity and taxonomy of *Trichoderma* and numerous novel species have emerged using DNA sequence data²–³. However, despite the various surveys aimed at covering the diversity of this genus, such studies have been concentrated mostly in Asia, Europe and the Americas⁶–¹³. In contrast, until now, Africa has been poorly covered in terms of assessing the diversity of *Trichoderma*, with the exception of some studies involving specific regions or ecological niches, such as soil in South Africa¹⁴. In the case of *Trichoderma* occurring as endophytes in *Coffea*, there is a single study covering species isolated from the rhizosphere of *C. arabica* in Ethiopia¹⁵.

Fungi belonging to the genus *Trichoderma* have a recognized role as decomposers¹⁶,¹⁷ and for a long time they were considered to be soil saprotrophs of little practical relevance¹⁸,¹⁹. Currently, it is widely accepted that such a generalization was erroneous and many species of *Trichoderma* are now recognized as mycoparasites, as well as endophytes of woody plants¹⁸,²⁰–²⁴. The endophytic interaction between *Trichoderma* and their host-plants is intimate and may be complex, involving many steps at each level from direct contact to internal colonization of...
Endophytic Trichoderma may behave as inocuous commensals or as true symbionts; stimulating the plant’s defence system in various ways: inducing host resistance to plant pests; promoting tolerance to abiotic stresses; increasing plant growth and photosynthetic capability; and, contributing towards the solubilization of nutrients for the host plant’s benefit.

Studies on Trichoderma as endophytes in perennial crop plants, particularly in their original wild to semi-wild situations, have revealed a considerable diversity of species—including a number of novel taxa—especially when compared with the same crops in cultivation. Notable examples are cacao, Theobroma cacao, and rubber, Hevea brasiliensis in their native Amazonian ranges.

Members of Trichoderma compete naturally in the wild with other groups of fungi to occupy niches and obtain nutrients and are capable of producing a range of secondary metabolites, including antibiotics and mycotoxins. Another characteristic of Trichoderma is the mycoparasitic ability of certain species which has led them to be considered as potential tools for the control of plant pathogenic fungi. There are several practical examples of the commercial application of mycoparasitic Trichoderma: notably, that of T. stromaticum for control of Monilinia fructigena—the causal agent of witches’ broom disease of cacao—the most important disease of the crop in the Neotropics. This mycoparasite colonizes the necrotic brooms of diseased plants, as well as the agoric fruit bodies of the fungus, decreasing inoculum production, and it has also been reported to be a common endophyte in healthy cacao trees. A product based on T. stromaticum (Tricovab) has been distributed to farmers in southern Bahia (Brazil) for a number of years, and now forms part of an integrated management strategy.

Trichoderma species, such as T. harzianum, can colonize and degrade resistant structures (sclerotia) of other plant pathogenic fungi, and have been mass-produced and used as commercial bio-fungicides. Although the known diversity of Trichoderma is already high, with more than 200 species names recognized, based on molecular phylogeny, most research on mycoparasitism has been undertaken with only a few of these species. Including T. harzianum sensu lato, T. atroviride, T. viride, T. asperellum and T. asperelloides, whilst mycoparasitism of rust fungi by Trichoderma has been little studied. The potential of Trichoderma as a tool for the management of plant diseases is now widely recognized, although this approach has virtually been unapplied for many diseases of tropical perennial crops. The main aim of this study was to collect, isolate and identify members of the genus Trichoderma from Coffea species and their associated Hemileia rusts in their centres of origin in Africa, with the long-term objective of assessing their potential as biocontrol agents of coffee leaf rust (CLR) caused by Hemileia vastatrix an increasing constraint to coffee production in the Americas.

Results
Phylogenetic analyses and GCPRS. A total of 94 Trichoderma isolates were obtained during this survey, 76 as endophytes in Coffea spp. and 18 as mycoparasites on coffee rusts (Table 1). The combined data set indicated that the 94 Trichoderma strains grouped into 16 highly supported monophyletic groups (Figs. 1 and 2). The concatenated trees generated in BI, ML and MP analyses shared a similar topology, providing high support to the phylogenetic species belonging to the clade Harzianum. Five isolates were grouped into three known species belonging to the clade Harzianum: T. brevem, T. guizhouense and T. aggressivum; one isolate was identified as T. viride in the clade Viride and three as T. spirale in the clade Stricpile (Fig. 1). Three isolates were grouped in T. parareesei, belonging to the Longibrachiatum clade and an additional isolate, obtained from Brazil (as a mycoparasite of CLR rusts), also fell within this clade, and was identified as T. andinense (Fig. 2). Twenty-one isolates were grouped into five species of the Viride clade: T. koningii, T. peterseni, T. theobromicola, T. hamatum and T. atroviride (Fig. 2). Fifty-nine isolates grouped in three phylogenetic species belonging to the clade Harzianum and one isolate belonging to the Viride clade did not correspond to any known species and were considered as new taxa, described in this work as: T. lentissimum sp. nov., T. caeruloviride sp. nov., T. botryosum sp. nov. and T. pseudopyramidale sp. nov. (Figs. 1 and 2). In order to clarify the phylogenetic relationship between T. pseudopyramidale sp. nov. and T. pyramidal, an analysis was performed with the addition of calmodulin sequences. The results of the analysis supported the distinction between T. pyramidal and the new species (Fig. 3). The isolates identified in this study as T. pseudopyramidale were positioned as paraphyletic with T. pyramidal reference isolates, in the tef and rpb2 tree (Fig. 4) and in the rpb2 tree (Fig. 5), the sequence of the single available reference isolate of T. pyramidal (S73) was distant from the T. pseudopyramidale clade.

The single tef and rpb2 trees for the Longibrachiatum and Viride clades were highly congruent (Figs. 6 and 7) with the topology of the concatenated tree (Fig. 2); the same was observed with the individual trees tef and rpb2 of the clade Harzianum (Figs. 4 and 5) with the topology concatenated (Fig. 1), except for the isolates identified as T. brevem (indicated with number 1, Fig. 5), when evaluated in the rpb2 tree. The reference isolates of this species were placed in two polyphyletic species and one of the isolates of this study attributed to T. brevem were positioned outside the monophyletic groups (they remained as singletons) in this analysis. In the tef tree the clades Stricpile and Viride divided the largest clade (Harzianum) (Fig. 4), however the grouping of the species identified in this study were similar to the concatenated tree (Fig. 1).

Diversity and distribution. Although the collections of plant material were not systematic—and, therefore, there was no purpose in quantifying the frequency of colonization of plants by species of Trichoderma in this study—it was possible to identify patterns of occurrence of taxa, in terms of region/locality, host Coffea species and plant tissue type. For example, it was observed that T. koningii was isolated only from leaves; whilst T. theobromicola, T. guizhouense and T. spirale were isolated exclusively from stems; and T. caeruloviride sp. nov.
| Taxon Isolate | Country | Substrate | Genbank accession numbers |
|--------------|---------|-----------|-------------------------|
| T. aggressivum COAD 2432 | Kenya | stem, Coffea sp. | MK044156 MK044249 |
| T. andinense COAD 2431 | Brazil | stem, Coffea sp. | MK044155 MK044248 |
| T. atroviride COAD 2396 | Kenya | leaf, Coffea sp. | MK044083 MK044177 |
| T. botryosum sp. nov. COAD 2422 | Ethiopia | berry, Coffeea arabica | MK044119 MK044212 |
| T. botryosum sp. nov. COAD 2401 | Cameroon | stem, Coffeea canephora | MK044088 MK044181 |
| T. botryosum sp. nov. COAD 2403 | Cameroon | stem, Coffeea arabica | MK044090 MK044183 |
| T. botryosum sp. nov. COAD 2505 | Ethiopia | stem, Coffea arabica | MK044112 MK044205 |
| T. botryosum sp. nov. COAD 2507 | Ethiopia | berry, Coffeea arabica | MK044116 MK044209 |
| T. botryosum sp. nov. COAD 2424 | Ethiopia | leaf, Coffea arabica | MK044121 MK044214 |
| T. botryosum sp. nov. COAD 2538 | Ethiopia | leaf, Coffea arabica | MK044222 MK044215 |
| T. botryosum sp. nov. COAD 2511 | Ethiopia | leaf, Coffea arabica | MK044126 MK044219 |
| T. botryosum sp. nov. COAD 2541 | Ethiopia | stem, Coffea arabica | MK044138 MK044231 |
| T. botryosum sp. nov. COAD 2542 | Ethiopia | stem, Coffeea arabica | MK044139 MK044232 |
| T. botryosum sp. nov. COAD 2520 | Ethiopia | stem, Coffeea arabica | MK044140 MK044233 |
| T. botryosum sp. nov. COAD 2543 | Ethiopia | stem, Coffea arabica | MK044141 MK044234 |
| T. botryosum sp. nov. COAD 2521 | Ethiopia | stem, Coffeea arabica | MK044142 MK044235 |
| T. botryosum sp. nov. COAD 2522 | Ethiopia | stem, Coffeea arabica | MK044143 MK044236 |
| T. botryosum sp. nov. COAD 2423 | Ethiopia | stem, Coffeea arabica | MK044144 MK044237 |
| T. botryosum sp. nov. COAD 2524 | Ethiopia | stem, Coffeea arabica | MK044145 MK044238 |
| T. botryosum sp. nov. COAD 2525 | Ethiopia | stem, Coffeea arabica | MK044146 MK044239 |
| T. botryosum sp. nov. COAD 2526 | Ethiopia | stem, Coffeea arabica | MK044147 MK044240 |
| T. botryosum sp. nov. COAD 2428 | Ethiopia | berry, Coffeea arabica | MK044148 MK044241 |
| T. botryosum sp. nov. COAD 2527 | Ethiopia | leaf, Coffea arabica | MK044149 MK044242 |
| T. botryosum sp. nov. COAD 2528 | Ethiopia | leaf, Coffea arabica | MK044151 MK044244 |
| T. botryosum sp. nov. COAD 2530 | Ethiopia | leaf, Coffea arabica | MK044152 MK044245 |
| T. breve COAD 2402 | Cameroon | stem, Coffeea canephora | MK044089 MK044182 |
| T. breve COAD 2429 | Ethiopia | berry, Coffeea arabica | MK044150 MK044243 |
| T. caeruloviride sp. nov. COAD 2416 | Ethiopia | berry, Coffeea arabica | MK044108 MK044201 |
| T. caeruloviride sp. nov. COAD 2415 | Ethiopia | berry, Coffeea arabica | MK044109 MK044202 |
| T. guizhouense COAD 2397 | Kenya | stem, Coffea sp. | MK044084 MK044176 |
| T. guizhouense COAD 2398 | Kenya | stem, Coffea sp. | MK044085 MK044178 |
| T. hamatum COAD 2417 | Ethiopia | stem, Coffeea arabica | MK044110 MK044203 |
| T. hamatum COAD 2418 | Ethiopia | stem, Coffeea arabica | MK044111 MK044204 |
| T. hamatum COAD 2423 | Ethiopia | berry, Coffeea arabica | MK044220 MK044213 |
| T. koningiopsis COAD 2405 | Cameroon | leaf, Coffeea canephora | MK044092 MK044185 |
| T. koningiopsis COAD 2502 | Cameroon | leaf, Coffeea canephora | MK044097 MK044190 |
| T. koningiopsis COAD 2537 | Cameroon | leaf, Coffeea canephora | MK044098 MK044191 |
| T. koningiopsis COAD 2409 | Cameroon | leaf, Coffeea canephora | MK044099 MK044192 |
| T. koningiopsis COAD 2503 | Cameroon | leaf, Coffeea canephora | MK044100 MK044193 |
| T. koningiopsis COAD 2410 | Cameroon | leaf, Coffeea canephora | MK044101 MK044194 |
| T. koningiopsis COAD 2411 | Cameroon | leaf, Coffeea canephora | MK044102 MK044195 |
| T. lentissimum sp. nov. COAD 2399 | Kenya | stem, Coffeea cf. arabica | MK044086 MK044179 |
| T. pararosei COAD 2485 | Ethiopia | Hemiticia sp. Mycoparasite | MK044082 MK044265 |
| T. pararosei COAD 2482 | Ethiopia | stem, Coffeea arabica | MK044153 MK044246 |
| T. pararosei COAD 2483 | Ethiopia | stem, Coffeea arabica | MK044154 MK044247 |
| T. peterseni COAD 2438 | Ethiopia | Hemiticia sp. Mycoparasite | MK044168 MK044261 |
| T. pseudopyramidalis sp. nov. COAD 2419 | Ethiopia | stem, Coffeea arabica | MK044113 MK044206 MK084875 |
| T. pseudopyramidalis sp. nov. COAD 2506 | Ethiopia | stem, Coffeea arabica | MK044114 MK044207 |
| T. pseudopyramidalis sp. nov. COAD 2420 | Ethiopia | stem, Coffeea arabica | MK044115 MK044208 MK084874 |
| T. pseudopyramidalis sp. nov. COAD 2508 | Ethiopia | leaf, Coffeea arabica | MK044117 MK044210 |
| T. pseudopyramidalis sp. nov. COAD 2421 | Ethiopia | leaf, Coffeea arabica | MK044118 MK044211 MK084873 |
| T. pseudopyramidalis sp. nov. COAD 2425 | Ethiopia | leaf, Coffeea arabica | MK044123 MK044216 MK084871 |
| T. pseudopyramidalis sp. nov. COAD 2509 | Ethiopia | leaf, Coffeea arabica | MK044124 MK044217 |
| T. pseudopyramidalis sp. nov. COAD 2510 | Ethiopia | leaf, Coffeea arabica | MK044125 MK044218 |
| T. pseudopyramidalis sp. nov. COAD 2540 | Ethiopia | leaf, Coffeea arabica | MK044127 MK044220 |
| T. pseudopyramidalis sp. nov. COAD 2512 | Ethiopia | leaf, Coffeea arabica | MK044128 MK044221 |

