Bacterial endosymbionts protect beneficial soil fungus from nematode attack

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Fungi of the genus Mortierella occur ubiquitously in soils where they play pivotal roles in carbon cycling, xenobiotic degradation, and promoting plant growth. These important fungi are, however, threatened by micropredators such as fungivorous nematodes, and yet little is known about their protective tactics. We report that threatened by these predators, fungi have developed a diverse set of defense strategies. These include the production of toxic proteins and nematocidal natural products, hyphal piercing, trapping, egg parasitism, and endoparasitism (13, 16). Information on defense strategies employed by Mortierella species against nematodes is, however, scarce. It is known that Mortierella globalpina traps nematodes by means of its hyphae and penetrates the nematode’s cuticle. In this way, M. globalpina may protect its host plants from plant-parasitic nematodes (e.g., Meloidogyne chitwoodi) (17). Antinematode activities have been implicated for some Mortierella species (18, 19), including Mortierella alpina [against Meloidogyne javanica or Heterodera sp. (20, 21)], but it is not a general trait of Mortierella (21, 22). Apart from the hyphal trapping strategy, insight into the molecular basis of the antinematode activities of Mortierella is missing. Furthermore, on a more general note, it is remarkable that thus far no Mortierella secondary metabolites have been associated with potential protective roles against nematodes.

Here, we report a so far unknown strategy of a Mortierella species to protect itself from nematode attack. We provide evidence that cytotoxic benzolactones initially isolated from fungal cultures are in fact produced by bacterial endosymbionts that have been overlooked thus far. We also show that the bacteria dwelling in the fungal hyphae protect their host from predatory nematodes.

Significance

Soil is a complex and competitive environment, forcing its inhabitants to develop strategies against competitors, predators, and pathogens. Identifying and understanding the molecular mechanisms has translational value for medicine, ecology, and agriculture. In this study, we show that a member of important soil-dwelling fungi (Mortierella) forms a tight alliance with toxin-producing bacteria (Mycosporum) that live within the fungal hyphae and protect their host from nematode attack. This discovery is relevant since Mortierella species correlate with healthy soils and are used as plant growth-promoting fungi in agriculture. Unraveling an ecological role for fungal endosymbionts in Mortierella, our results contribute to the understanding of a mainspring in fungal–endobacterial symbioses and open the possibility for the development of new biocontrol agents.

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Results and Discussion

*Mortierella* Fungus Harbors Bacterial Endosymbionts Producing Toxic Macrolactones. We reasoned that macrolactones CJ-12,950 and CJ-13,357 (Fig. 1A) (23) from cultures of *Mortierella verticillata* [synonym *Podila verticillata* (24)] could play a role as nematode defense metabolites. Although the initial report on CJ-12,950 and CJ-13,357 only stated that these compounds enhance the expression of the low-density lipoprotein receptor in human hepatocytes (23), they share the macrolactone enamide architecture with structurally related vATPase inhibitors (25, 26). Moreover, the architectures of CJ-12,950 and CJ-13,357 specifically resemble those of *Burkholderia* sp. strain B8 produced necroximes A to D (1 to 4), which proved to be cytotoxic (27). Since only the two-dimensional structures of CJ-12,950 and CJ-13,357 had been reported (23), we assigned their absolute configurations by examining the structural relationships with necroximes C and D. Optical rotation comparison, high-performance liquid chromatography (HPLC)–based coelution experiments and comparison of tandem mass spectrometry (MS/MS) fragmentation indicated that necroxime D (4) is identical to CJ-12,950, and necroxine C (3) is identical to CJ-13,357 (Fig. 1B and SI Appendix, Table S4). These assignments were corroborated by comparison of the NMR spectra of purified metabolites (SI Appendix, Table S9).