Continued
The other species were distributed in more than one plant tissue type. The pre-
Table 1. Trichoderma strains obtained in the survey and used in the phylogenetic analyses, with their corresponding geographic origin, host and tissue source. Ex-type strains are indicated in bold. *Close to H. caffecola, but currently being assessed as a new species of Hemileia (Authors, unpublished). †Identified as a wild and geographically-isolated population of Coffea arabica, common in the understorey forest. Kew Herbarium (Herb K) records reflect uncertainty about its true identity; botanical specimens from present survey deposited in Herb K.

| Taxon                  | Isolate | Country      | Substrate          | Genbank accession numbers |
|------------------------|---------|--------------|--------------------|--------------------------|
| T. pseudopyramidale sp. nov. | COAD 2513 | Ethiopia | Leaf, Coffea arabica | Endophyte MK044129 MK044222 - |
| T. pseudopyramidale sp. nov. | COAD 2514 | Ethiopia | Leaf, Coffea arabica | Endophyte MK044130 MK044223 - |
| T. pseudopyramidale sp. nov. | COAD 2426 | Ethiopia | Leaf, Coffea arabica | Endophyte MK044131 MK044224 MK084870 |
| T. pseudopyramidale sp. nov. | COAD 2515 | Ethiopia | Leaf, Coffea arabica | Endophyte MK044132 MK044225 - |
| T. pseudopyramidale sp. nov. | COAD 2516 | Ethiopia | Leaf, Coffea arabica | Endophyte MK044133 MK044226 - |
| T. pseudopyramidale sp. nov. | COAD 2517 | Ethiopia | Leaf, Coffea arabica | Endophyte MK044134 MK044227 - |
| T. pseudopyramidale sp. nov. | COAD 2518 | Ethiopia | Leaf, Coffea arabica | Endophyte MK044135 MK044228 - |
| T. pseudopyramidale sp. nov. | COAD 2427 | Ethiopia | Leaf, Coffea arabica | Endophyte MK044136 MK044229 MK084872 |
| T. pseudopyramidale sp. nov. | COAD 2519 | Ethiopia | Leaf, Coffea arabica | Endophyte MK044137 MK044230 - |
| T. pseudopyramidale sp. nov. | COAD 2433 | Cameroon | Hemileia sp. Mycoparasite | MK044157 MK044230 MK084869 |
| T. pseudopyramidale sp. nov. | COAD 2434 | Ethiopia | Hemileia sp. Mycoparasite | MK044158 MK044231 MK084868 |
| T. pseudopyramidale sp. nov. | COAD 2529 | Ethiopia | Hemileia sp. Mycoparasite | MK044159 MK044232 - |
| T. pseudopyramidale sp. nov. | COAD 2435 | Ethiopia | Hemileia sp. Mycoparasite | MK044160 MK044233 MK084867 |
| T. pseudopyramidale sp. nov. | COAD 2530 | Ethiopia | Hemileia sp. Mycoparasite | MK044161 MK044234 - |
| T. pseudopyramidale sp. nov. | COAD 2436 | Ethiopia | Hemileia sp. Mycoparasite | MK044162 MK044235 MK084865 |
| T. pseudopyramidale sp. nov. | COAD 2531 | Ethiopia | Hemileia sp. Mycoparasite | MK044163 MK044236 - |
| T. pseudopyramidale sp. nov. | COAD 2532 | Ethiopia | Hemileia sp. Mycoparasite | MK044164 MK044237 - |
| T. pseudopyramidale sp. nov. | COAD 2437 | Ethiopia | Hemileia sp. Mycoparasite | MK044165 MK044238 MK084866 |
| T. pseudopyramidale sp. nov. | COAD 2533 | Ethiopia | Hemileia sp. Mycoparasite | MK044166 MK044239 MK084865 |
| T. pseudopyramidale sp. nov. | COAD 2534 | Ethiopia | Hemileia sp. Mycoparasite | MK044167 MK044240 - |
| T. pseudopyramidale sp. nov. | COAD 2535 | Ethiopia | Hemileia sp. Mycoparasite | MK044169 MK044242 - |
| T. pseudopyramidale sp. nov. | COAD 2536 | Ethiopia | Hemileia sp. Mycoparasite | MK044170 MK044243 - |
| T. pseudopyramidale sp. nov. | COAD 2439 | Ethiopia | Hemileia sp. Mycoparasite | MK044171 MK044244 MK084864 |
| T. pseudopyramidale sp. nov. | COAD 2591 | Ethiopia | Stem, Coffea arabica | Endophyte MK044174 MK044268 - |
| T. pseudopyramidale sp. nov. | COAD 2592 | Ethiopia | Stem, Coffea arabica | Endophyte MK044175 MK044269 - |
| T. spiral  | COAD 2404 | Cameroon | Stem, Coffea canephora | Endophyte MK044091 MK044184 - |
| T. spiral  | COAD 2408 | Cameroon | Stem, Coffea canephora | Endophyte MK044096 MK044189 - |
| T. theobromicola  | COAD 2406 | Cameroon | Stem, Coffea canephora | Endophyte MK044093 MK044186 - |
| T. theobromicola  | COAD 2407 | Cameroon | Stem, Coffea canephora | Endophyte MK044094 MK044187 - |
| T. theobromicola  | COAD 2501 | Cameroon | Stem, Coffea canephora | Endophyte MK044095 MK044188 - |
| T. theobromicola  | COAD 2504 | Cameroon | Stem, Coffea canephora | Endophyte MK044103 MK044196 - |
| T. theobromicola  | COAD 2412 | Cameroon | Stem, Coffea canephora | Endophyte MK044104 MK044197 - |
| T. theobromicola  | COAD 2440 | Cameroon | Stem, Coffea canephora | Endophyte MK044106 MK044199 - |
| T. theobromicola  | COAD 2414 | Cameroon | Stem, Coffea canephora | Endophyte MK044107 MK044200 - |
| T. theobromicola  | COAD 2589 | Cameroon | Stem, Coffea canephora | Endophyte MK044172 MK044266 - |
| T. theobromicola  | COAD 2590 | Cameroon | Stem, Coffea canephora | Endophyte MK044173 MK044267 - |
| T. virens  | COAD 2400 | Cameroon | Stem, Coffea brevipes | Endophyte MK044087 MK044180 - |

was isolated from berries only. The other species were distributed in more than one plant tissue type. The pre-

Table 2. T. botryosum sp. nov. 20 isolates from T. botryosum sp. nov. from leaves, stems and berries; 34 isolates of T. pseudopyramidale sp. nov. from leaves and stems, and as myco-

was isolated from berries only. The other species were distributed in more than one plant tissue type. The pre-

dominant taxon in all plant tissues was T. botryosum sp. nov., but this species was never found as a mycoparasite, whereas T. pseudopyramidale sp. nov. was isolated both as an endophyte and as a mycoparasite—almost exclu-
sively from Ethiopia—being common on Hemileia rust in forest coffee populations. The species T. aggressivum, T. andinensis, T. parareesei and T. peterseni, were isolated only once during the survey, from Kenya, Brazil, and

In Ethiopia, 64 isolates belonging to seven species were isolated, namely: 20 isolates of T. botryosum sp. nov. from leaves, stems and berries; 34 isolates of T. pseudopyramidale sp. nov. from leaves and stems, and as myco-

parasites; three isolates of T. parareesei from stems and one as a mycoparasite; three isolates of T. hamatum from stems and berries; two isolates of T. caeruloviride sp. nov. from berries; one isolate of T. peterseni growing as a
mycoparasite and one isolate of \textit{T. breve} from berries (Table 2). In Kenya, four species were collected, namely: one isolate of \textit{T. aggressivum} growing as a mycoparasite; one isolate of \textit{T. atroviride} from leaves; one isolate of \textit{T. guizhouense} and one of \textit{T. lentissimum} sp. nov. from stems (Table 2). When the diversity of \textit{Trichoderma}

Figure 1. Bayesian phylogenetic tree of clades \textit{Harzianum}, \textit{Strictipile} and \textit{Virens}. The tree was based on a concatenated \textit{tef1} and \textit{rpb2} sequence dataset. Bootstrap values ($\geq 70\%$) of the ML and MP analyses, as well as posterior probability scores ($\geq 0.9$) from a Bayesian analysis of the same dataset, are indicated at well supported nodes together with thickened branches. The isolates belonging to known species, obtained in this study, are in bold. Isolates of new species, described in this study, are in bold red. The tree was rooted with \textit{Trichoderma asperellum} (TR3). The phylogenetic tree was edited using Inkscape 1.0 (https://inkscape.org/pt-br/).
from coffee in West Africa (Cameroon) is compared with that of East Africa (Kenya and Ethiopia), 
*T. breve*, *T. pseudopyramidale* sp. nov. and *T. botryosum* sp. nov. are the only species which are present both in Ethiopia and Cameroon. The four species found in Kenya were absent from coffee samples in the other countries. Although

Figure 2. Bayesian phylogenetic tree of clades *Longibrachiatum* and *Viride*. The tree was based on a concatenated tef1 and rpb2 sequence dataset. Bootstrap values (≥70%) of the ML and MP analyses, as well as posterior probability scores (≥0.9) from a Bayesian analysis of the same dataset, are indicated at well supported nodes together with thickened branches. The isolates belonging to known species, obtained in this study, are in bold. Isolates of new species, described in this study, are in bold red. The tree was rooted with *Protocrea pallida* (CBS 121552). The phylogenetic tree was edited using Inkscape 1.0 (https://inkscape.org/pt-br/).
this suggests strong endemism and isolation of the Trichoderma mycobiota of coffee in Kenya, this should be treated with caution since sampling in Kenya was limited and, thus, this may be an artefact. However, the fact that there was no commonality in species between Kenya and the other countries is intriguing and warrants further investigation. The only Brazilian isolate was identified as T. andinense. The occurrence of T. andinense appeared in an ad hoc isolation during a search for mycoparasites of CLR pustules in Brazil.

The coffee survey in Brazil—aimed at obtaining endophytic Trichoderma from plants growing in semi-wild conditions at eight localities in four states (Table 3)—yielded isolates comprising a range of different genera but just a single isolate was identified as belonging to the genus Trichoderma.

When comparing the number of taxa of Trichoderma present in C. canephora and C. arabica, it was found that both harbour five species but these are distributed differently in the host tissues. The only Trichoderma spp. occurring in both C. arabica and C. canephora were T. breve and T. botryosum sp. nov. In C. arabica, the highest diversity of Trichoderma was found in berries (four species), whereas in C. canephora the highest diversity was found in stems (four species); however, since most of the sampling was focused on C. arabica, it would be premature to reach a firm conclusion. Stems yielded the highest number of isolates from the aerial tissues of Coffea, with 40 (42.6% of the total), followed by leaves with 29 (30.8%), and seven isolates (7.4%) from berries. Mycoparasites, with 18 isolates, formed 19.2% of the total (Tables 1 and 2).

**Taxonomy.** Four additions to the genus Trichoderma emerged from the phylogenetic study of the isolates obtained during this survey of Coffea in Africa. Morphological and cultural information proved useful to confirm their separation from closely-related known species of Trichoderma; providing further evidence for their recognition as novel species. Type cultures were deposited in the internationally-recognized culture collection of the Universidade Federal de Viçosa (COAD). The following species were collected and identified, with authority names and publication details as recommended by Bissett et al.12.

*Trichoderma aggressivum*—Samuels & W. Gams—in Samuels et al., *Mycologia* 94: 167, 2002. MycoBank: MB484638.

Description and illustration, see5,56.

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**Figure 3.** Bayesian phylogenetic tree of *T. pyramidal*e and *T. pseudopyramidale* sp. nov. The tree was based on a concatenated tef1, rpb2 and cal sequence dataset. Bootstrap values (≥ 70%) of the ML and MP analyses, as well as posterior probability scores (≥ 0.9) from a Bayesian analysis of the same dataset, are indicated at well supported nodes together with thickened branches. The isolates belonging to known species, obtained in this study, are in bold. Isolates of new species, described in this study, are in bold red. The tree was rooted with Trichoderma lentiforme (DIS 167C and DIS 167E). The phylogenetic tree was edited using Inkscape 1.0 (https://inkscape.org/pt-br/).
Material examined. KENYA: Eastern Province, Marsabit National Park, Lake Paradise, primary forest, alt 1340 m: isolated as a mycoparasite of *Hemileia vastatrix* on leaves of *Coffea cf. arabica*, H.C. Evans & R.W. Barreto (culture COAD 2432).

**Figure 4.** Bayesian phylogenetic tree of clades *Harzianum*, *Strictipile* and *Virens*. The tree was based on tef1 sequence dataset. Bootstrap values (≥70%) of the ML analyses, as well as posterior probability scores (≥0.9) from a Bayesian analysis of the same dataset, are indicated at well supported nodes together with thickened branches. The isolates belonging to known species, obtained in this study are in bold. Isolates of new species, described in this study, are in bold red. The tree was rooted with *Trichoderma asperellum* (TR3). The phylogenetic tree was edited using Inkscape 1.0 (https://inkscape.org/pt-br/).
Notes: *Trichoderma aggressivum*, in the *Harzianum* clade, has been reported to cause the green mould epidemic in commercially grown *Agaricus bisporus* in North America. This is the first report of *T. aggressivum* as a mycoparasite of *Hemileia* and of rusts, in general; having been recorded previously only in mushroom farms in both North America and Europe. Thus, this appears to be the first record of *T. aggressivum* from the tropics. It has been shown to produce various antifungal compounds, and falls within the clade defined by a mycoparasitic mode of nutrition in a consensus phylogenetic tree.

**Figure 5.** Bayesian phylogenetic tree of clades *Harzianum, Strictipile* and *Virens*. The tree was based on rpb2 sequence dataset. Bootstrap values (≥70%) of the ML analyses, as well as posterior probability scores (≥0.9) from a Bayesian analysis of the same dataset, are indicated at well supported nodes together with thickened branches. The isolates belonging to known species, obtained in this study, are in bold. Isolates of new species, described in this study, are in bold red. The tree was rooted with *Trichoderma asperellum* (TR3). The phylogenetic tree was edited using Inkscape 1.0 (https://inkscape.org/pt-br/).
**Figure 6.** Bayesian phylogenetic tree of clades *Longibrachiatum* and *Viride*. The tree was based on tef1 sequence dataset. Bootstrap values (≥ 70%) of the ML analyses, as well as posterior probability scores (≥ 0.9) from a Bayesian analysis of the same dataset, are indicated at well supported nodes together with thickened branches. The isolates from known species obtained in this study are in bold. Isolates belonging to new species, described in this study, are in bold red. The tree was rooted with *Protocrea pallida* (CBS 121552). The phylogenetic tree was edited using Inkscape 1.0 (https://inkscape.org/pt-br/).

*Trichoderma andinense* (Samuels & O. Petrini) Samuels, Jaklitsch & Voglmayr—in Jaklitsch & Voglmayr, *Mycotaxon* 126:146, 2014. MycoBank: MB807417.

Synonym: *Hypocrea andinensis* Samuels & O. Petrini—in Samuels et al., *Stud. Mycol.* 41:13, 1998.

Description and illustration, see2,64.
Material examined. BRAZIL: Rio de Janeiro, Duas Barras, Fazenda do Campo, Sítio Recanto do Sossego, coffee farm, isolated as a mycoparasite of *Hemileia vastatrix* on leaves of *Coffea arabica*, 2 May 2015, R. W. Barreto (culture COAD 2431).

Notes: This fungus, in the *Longibrachiatum* clade, was originally described from its sexual morph collected from a log in the Venezuelan Andes. Similar isolates from soil in Saudi Arabia, Amazonian Peru and Hawaii were reported later but were considered to represent new taxa within the *T. andinense* sub-clade and this species “remains known only from a single collection”. Thus, this is the first report of *T. andinense* as a mycoparasite of H. vastatrix.
H. vastatrix, and as a mycoparasite, in general. It was shown to group in a plant saprotrophy clade in the study by Chaverri and Samuels24.

Trichoderma atroviride P. Karst.—in Finl. Mögelsvamp. 21, 1892. MycoBank: MB451289.

Synonym: Hypocrea atroviridis Dodd, Lieckfeldt & Samuels, Mycologia 95:36, 2003.

Description and illustration, see5.

Material examined. KENYA: Western Region, Nandi County, Kakamega Forest Reserve, Isecheno, coffee farm, alt 1600 m; isolated as an endophyte from leaf of Coffea arabica, 23 January 2015, H. C. Evans & R.W. Barreto (culture COAD 2396).

Notes: This is a cosmopolitan species, in the Viride clade, but it is more commonly isolated from soil in tropical regions and its sexual morph is rarely formed. Chaverri and Samuels show its habitat preference is soil, but in their phylogenetic tree, T. atroviride groups with species having a mycoparasitic mode of nutrition. In our study, it was isolated from a leaf of C. arabica and this is the first record of T. atroviride as an endophyte of coffee.

Trichoderma atroviride is more associated with the plant rhizosphere, rather than aerial tissues, and it has been shown recently that colonization of both maize and tomato roots by this fungus induces foliar herbivory resistance. Previously, it has been known to protect plants against root pathogens (Pythium, Rhizoctonia) through induced resistance and antibiosis.

Trichoderma botryosum M.C.H. Rodríguez, H.C. Evans & R.W. Barreto sp. nov.—MycoBank: MB832341 (Figs. 8g–i, 9).

Etymology: referring to the grape-like clusters of conidia.

Holotype. ETHIOPIA: Southern Nations, Nationalities and Peoples Region, Kaffa Zone, Bonga District, Gela Wild Coffee Biosphere Reserve, cloud forest, alt 1900 m; isolated as an endophyte from berries of Coffea arabica; 24 November 2015, H.C. Evans, K. Belachew & R.W. Barreto. Ex-type culture: COAD 2422. GenBank: TEF1 = MK044119; RPB2 = MK044212.

Culture characteristics—Colonies on PDA: Optimum growth temperature at 30 °C, 65 mm after 72 h, 56–61 mm at 25 °C, 25 mm at 35 °C; covering the plate after 96 h; cottony white aerial mycelium, green sporulation starting from the centre of the plate, with the formation of concentric rings; with a sweet odour but absence of exudates and soluble pigments. Colonies on CMD: Optimum growth temperature at 30 °C; 63 mm after 72 h, 55–58 mm at 25 °C, 43 mm at 35 °C; mycelium hyaline, poor sporulation, concentric rings absent and no odour. Colonies on SNA: Optimum growth temperature at 30 °C, 61 mm after 72 h, 49–53 mm at 25 °C, 28 mm

Table 2. Number of taxa collected in this survey per country-source.

| Species                     | Ethiopia | Cameroon | Kenya |
|-----------------------------|----------|----------|-------|
| Leaf | Stem | Berry | Mycoparasite | Leaf | Stem | Berry | Mycoparasite | Leaf | Stem | Berry | Mycoparasite |
| T. aggressivum               | 1        | 0        | 0      |       |       |       |       |       |       |       |       |
| T. atroviride                |          | 1        | 0      |       |       |       |       |       |       |       |       |
| T. botryosum sp. nov.        | 6        | 11       | 3      | 2     |       |       |       |       |       |       |       |
| T. caeruloviride sp. nov.    |          | 2        |       |       |       |       |       |       |       |       |       |
| T. guizhouense               | 2        | 1        | 0      |       |       |       |       |       |       |       |       |
| T. hamatum                  | 2        | 3        |       |       |       |       |       |       |       |       |       |
| T. koningopis               |          | 6        | 1      |       |       |       |       |       |       |       |       |
| T. breve                    |          | 1        | 1      |       |       |       |       |       |       |       |       |
| T. lentissimum sp. nov.      | 2        | 1        |       |       |       |       |       |       |       |       |       |
| T. parareesi                |          | 1        |       |       |       |       |       |       |       |       |       |
| T. petersenii               |          | 1        |       |       |       |       |       |       |       |       |       |

Table 3. Survey sites in Brazil of Coffea arabica, in semi-wild or forest situations, sampled for presence of Trichoderma endophytes.
at 35 °C; mycelium hyaline and smooth, green sporulation; forming concentric rings; exudates and soluble pigments absent.

Conidiophores pyramidal; phialides in whorls or pairs, lateral or terminal, lageniform to ampulliform, 4.0–8.0 (−8.6) × (1.9−) 2.3–3.1 µm (L/W), 1.2–2.4 µm in width at the base; supporting cells 5.4–15.6 × 1.8–3.4 µm (L/W); conidia globose to broadly ovoid, 1.4–3.3 × 1.6–2.8 µm (L/W), green, smooth; chlamydospores abundant, globose to ellipsoidal, terminal and intercalary, 4.4–8.1 × 3.8–7.3 µm (L/W).

Notes: *Trichoderma botryosum* grouped phylogenetically close to *T. afarasin* and *T. endophyticum* in the *Harzianum* clade. The new species is morphologically similar to its close relatives in: the pyramidal-type conidiophores; size of conidia and ampulliform phialides; and growth rate on PDA at 25 °C. The most distinctive morphological features in this species are the presence of chlamydospores and the grape-like clusters of conidia.

Figure 8. Colony characteristics of the new *Trichoderma* species on PDA, CMD and SNA. All colonies incubated at 25 °C under a 12 h day/night light regime and photographed on day seven. (a–c) *T. lentissimum* sp. nov.; (d–f) *T. caeruleoviride* sp. nov.; (g–i) *T. botryosum* sp. nov.; (j–l) *T. pseudopyramidale* sp. nov.
This appears to be a common endophyte of wild *C. arabica* in Ethiopia; being isolated from all the aerial tissues, with 20 isolates recorded (Table 2). It was uncommon in Cameroon with only two isolates from coffee stems.

*Trichoderma breve* K. Chen & W.Y. Zhuang—Chen & Zhuang., *Sci. Rep.* 7 (no. 9090): 7, 2017. MycoBank: MB809992.

Description and illustration, see12.

**Material examined.** CAMEROON: South-West Province, Busumbo, Mt. Etinde, coffee farm, 450 m; isolated as endophyte from stem of *Coffea canephora*, 17 November 2015, H.C. Evans, R.W. Barreto & M. K. Ndacnou (culture COAD 2402). ETHIOPIA: Southern Nations, Nationalities and Peoples Region, Kaffa Zone, Bonga District, Komba Wild Forest Reserve, cloud forest, 2000 m; isolated as an endophyte from a berry of *Coffea arabica*, 25 November 2015, H.C. Evans, R.W. Barreto & K. Belachew (culture COAD 2429).

**Notes:** *Trichoderma breve* is similar morphologically to the *T. harzianum* complex, and has previously been isolated from soil in the north of China12. Phylogenetic analyses indicate that *T. breve* is closely related to *T. bannanense*12, and in our study it lies close to *T. lentiforme*. This is the first report of *T. breve* in Africa and also as an endophyte of coffee; being recorded from both *C. canephora* and *C. arabica* in Cameroon and Ethiopia, respectively, with a single isolate from each country (Table 2).

*Trichoderma caeruloviride* M.C.H. Rodríguez, H.C. Evans & R.W. Barreto, sp. nov.—Mycobank: MB832340 (Figs. 8d–f, 10).

**Etymology:** referring to the blue-green colour of the conidial mass.

**Holotype:** ETHIOPIA: Southern Nations, Nationalities and Peoples Region, Kaffa Zone, Bonga District, Gedam Village, coffee farm, alt 1550 m; isolated as an endophyte from berries of *Coffea arabica*. 25 November 2015, H.C. Evans, K. Belachew & R.W. Barreto. Ex-type culture: COAD 2415. GenBank: TEF1 = MK044109; RPB2 = MK044202.

**Culture characteristics—**Colonies on PDA: Optimum growth at 25 °C, 51–53 mm after 72 h, 34 mm at 30 °C, no growth at 35 °C, covering the plate after 96 h; mycelium white, aerial, low with green sporulation, no formation of concentric rings; lacking pigmentation and odour. Colonies on CMD: Optimum growth temperature at 25 °C, 42–44 mm after 72 h, 19 mm at 30 °C, no growth at 35 °C; mycelium hyaline, poor sporulation, green conidia, no concentric rings present and no odour. Colonies on SNA: Optimum growth temperature at 25 °C, 29–36 mm after 72 h, 12 mm at 30 °C, no growth at 35 °C; mycelium hyaline and smooth, poor sporulation; green to blue conidia, with the formation of concentric rings; no pigmentation in the medium.

Conidiophores pyramidal with verticillate, paired lateral branches; phialides generally formed on terminal branches, in divergent whorls of three to four, (5.2–) 5.3–12.2 (–13.2) × (1.7–) 2.0–2.8 (–3.48) µm, mean...
7.4 × 2.5 µm (L/W); supporting cells (5.4–) 7.9–9.7 (–10.1) × 1.7–2.0 (–3.0) µm, mean 8.5 × 1.9 µm (L/W); conidia ellipsoidal to ovoid, green, smooth, 2.2–3.0 (–3.2) × (1.9–) 2.3–3.1 (–3.4) µm, mean 2.8 × 2.8 µm (L/W); chlamydospores terminal and intercalary, globose, 3.3–5.0 (–6.6) × 3.0–4.6 (–5.3) µm, mean 4.4 × 3.7 µm (abundantly formed on CMD after 4 days).

Notes: Phylogenetic analyses placed T. caeruloviride close to T. amazonicum and T. pleuroticola in the Harzianum clade. The new species can be distinguished from its nearest relatives by: no growth at 35 °C; the presence of a coconut-like odour on PDA; and the blue-green conidia in SNA microculture. Morphologically, T. caeruloviride is distinct from T. amazonicum which has a branching pattern of the pachybasium type, elliptical to subglobose conidia, ampulliform phialides, and chlamydospore-like structures in clusters. Trichoderma caeruloviride shares some morphological characteristics with T. pleuroticola, such as the pyramidal-type branching pattern, the globose conidia and the formation of chlamydospores; but can be separated by the larger, lageniform phialides of T. caeruloviride. Both isolates were from berries of C. arabica in Ethiopia and it appears to be a rare species (Table 2).

**Trichoderma guizhouense** Q.-R. Li, E.H.C. McKenzie & Yong Wang—in Li et al., Mycol. Prog. 12:170, 2012. MycoBank: MB563664.

Description and illustration, see4,55. Material examined. KENYA: Eastern Province, Marsabit National Park. Lake Paradise, primary forest, alt 1340 m; isolated as an endophyte from stem of Coffea sp. (wild population close to C. arabica and C. canephora, but identity uncertain, even after examination by an authority on the genus Coffea) H.C. Evans & R.W. Barreto, 28 January 2015 (cultures COAD 2397 and COAD 2398).

Notes: Trichoderma guizhouense, in the Harzianum clade, was first isolated from soil in Guizhou Province of China38, and has since shown promise as a biocontrol agent of Rhizoctonia root rot27. It was also isolated as an endophyte from the woody liana, Ancistrocladus korupensis (Ancistrocladaceae)—extracts of which are active against HIV59—and from the sapwood of Cola spp. (Malvaceae) in rainforest of the Cameroon Republic62 (H.C. Evans unpubl.). This is the first report of T. guizhouense as a coffee endophyte.

**Trichoderma hamatum** (Bonord.) Bainier—Bull. Soc. Mycol. Fr. 22:131, 1906. MycoBank: MB165799.
Synonyms: *Verticillium hamatum* Bonord., *Handb. Allgem. Mykol.*, 97, 1851. *Pachybasium hamatum* (Bonord.) Sacc., *Rev. Mycol. Toulouse* 7:161, 1885. *Rhymatotrichum hamatum* (Bonord.) Oudem., *Ned. Kruidk. Archf.* 3:908, 1903.

**Description and illustration.** See1,3,5. **Material examined.** EThIOPIA: Southern Nations, Nationalities and Peoples Region, Kaffa Zone, Bonga District, Gedam Village, coffee farm, alt 1550 m, isolated as an endophyte from stems of *Coffea arabica*, 25 November 2015, H.C. Evans, R.W. Barreto & K. Belachew (cultures COAD 2418 and COAD 2417). EThIOPIA: Southern Nations, Nationalities and Peoples Region, Kaffa Zone, Bonga District, Gela Wild Coffee Biosphere Reserve, cloud forest, alt 1700 m; isolated as an endophyte from berry of *Coffea arabica*, 25 November 2015, H.C. Evans, R.W. Barreto & K. Belachew (COAD 2423).

**Notes:** *Trichoderma hamatum*, in the Viride clade, is a cosmopolitan species, originally isolated from soil but it has also been reported as an endophyte in both stems and pods of wild *Theobroma gleri* from sub-montane forest in western Ecuador20. In addition, it was identified as a mycoparasite of frosty pod disease (*Monilithophthora roeri*) on the same host in this ecosystem20. It has also been isolated from the rhizosphere of *C. arabica* in Ethiopia15. In our study, *T. hamatum* was obtained from stems and berries of *C. arabica* in both cultivated and wild coffee plants, also in Ethiopia. It is most frequently cited as a colonizer of the rhizosphere, and some soil strains have been shown to promote crop growth, to activate biocontrol mechanisms against root pathogens and to induce systemic resistance to foliar pathogens75. Previously, an endophyte strain from the pod of a wild *Theobroma* species was found to promote the growth and delay drought symptoms in cacao plants77.

**Trichoderma koningiopsis** Samuels, C. Suárez & H.C. Evans—in Samuels et al., *Stud. Mycol.* 56:117, 2006. MycoBank: MB487454.

**Description and illustration.** See1,6,9. **Material examined.** CAMEROON: Eastern Province, Zemele Village, coffee farm, alt 660 m; isolated as an endophyte from leaves of *Coffea canephora*, 22 November 2015, H.C. Evans, R.W. Barreto & M. K. Ndacnou (Cultures COAD 2537; COAD 2405; COAD 2502; COAD 2503; COAD 2410; COAD 2411 and COAD 2409).

**Notes:** *Trichoderma koningiopsis*, in the Viride clade, is a cosmopolitan species, but it is more frequently recorded in tropical rather than temperate regions, and mostly from soil. During a survey of *Trichoderma* diversity in soil and leaf litter from the Amazonian rainforest of Colombia, *T. koningiopsis* was amongst the commonest species isolated73. In the Atlantic rainforest of Brazil, *T. koningiopsis* was found to be the dominant *Trichoderma* species in leaves being carried by *Atta* leaf-cutting ants, and subsequently rejected by them from the nest: it was posited that the ants recognized the threat posed by this mycoparasite to the fungal garden74. It has also been reported as a common stem endophyte in a species of *Theobroma* in sub-montane forest in western Ecuador24. It has also been shown to colonize cacao plants via the leaf trichomes75. An isolate of *T. koningiopsis* from the stem of a *Vinca* species in Iran was found to produce a range of anti-microbials, including trichodermin, as well as cytotoxic compounds76. *Trichoderma koningiopsis* has also previously been isolated from the rhizosphere of *C. arabica* in Ethiopia15. Here, it is reported for the first time as an endophyte of *Coffea*: all seven isolates being recovered from the leaves and stem of both cultivated and wild *C. canephora* in Cameroon (Table 2), where it appears to be common.

**Trichoderma lentissimum** M.C.H. Rodriguez, H.C. Evans & R.W. Barreto sp. nov. MycoBank: MB832339 (Figs. 8a–c, 11).

**Etyymology:** referring to its slow growth in culture.

**Holotype**: KENYA: Eastern Province, Marsabit National Park, Lake Paradise, primary forest, alt 1340 m; isolated as an endophyte from stem of *Coffea cf. arabica*, 28 January 2015, H.C. Evans & R.W. Barreto. Ex-type culture: COAD 2399. Genbank: TEFI = MK044086; RPB2 = MK044179.

**Culture characteristics—Colonies on PDA:** Optimum growth temperature at 25 °C, 28–30 mm after 72 h, 7 mm at 30 °C; no growth at 35 °C; at 25 °C mycelium mostly on surface, greyish-white aerial, olive-green sporulation, beginning in the colony centre with the formation of concentric rings; no pigmentation in the medium; odour lacking. Colonies on CMD: Optimum growth temperature at 25 °C, 29–31 mm after 72 h, 1 mm at 30 °C, no growth at 35 °C; mycelium mainly hyaline, low with olive-green sporulation, no formation of concentric rings, lacking pigmention and odour. Colonies on SNA: Optimum growth temperature at 25 °C, 30 mm after 72 h, 9 mm at 30 °C, no growth at 35 °C; mycelium hyaline and smooth, low with olive-green sporulation; absence of concentric rings and no pigmentation; forming amorphous and cottony pustules, 1–3.5 mm in diam.

Conidiophores pyramidal with phialides in whorls; phialides lageniform, (4–) 5–9.7 (–10.7) × (1.9–) 2–3 (–3.2) μm, mean 7.6 × 2.7 μm (L/W); supporting cells (4.9–) 5.7–11.9 (–12.5) × (1.6–) 1.7–2.9 (–3) μm, mean 8.1 × 2.1 μm (L/W); conidia globose to broadly ellipsoid, 2.2–3.9 (–4.3) × (1.9–) 2–2.9 (–3.2) μm, mean 2.8 × 2.6 μm, green-coloured, smooth; chlamydospores globose to subglobose, abundant, intercalary and terminal, (5.4–) 6.9–12.3 (–13.5) × (3.6–) 4–7.1 (–10.2) μm, mean 9.4 × 6.6 μm on CMD and PDA.

**Notes:** *Trichoderma lentissimum* is phylogenetically close to *T. gamsii*77 and *T. lieckfeldtiae*78 in the Viride clade. The new species is also morphologically similar to *T. gamsii* in the branching pattern (pyramidal type); the lageniform phialides; and the formation of chlamydospores; but can be separated based on the smaller phialides and conidia of *T. lentissimum* as compared with those of *T. gamsii*. The most prominent differences between *T. lentissimum* and *T. lieckfeldtiae*, is the pachybasium-type branching pattern which is found only in *T. lieckfeldtiae* and the absence of chlamydospores in the latter. This species appears to be rare and it was only isolated once during the survey, as a stem endophyte in Kenya.

**Trichoderma parareesei** Jaklitsch, Druzhin. & Atanasova—in Atanasova et al., *Appl. Environ. Microbiol.* 76:7261, 2010. MycoBank: MB515503.

**Description and illustration.** See1,9.

In a paper of the same year (2010)78, this species is listed as *T. parareesei* nom. prov. and, subsequently, registered as *T. parareesi* sp. nov. Jaklitsch, Druzhinina & Atanasova in MycoBank79. However, the authorities
Material examined. ETHIOPIA: Southern Nations, Nationalities and Peoples Region, Kaffa Zone, Bonga District, Komba Wild Forest Reserve, cloud forest, alt 2000 m; isolated as a mycoparasite of *Hemileia* cf.*coffeicola*, from leaf of *Coffea arabica*, 19 January 2018, H.C. Evans & K. Belachew (culture COAD 2483); Ibid, Maakira, coffee farm, alt 1450 m; isolated as an endophyte from stem of *Coffea arabica*, 21 January 2018, H.C. Evans & K. Belachew (culture COAD 2482); Kaffa Zone, Gesha District, coffee farm, alt 1600 m; isolated as an endophyte from stem of *Coffea arabica*, 22 January 2018, H.C. Evans & K. Belachew (culture COAD 2485).

Notes: *Trichoderma parareesei*, in the *Longibrachiatum* clade, was originally isolated from soil of a subtropical rainforest near Iguazu Falls, Argentina, and is reported to have a pantropical distribution in both rainforest and agricultural soils. During a survey of *Trichoderma* species in the sapwood and dead branches of cacao trees in south-eastern Brazil, it was exclusively isolated from dead wood—where it was the dominant species—and was never recorded from the sapwood. In our study, *T. parareesei* was isolated as an endophyte from stems of *C. arabica* and also as a mycoparasite of *Hemileia* cf.*coffeicola*. This species has never been reported before either as an endophyte in coffee or as a mycoparasite, and probably this is the first record as an endophyte, in general (see). *Trichoderma parareesei* was described as a sympatric, clonal, agamospecies (reproducing only asexually) closely related to *T. reesei* and is its likely ancestor. The latter is a critically important species in the biotechnology industry as a producer of cellulases and hemicellulases and—because of its ability to express recombinant proteins—it is now being targeted for a role in the production of biofuels. Many of these industrial strains have been shown to be *T. parareesei* by Druzhinina et al. They also found that this species is strongly mycoparasitic, compared to *T. reesei*, showing significant antagonism to a range of aerial plant pathogens in dual-culture tests. It has also been demonstrated that *T. parareesei* has biocontrol potential against both fungal and oomycete plant pathogens and, moreover, that it enhances root development and promotes growth, in general, of tomato plants. Finally, it was posited that the ecological niche of *T. parareesei* is not soil but the canopy of tropical forest. Our results confirm their supposition since this species was recorded in stems of wild to semi-wild *C. arabica*, as well

Figure 11. Morphological features characteristic of *Trichoderma lentissimum* sp. nov. (COAD 2399). (a–g) Conidiophores and phialides formed on SNA and CMD. (d, e) Chlamydospores on CMD. (f) Conidia. Bars: (a, b, d–f) = 10 µm; (c, g) = 20 µm.
Trichoderma pseudopyramidale

T. pyramidale spores on CMD and produces a yellow reverse on PDA; and, unlike has larger phialides and conidia, compared to T. pseudopyramidale, whilst the latter species forms chlamydospores close to H. coffeicola (Table 1). The latter species, however, is a pathogen of C. canephora in the lowland tropics of West Africa and thus this high-altitude rust of wild Arabica coffee in Ethiopia is considered to be undescribed.

Trichoderma pseudopyramidale M.C.H. Rodriguez, H.C. Evans & R.W. Barreto sp. nov.—MycoBank: MB832342 (Figs. 8j–l, 12).

Etymology: indicating its phylogenetic proximity to T. pyramidale.

Holotype: ETHIOPIA: Southern Nations, Nationalities and Peoples Region, Kaffa Zone, Bonga District, Komba Wild Forest Reserve, cloud forest, alt 1900 m; isolated as a mycoparasite of Hemileia cf. coffeicola on leaf of Coffea arabica, 26 November 2015, H.C. Evans & R.W. Barreto (culture COAD 2438).

Notes: With the exception of the soil isolate DAOM 165782 (North Carolina), T. petersenii was known previously only from ascospore isolations. However, it has since shown to be common on woody hosts in Southern Europe, especially in Spain, and is reported to be ubiquitous on the Canary Islands, being found in the sexual stage on stromata of the Xylariaceae, presumably as a mycoparasite. In our study, T. petersenii was found for the first time in Africa, and also this is the first report of it as a rust mycoparasite, specifically, on a Hemileia species close to H. coffeicola (Table 1). The latter species, however, is a pathogen of C. canephora in the lowland tropics of West Africa and thus this high-altitude rust of wild Arabica coffee in Ethiopia is considered to be undescribed.

Material examined. ETHIOPIA: Southern Nations, Nationalities and Peoples Region, Kaffa Zone, Bonga District, Maakira-Grugutto, semi-wild coffee farm, alt 1600 m; isolated as an endophyte from leaves and stems of Coffea arabica, 25 November 2015, H.C. Evans & R.W. Barreto. Ex-type culture: COAD 2426.

Culture characteristics—Colonies on PDA: Optimum growth temperature at 25 °C, 43–45 mm after 72 h, 40 mm at 30 °C, 22 mm at 35 °C, covering the plate after 96 h; mycelium cream, concentric rings absent and no odour; yellow pigmentation in the central reverse of plate. Colonies on CMD: Optimum growth temperature at 25 °C and 30 °C, 43–44 mm after 72 h, 44 mm at 30 °C, 25 mm at 35 °C; mycelium hyaline, no sporulation, absence of concentric rings and odour. Colonies on SNA: Optimum growth temperature at 25 °C, 40–42 mm after 72 h, 33 mm at 30 °C, 23 mm at 35 °C, mycelium hyaline and smooth, sporulation sparse; absence of concentric rings and pigmentation; forming amorphous, cottony pustules, measuring 1–3.5 mm in diam, turning yellowish then dark green centrally.

Conidiophores pyramidal- to tree-type; phialides ampulliform to lageniform, usually formed in whorls, (5–) 5.3–8.6 (–9.1) × (1.9–) 2.2–2.9 (–3.2) μm, mean 6.4 × 2.6 μm (L/W), base (1.3–) 1.3–2.3 (–2.5) μm wide; supporting cells (2.5–) 3.3–7.1 (–8.2) × 1.9–2.8 (–3.2) μm, mean 6.0 × 2.5 μm (L/W); conidia globose, sub-globose or ovoid, green, smooth, (1.8–) 2.1–2.9 (2.1–) 2.3–2.9 (–3.2) μm, mean 2.5 × 2.6 μm (L/W); chlamydospores globose to sub-globose (3.21–) 3.2–8.1 (–9.07) × (3.5–) 3.9–7.5 (–8.5) μm, mean 6.0 × 6.1 μm (L/W).

Additional strain examined. CAMEROON: Eastern Province, Somalomo Village, coffee farm, alt 700 m; isolated as a mycoparasite from pustules of Hemileia coffeicola on leaves of Coffea canephora, 22 November 2015, H.C. Evans & R.W. Barreto, COAD 2433. GenBank: TEF1 = MK044131, MK044157; RPB2 = MK044224, MK044250; CAI = MK084870, MK084869.

Notes: Most of the species found in this study could be identified with high support using the combination of tef1 and rpb2 genes. However, for the isolates assigned to T. pseudopyramidale, it was also necessary to include the calmodulin gene in the analysis in order to separate the novel species from the closely related T. pyramidale. Trichoderma pseudopyramidale grouped phylogenetically close to T. pyramidale in the Harzianum clade. The two species share several characteristics in common, such as pyramidal conidiophores, a similar growth rate on PDA and SNA, and the formation of amorphous pustules with white-yellow borders. Trichoderma pyramidale has larger phialides and conidia, compared to T. pseudopyramidale, whilst the latter species forms chlamydospores on CMD and produces a yellow reverse on PDA; and, unlike T. pyramidale, it is able to grow at 35 °C.

Trichoderma pseudopyramidale forms two monophyletic subclades, one containing endophytic isolates and another including isolates obtained directly from CLR pustules (as mycoparasites). Since both subclades come from a phylogenetically well-supported clade by ML, MP and BI, we decided to keep them in a single species and consider them to represent an infra-specific grouping not warranting taxonomic recognition at this stage. No significant differences in morphology or in the growth rates for isolates belonging to these subclades were found. In our study, T. pseudopyramidale represented by far the commonest isolate (35 isolates), the greater majority as an endophyte in both stems and leaves of C. arabica, as well as a mycoparasite of Hemileia cf. coffeicola on wild Arabica coffee in Ethiopia (Tables 1 and 2). There was a single record from Cameroon, as a mycoparasite of Hemileia coffeicola on C. canephora.

Trichoderma spirale

Bissett—Can. J. Bot. 69:2408, 1991. MycoBank: MB359087 Description and illustration, see.

Material examined. CAMEROON: Eastern Province, Somalomo, Dja Forest Reserve, rainforest, alt 700 m; isolated as an endophyte from stems of C. canephora, 21 November 2015, H.C. Evans & R.W. Barreto (cultures COAD 2408; COAD 2404 and COAD 2413).

Notes: Trichoderma spirale, in the Strictipile clade, is a cosmopolitan species. It was isolated for the first time from soil in Thailand, but has since found to be common in sapwood of cacao and other Theobroma spp. It was also the dominant Trichoderma species isolated from roots of Pinus densiflora in South Korea and it was suggested that Trichoderma may be playing a role in stimulating plant colonization by the ectomycorrhizal fungus Tricholoma matsutake. In our study, it was found for the first time as an endophyte of coffee; all three isolates were from stems of wild C. canephora in Cameroon rainforest of the Congo basin.

Trichoderma theobromicola

Samuels & H.C. Evans—in Samuels et al., Mycol. Res. 110:390, 2006. MycoBank: MB356642.
Material examined. CAMEROON: Eastern Province, Somalomo, Dja Forest Reserve, rainforest, 700 m; as an endophyte from stems of Coffea canephora, 21 November 2015, H.C. Evans & R.W. Barreto (cultures COAD 2406; COAD 2407; COAD 2501; COAD 2504; COAD 2412; COAD 2440; COAD 2414; COAD 2589 and COAD 2590).

Notes: Trichoderma theobromicola, in the Viride clade, was found for the first time growing as an endophyte in the trunk of wild cacao in Amazonian Peru and, in subsequent greenhouse studies, it demonstrated promise as an antagonist against frosty pod disease (Moniliophthora roreri) after being inoculated into and re-isolated from seedlings of cacao22. Similarly, an isolate of T. theobromicola from Cola sp. in the Cameroon38 revealed biocontrol potential after it was shown to be parasitic on and reduced the disease incidence of Phytophthora in Capsicum annuum84. All the isolates in our study were from stems of wild C. canephora in Cameroon rainforest (Table 1). This is the first report of this species as an endophyte of coffee.

Trichoderma virens (J.H. Hill, Giddens & A.A. Foster) Arx—Nova Hedwig. Beih. 87: 288, 1987. MycoBank: MB128198.

Synonym: Gliocladium virens J.H. Mill., Giddens & A.A. Foster, Mycologia 49:792, 1957.

Material examined. CAMEROON: South-West Province, Ekonjo, Mt. Etinde, rainforest, alt 700 m; isolated as an endophyte from stem of wild Coffea brevipes, 17 November 2015, H.C. Evans, R.W. Barreto & M.K. Ndacnou (culture COAD 2400).

Notes: Trichoderma virens is a cosmopolitan species, commonly isolated from soil samples, but its sexual morph appears to be rare; having been found only once on dead wood1. In our study, it was isolated from the stem of a wild species of Coffea in Cameroonian rainforest on a single occasion. This is the first record of T. virens as an endophyte of coffee and, seemingly, as an endophyte of aerial plant tissues. Previously, it has been shown to colonize sugar-cane roots; forming dense mycelium in the intercellular spaces85. It was also reported that T. virens secretes proteins to facilitate colonization of maize roots in which plant-host immune responses are suppressed86. Earlier, it was demonstrated that T. virens promotes growth of Arabidopsis by stimulating the

Figure 12. Morphological features characteristic of Trichoderma pseudopyramidale sp. nov. (COAD 2426). (a, d) Stereo microscope images on SNA. (b, c, e, f, i) Conidiophores and phialides formed on SNA. (h) chlamydospores on CMD. (g) Conidia. Bars: (e, c, f, h) = 10 µm; (b, i) = 20 µm.
root system through an auxin-dependent mechanism\(^{87}\). The isolate from our study may have additional mechanisms to colonize woody stems and, perhaps, to form a similar beneficial interaction with its wild coffee host.

**Discussion**

Previous studies have investigated the diversity of endophytic fungi associated with coffee\(^{88–92}\), but these were based on surveys restricted to the Americas and Hawaii, where coffee is an exotic introduced species. The endophytic mycobionta found in these studies is dominated by genera such as *Colletotrichum*, *Fusarium*, *Penicillium*, *Pestalotia* and *Xylaria*. Such assemblages consist mainly of opportunistic endophytes—seemingly, of little biological significance to their hosts\(^{85–89}\)—with *Trichoderma* appearing only infrequently. Only one study involved sampling of all the coffee tissues (leaf, berry, stem, root system)\(^{93}\) and, of the 843 isolates obtained, only four were identified as belonging to the genus *Trichoderma*. Conversely, and in sharp contrast, the *Coffeea* samples from Africa in this study yielded 76 endophytic isolates of *Trichoderma* from the aerial plant tissues of a relatively small sample size, with a highly diverse taxonomic range, including four new species. At this stage, it is not possible to determine whether the new taxa described herein are geographically restricted to Africa or even to coffee. Nevertheless, we find it significant that a far richer diversity of *Trichoderma* was found in association with coffee in its African centre of origin compared to that elsewhere, especially in the Neotropics. We also find particularly relevant the complete absence of endophytic *Trichoderma* species isolates amongst the plethora of fungal isolates obtained from our sampling in semi-wild situations in Brazil. This was entirely unexpected and may indicate the existence of a ‘*Trichoderma* void’ in the coffee endophyte mycobionta outside of Africa.

The occurrence of *Trichoderma* in association with *C. arabica* has been reported previously in Ethiopia\(^{15,94}\), but these studies focused on strains isolated from the rhizosphere and root tissues. The isolates included: *T. harzianum* sensu lato, *T. hamatum*, *T. asperelloides*, *T. spirale*, *T. atroviride*, *T. koningiopsis*, *T. gamsii* and *T. longibrachiatum*. Only three of these taxa were isolated during our study of stems, leaves and berries: namely, *T. hamatum*, *T. spirale* and *T. koningiopsis*. These are cosmopolitan species that are frequently isolated from tropical habitats, especially from soil\(^{95–97}\). Certain *Trichoderma* species were isolated from more than one plant tissue type: *T. koningiopsis* and *T. spirale* from the leaves and stems of *C. canephora*, *T. hamatum* from the stems and berries of *C. arabica*, *T. hamatum*, *T. koningiopsis* and *T. spirale* have also been reported as endophytes in other tropical woody plants, notably cacao and rubber\(^{7}\). Nevertheless, only *T. hamatum* had previously been reported as endophytic in *C. arabica*; occurring in the root system\(^{15,94}\). Apart from the four novel species described here, other *Trichoderma* species were found for the first time as endophytes in coffee: *T. atroviride*, *T. guizhouense*, *T. breve* and *T. theobromicola*. These species were known from other habitats, such as: tropical soils; decaying wood and bark; as mycoparasites; on mushroom compost; in leaf-cutting ant colonies; and as endophytes in *Theobroma* spp. (Malvaceae)\(^{55,56,96,97}\). *Trichoderma guizhouense* has a worldwide distribution, mainly in soil, and had only been reported previously as an endophyte in the endemic woody liana, *Ancistrocladus korupensis*, and in the stems of *Cola* trees in primary forest in south-west Cameroon\(^{4}\). Previously, *T. theobromicola* was known only from South America, and reported to be a common endophyte in sapwood of cacao\(^{62,79}\) whilst *T. breve*, a recently described species isolated from soil, was previously known only from northern China\(^{4,2}\). These two species are new geographical and host records for Africa, but this may simply reflect the poor sampling of *Trichoderma* in the region, particularly for endophytes. The results of the surveys also suggest that many species of *Trichoderma* are either cosmopolitan or pantropical.

Mycoparasitism—the ecological relationship where one fungus parasitizes another\(^{88}\)—has now been reported for a number of species of *Trichoderma*, notably: *T. atroviride*, *T. hamatum*, *T. longibrachiatum*, *T. reesei* and *T. virens*, and it has recently been established that mycoparasitism is an ancestral trait of the genus\(^{19,99}\). Mycoparasitic *Trichoderma* spp. have a wide range of hosts, including true fungi, such as *Alternaria alternata* and *Fusarium* spp., as well as Oomycetes, such as *Pythium ultimum*\(^{68,98}\). However, the species found as mycoparasites of *H. vastatrix* reported here—*T. aggressivum*, *T. andinense*, *T. parareesei*, *T. petersenii* and *T. pseudopyramidale*—are the first in the genus to be reported attacking the *Hemileia* rusts associated with coffee. Three of the species of *Trichoderma* obtained during the surveys are well-known mycoparasites, but were found here only growing as endophytes in coffee, namely: *T. atroviride*, *T. hamatum* and *T. virens*. *Trichoderma pseudopyramidale* may deserve special attention as a potential biocontrol agent of CLR, since it was the most common mycoparasitic species obtained from both Cameroon and Ethiopia (77.8% of total mycoparasites). In Ethiopia this species was commonly associated with a purported new species of *Hemileia* (cf. *H. coffeicola*) on wild *C. arabica* in cloud forest (ca. 2000 m). It was also frequently isolated as an endophyte from the leaves and stems of both semi-wild and wild *C. arabica* in Ethiopia (see Table 1). It may encompass dual roles as an endophytic bodyguard of coffee and also as a contact mycoparasite of CLR.

Mycoparasitic fungi associated with coffee rust have been studied in regions of the world where coffee is not a native species, such as in Mexico\(^{100}\). It is interesting to note that this Mexican survey identified six purported mycoparasites: *Acremonium byssoides*, *Calcarisporium ovalisporum*, *C. arbuscula*, *Fusarium palididoseum*, *Sporothrix guttiformis* and *Verticillium (= *Akanthomyces lecanii*). A more recent publication reporting the results of an investigation in Mexico and Puerto Rico, involving the use of single-molecule DNA sequencing of fungal rRNA gene barcodes to identify putative mycoparasites in pustules of *H. vastatrix*, yielded 15 fungal taxa associated with CLR, none of which belonged to *Trichoderma*\(^{49}\). Information on the ecology of the new *Trichoderma* species described here, and their role in nature, is limited because relatively few strains of each species were isolated during the survey; the exceptions being *T. botryosum* and *T. pseudopyramidale*, which constituted over 60% of the total isolations and seem to have a close association with their *Coffeea* hosts, in both Cameroon and Ethiopia, in wild, semi-wild and cultivated situations.

The aim of the present study was to collect and catalogue endophytes of *Coffeea* species—as well as the mycoparasites of the associated *Hemileia* rusts—in their African centres of origin, as part of a project to screen and
assess these isolates as potential biological control agents of CLR. The target area is Central America where the rust has become a critical constraint to coffee production, as well as causing a socio-economic crisis, over the past decades. The work presented here covers only the taxonomy with some observations on the ecology of the Trichoderma isolates resulting from the surveys in Africa, but these data will be pivotal for selecting candidate biocontrol agents for the potential management of H. vastatrix in the Americas.

The philosophy behind the overall project is based on the concepts of classical biological control and, in the case of CLR, on the Enemy Release Hypothesis which posits that exotic species become invasive and achieve pest status because of increased fitness in the absence of their coevolved natural enemies. One solution to address the problem of invasive alien pests is to source, import and release coevolved natural enemies from the centres or regions of origin of the target species in order to reduce ‘pest’ fitness: the classical biological control strategy. This approach using fungal natural enemies, such as entomopathogens and plant pathogens, has been employed successfully to control invasive alien arthropod pests and weeds, but never against alien plant diseases using mycoparasites. The evidence from our study indicates that there is a guild of Trichoderma species, potentially antagonistic to H. vastatrix in Africa, which could be exploited for biological control of CLR in Central America following the classical approach. There are claims that non-specific, indigenous mycoparasites; notably, Lecanicillium lecanii—now Akanthomyces lecanii—can reduce the impact of CLR in the Americas, but this is not evident based on the continuing rust outbreaks.

Another scenario has been suggested to further explain the invasiveness of alien plant species: the Endophyte-Enemy Release Hypothesis, which posits that alien plants arrive not only without their coevolved natural enemies but also deficient in, or completely lacking, coevolved endophytes, some of which may be acting as symbionts (‘bodyguards’); protecting their hosts against adverse abiotic and biotic factors. Thus, in their absence, exotic crops thrive and alien weeds invade, with no natural enemies reducing plant fitness and fecundity and no bodyguards to ‘pay’ for protection. In crop species, the consequences can be catastrophic when coevolved natural enemies—lacking their own natural enemies, such as mycoparasites (in the case of fungal pathogens)—eventually catch up with their endophyte-deficient plant hosts. Such may be the case with H. vastatrix in Central America—and, of course, this may explain the devastating rust epitaphiastics that destroyed coffee cultivation in Sri Lanka (Ceylon) in the nineteenth century, as well as in all the global regions where the rust has invaded.

Thus, the ideal classical biological control agent for CLR would combine the best of both worlds in the form of an endophytic mycoparasite, and—as our results indicate—the genus Trichoderma contains such candidates. Potentially, these would not only be used to colonize the coffee leaf and parasitize the external rust pustules—as well as to target the invasive, intercellular mycelium of the rust—but also to bolster host defences through induced resistance. There is increasing evidence that, in addition to induced resistance to diseases and pests, endophytic Trichoderma species confer a range of other benefits to their plant hosts, in particular, drought tolerance, resistance to abiotic factors such as salt stress and growth stimulation.

Preliminary data, using Trichoderma isolates from the survey, are showing positive results in the laboratory with evidence of reduction in rust disease severity (Authors, unpublished). Greenhouse screening of four isolates of Trichoderma (COAD 2418, COAD 2417, COAD 2535 and COAD 2439), belonging to T. hamatum and T. pseudopyramidale sp. nov., showed their ability to inhibit the germination of H. vastatrix urediniospores above 70% in vitro. Isolate COAD 2396 (T. atroviride) reduced the severity of the disease to less than 50% of the levels observed in the controls when applied before or simultaneously with H. vastatrix on coffee leaf discs. In addition, an isolate of T. parareesei (COAD 2482) promoted the growth and increased the biomass of tomato roots by 33% and 57%, respectively; whilst others are now showing the ability to increase drought tolerance (Authors, unpublished).

The methodology employed during the survey for the isolation of endophytes has proven to be robust. It has been emphasized previously that endophyte isolation is a method-dependent process and this will determine the quality and quantity of fungi obtained. In our experience, isolating in situ—directly in the field from tree stems—or immediately after collection, eliminates or reduces contamination by many of the opportunistic endophytes and favours the slower-growing, potentially obligate endophytes. This has consistently been demonstrated not only during the present coffee survey in Africa, but also from previous surveys of wild species of Coffea and Hevea in South American rainforests where this approach was pioneered. These surveys resulted not only in the discovery of numerous new Trichoderma taxa—which are still being described—but in many other taxonomic novelties, including new endophytic lineages of Topolyphadium and a new class of Pezizomyxomycota. Moreover, they reveal the paucity of endophytes in cultivated exotic plants—in this instance, cacao and rubber—compared to wild populations of Theobroma and Hevea in natural ecosystems. This has been confirmed during the present survey, when a survey of coffee endophytes in four states of Brazil, failed to isolate any species of Trichoderma, providing compelling evidence that centres of origins or diversity of plants harbour unique guilds of endophytic Trichoderma species—as well as other genera—that could be exploited not only for classical biological control but also as potential reservoirs of novel metabolites.

In conclusion, our surveys in Africa for endophytes and mycoparasites associated with the genus Coffea and with its Hemileia rusts have revealed a highly diverse range of fungi, with many novel species; Trichoderma being just one component. Because of the relatively few countries (3) and localities (18) visited, and the restricted number of host plants sampled, this can only be viewed as a snapshot of the actual diversity of endophytes, as well as of mycoparasites, associated with Coffea in Africa, especially in forest ecosystems. Potentially, in Madagascar, where the diversity of the genus is richer with 59 confirmed species, this still-untapped diversity could be even higher. Loss of forest habitats in Africa and Madagascar means that many of these fungi will go extinct, along with their host plants, before being described. The potential loss of such key antagonists of the CLR fungus—as well as of Coffea germlaspam—should be cause for concern to coffee stakeholders.
Materials and methods
Sampling and isolation. The fungal isolates were all obtained during survey collections in Africa, namely: Kenya (May–June, 2015); Cameroon (November, 2015); Ethiopia (November, 2015; May–June, 2017; January, 2018). In addition, surveys were made in coffee farms in Brazil to compare and contrast the guilds of Tricho-derma present in the native and exotic ranges of coffee. Surveys were undertaken in cooperation with African scientists from local research organizations; notably, IRAD (Institut de Recherche Agricole pour le Developpe-ment), in Cameroon; Jimma University and Ethiopian Institute of Agricultural Research, in Ethiopia. Ad hoc surveys were also undertaken by the local scientists. The surveys were targeted at areas where wild species of Coffea occur and, specifically, where the main species of commerce—Coffea arabica (Kenya and Ethiopia) and Coffea canephora (Cameroon—Congo Basin)—are present in the wild, or are cultivated in semi-wild conditions (Figs. 13a,b, 14a–c, 15a,b). At each selected site, Coffea plants were examined for rust pustules—with particular attention to collecting rust colonies exhibiting mycoparasitism, or appearing to be abnormal (unusual colour, poor sporulation) (Figs. 14d, 15c,d). Specimens were dried in a plant press for later processing in the labora-tory (preliminary identification and isolation). Also, at each site, samples of at least three separate adult plants were collected, consisting of healthy leaves, berries and 3-cm diam or thicker stem sections of each individual, and bagged for examination and processing later the same day. Isolations were made from healthy leaves, stems and berries of C. arabica, C. brevipes, C. canephora and C. eugenioides (Figs. 13c–f, 15e). The isolation protocol followed the procedure described by Evans et al.20 with modifications, and were performed as described below.

1. Stems in situ were thoroughly rubbed with cotton wool soaked in 70% alcohol and, after the alcohol had evaporated, the bark was removed using a flaming knife or machete blade (Fig. 13a). The exposed panel was then cleaned with a scalpel (Swann Morton 10) and the surface further pared with a smaller blade (Swann Morton 11). Nine, triangular slivers of sapwood (ca. 8 × 5 mm) were excised with a scalpel (Swann Morton 10A) from the panel and transferred individually with fine forceps to three plastic Petri plates (3- or 5-cm diam; 3 samples/plate), containing selective media: potato dextrose agar (PDA), one-fifth strength (20% PDA), supplemented with 10 mg/l penicillin–streptomycin solution. These were sealed immediately with electrical tape and stored in plastic boxes. During these procedures, all instruments were surface sterilized in 90% ethanol and flamed using a portable, alcohol burner. On arrival at the laboratory, the plates were transferred to a 25 °C incubator and examined regularly over an 8-week period. Hyphal tips or spores were excised or picked from colonies as they appeared on or around the wood samples and transferred to 5-cm diam, plastic Petri plates containing 20% PDA or potato carrot agar (PCA) and incubated under black light at 25 °C to promote sporulation. This procedure was firstly described for the isolation of endophytic fungi by Evans et al.20 but was applied here for the first time for endophytic fungi from coffee.

2. Young mature healthy leaves (third from the branch tip) were thoroughly rubbed with cotton wool soaked in 70% alcohol and, after the alcohol had evaporated, three small (ca. 5 × 5 mm) square fragments were excised from the leaf centre (including the midrib) and were surface sterilized for 3 min by immersion in 10% bleach, followed by immersion in sterile water in stoppered plastic tubes and, following a thorough agitation, were plated as described for stems. Subsequent processing was as described for stems.

3. Whenever available, Coffea berries were also sampled and treated similarly as described above for leaf samples but, after surface cleaning with alcohol, each fruit was skinned and inner parts were divided into three slices which were then surface sterilized before plating. Further steps followed the same procedure as described above.

For the isolation of mycoparasites, conidia from parasitized rust pustules were selected and picked-off with a sterile needle, using a dissecting microscope, and transferred to PDA plates. The dried samples were processed within 2 weeks of collection after transport to the laboratory in the UK or Brazil. The same endophyte isolation protocol described above was utilized for samples collected at eight localities in four Brazilian states (Espírito Santo, Minas Gerais, São Paulo and Rio de Janeiro). Survey sites closest to those where coffee was sampled in Africa were selected; concentrating on those where coffee plants were growing in semi-wild or forest situations, such as abandoned coffee farms and invasive populations in Atlantic rainforest (Fig. 16).

DNA extraction, PCR amplification and sequencing. Strains were grown in 3-cm diam plates contain-ing 5 mL of potato dextrose broth (PDB) at 25 °C in the dark for 4–5 days. DNA was extracted from the mycelium grown on the surface of the broth. DNA was extracted with the Wizard Genomic DNA Purifica-tion kit (Promega, Madison, EAU) by following the manufacturer’s instructions. The fragments rpb2 (primers fRPB27–r—RPB25F2)211 and tef1 (primers EF2—EF1728M) were amplified for all isolates and additionally cal (primers CAL228–CAL737)212 was amplified for a subset of 12 isolates.

The polymerase chain reaction (PCR) amplifications were performed in a total reaction volume of 12.5 µL, including 0.25 µL of each primer, 1.25 µL of BSA, 6.25 µL of Taq polymerase [including dNTPs], 0.25 µL of genomic DNA [30 ng/µl] 0.25 µL DMSO and 4 µL of sterile ultrapure water. PCR conditions for rpb2 were 95 °C/5 min., followed by 38 cycles at 95 °C/1 min., 58 °C/2 min., 72 °C/2 min. and 72 °C/10 min. For tef1, conditions were 94 °C/2 min., followed by 9 cycles at 94 °C/35 s, 66 °C/55 s, and 35 cycles at 94 °C/35 s, 56 °C/55 s and 72 °C/1 min 30 s. Conditions for cal were 95 °C/8 min., followed by 35 cycles at 95 °C/15 s, 55 °C/20 s, 72 °C/1 min and extension at 72 °C/5 min. PCR products were visualized by Gelred (Thermo Fisher Scientific) staining following electrophoresis of 4 µl of each product in 1% agarose gel. The PCR products were sequenced by Macrogen Inc., South Korea (http://www.macrogen.com).
Phylogenetic analysis. Consensus sequences were assembled from forward and reverse sequencing chromatograms using SeqAssem\textsuperscript{123} tef1, rpb2 and cal contigs of all strains were compared to homologous sequences deposited in NCBI GenBank. Sequences generated in the present study were deposited in the NCBI GenBank database (Table 1) and sequences obtained in other studies were used in our phylogenetic analyses and were retrieved from the NCBI GenBank database (Supplementary Table S1). T. Sequence alignments were performed using MUSCLE implemented in MEGA \textsuperscript{124}. In total, the dataset comprised 324 partial tef1 (sequences 664 pb); 169 partial rpb2 sequences (951 pb) and 25 partial cal sequences (443 pb). Two concatenated trees with tef1 and rpb2 sequences were created, one with taxa of the clades Harzianum (more numerous), Stricpile and Virens, and the other with the rest of the taxa (Figs. 1 and 2); a third concatenated analysis with partial sequences of three genes, tef, rpb2 and cal, was constructed with a subgroup of sequences to clarify the phylogenetic relationships of...
some species within the clade Harzianum (Fig. 3), such trees containing 168 taxa with 2515 characters, 86 taxa with 2422 characters and 25 taxa with 1927 characters, respectively. The concatenated alignments were generated in Sequence matrix v1.8125. Single-gene trees were also generated. Maximum parsimony (MP), Maximum likelihood (ML) and Bayesian Inference (BI) were performed for the concatenated and single-gene trees. Prior to phylogenetic analyses, the most appropriate nucleotide substitution model for each locus was selected using MRMODELTEST v.2126. Nucleotide substitution models in the two-gene concatenated trees were HKY + I + G and SYM + I + G (Figs. 1 and 2), for tef1 and rpb2, respectively. For the three-gene concatenated tree, the models were HKY + I, K80 + I and K80 + G (Fig. 3) for tef, rpb2 and cal, respectively. For all trees the BI and ML analysis were estimated in the CIPRES Science Gateway Platform using Mr. Bayes 3.2.6 and RaxML-HPC v.8, respectively127,128 and MP in MEGA 10. Phylogenetic trees were visualized using FigTree (http://tree.bio.ed.ac.uk/software/figtree/). Phylogenetic species were recognized based on two main previously accepted criteria129 Genealogical Concordance (the clade was present in the majority of the single-locus genealogies, as revealed by a majority-rule consensus tree) and Genealogical Non-discordance (the clade was well supported in the least one single-locus genealogy, as judged both by MP and BI and was not contradicted in any other single-locus genealogy at the same level of support).

Morphological characteristics. The results of the phylogenetic analysis of the assemblage of Trichoderma isolates guided the selection of isolates to be included in the morphological analysis and characterization of novel taxa. One or two isolates of each new taxon were examined. Procedures for morphological observation of Trichoderma followed the protocol established by Samuels and Hebbar5. Macroscopic characteristics of colonies—mycelium colour, radial growth, presence/absence of concentric rings, sporulation “pustules”62, pigmentation and presence/absence of odour—were evaluated on PDA, CMD (Corn-meal Agar) and SNA (Synthetic Nutrient Deficient Agar) after 7 days at 25 °C under a 12-h daily light regime (light provided by two white and one near-UV lamps placed 35 cm above the plates). Rates of growth were evaluated at 72 and 96 h on the three culture media at 25, 30 and 35 °C in the dark. Observations of fungal structures were made using an Olympus BX 51 microscope and were based on slide cultures prepared with colonies of each isolate growing from PDA and CMD blocks130. After 4–5 days of growth at 25 °C under the same light regime described above, the slides were mounted in 3% KOH for observation and illustration. Descriptions included biometric data of phialides, conidia and chlamydospores. Measurements were taken from images generated with a digital camera Olympus Q-Color 3.
Statement. All experimental protocols adopted during this research were approved by the Comissão de Pesquisa do Departamento de Fitopatologia—Universidade Federal de Viçosa.

Informed consent. Informed consent was obtained to publish the names/information/images of all study participants appearing in the publication.

Ethical approval. All experimental protocols were approved by a named institutional and/or licensing committee/s. All methods were carried out in accordance with relevant guidelines and regulations.
Figure 16. Survey sites in semi-wild situations in Brazil. (a) Miraine Ndacnou collecting a sample of mature *Coffea arabica* from the forest reserve Mata do Paraíso (Universidade Federal de Viçosa, state of Minas Gerais, Brazil). Notice the well-established secondary forest (Mata Atlântica) under which the coffee plants exist. (b) Miraine Ndacnou selecting a mature *Coffea arabica* growing in a forest fragment (Mata Atlântica) of the Fazenda Camocim (Pedra Azul, Domingos Martins, state of Espírito Santo, Brazil) for sampling.

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**Author contributions**

All authors contributed to the study conception and design. M.C.H.R., H.C.E. and R.W.B wrote the paper, with inputs from all authors, and carried out the experiments. D.M.M. undertook sample preparation. L.M.A. contributed to the interpretation of the results and data analysis. K.B.B. and M.K.N. facilitated and participated in material collection, isolations and analysis of specimens. All authors provided critical feedback and helped to shape the research.

**Competing interests**

The authors declare no competing interests.

**Additional information**

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