Given the bacterial origin of the necroximes (27) and related macrolactones (25, 28–30), we questioned the biosynthetic capability of *M. verticillata* and sought to identify the true producer. Since several *Mortierella* spp. have been reported to live in symbiosis with bacteria (31, 32), we suspected an endosymbiont to be the true source of 3 and 4. Yet, a 2018 report investigating the prevalence of *Burkholderiaceae*-related bacteria within *Mortierella* spp. stated that strain NRRL 6337 was devoid of endosymbionts (32). Nonetheless, we re-examined the same strain for endosymbionts by staining fungal hyphae with the chitin-binding Calcofluor White dye, and tentatively endobacteria with the nucleic acid dye Syto9 Green (Fig. 1C). Fluorescence microscopy revealed the presence of endosymbiotic organisms in *M. verticillata* NRRL 6337 (SI Appendix, Fig. S1).

To identify the observed bacterial endosymbionts, we cut a small piece of fungal mycelium and extracted holobiont DNA, followed by PCR amplification of the 16S ribosomal DNA (rDNA) region using universal primers. Sequencing of the 16S rDNA region (SI Appendix, Table S1) and BLAST analysis indicated that the symbiont of *M. verticillata* NRRL 6337 is a *Mycoavidus* species. Notably, members of this genus have been reported as symbionts of soil-dwelling fungi (31, 32). However, the full genomes of only three *Mycoavidus* cystinexigenus strains from *Mortierella elongata* and *Mortierella parvispora* have been sequenced (31, 33–35). PCR-amplified bacterial 16S rDNA sequences from other *Mortierella* fungi, however, revealed further *Mycoavidus* endosymbionts with three phylogenetically distant clades (*Mortierella*-associated *Burkholderia*-related endosymbionts [MorBRE] groups A to C) (32). Through phylogenetic analysis, we found that the *Mycoavidus* symbiont of *M. verticillata* NRRL 6337 falls into MorBRE group A (Fig. 1D and SI Appendix, Fig. S3) comprising symbionts of *Mortierella humilis*, *Mortierella gamsii*, *Mortierella basiparvispora*, and *M. elongata* (*M. cystinexigenes*). To better understand the occurrence of *Mycoavidus* endosymbionts in *M. verticillata* strains, we investigated five additional *M. verticillata* strains for the presence of endosymbionts. Amplification of the 16S rDNA regions from gDNA of symbionts of these strains revealed a conserved occurrence of *Mycoavidus* endosymbionts in *M. verticillata* strains. Interestingly, these additional endosymbionts all fall into another phylogenetic group together with *Burkholderia* sp. strain B8. Furthermore, analysis of the metabolic profiles of the respective fungi did not show any production of necroximes (Fig. 1D and E). This finding shows that endosymbionts may frequently occur in *Mortierella* and other species of the order Mucorales, but they can be phylogenetically different.

To clarify whether bacterial endosymbionts are the true producers of 3 and 4, we aimed at curing *M. verticillata* NRRL 6337 of its symbiont through the addition of antibiotics (36). Over the course of several months, we subcultivated the fungal strain on agar plates containing kanamycin, ciprofloxacin, or chloramphenicol. During treatment, changes of the fungal growth were noticeable (Fig. 1F). Finally, we confirmed the absence of the symbions by fluorescence staining, microscopic inspection, and PCR analysis (SI Appendix, Figs. S2 and S4). The metabolic profiling of the symbiont-free fungal strain by liquid chromatography (LC) combined with high-resolution electrospray ionization revealed the complete absence of 3 and 4 (Fig. 1B). These findings indicate that *Candidatus* Mycoavidus necroximicus is the true producer of the macrolactones.

*Ca. M. necroximicus* Dedicates 12% of Its Genome to Secondary Metabolism. To gain insight into the symbiont’s biosynthetic potential, with particular focus on the molecular basis of necroxime biosynthesis, we aimed at sequencing the genome of the endosymbiont. Attempts to isolate and cultivate the endosymbiont in the absence of the fungal host, however, proved to be futile. Methods previously used to axenically cultivate similar fungal endobacteria did not enable growth of the endosymbionts (33, 37), indicating a strong dependence of the bacterial symbiont on the host environment. Thus, we sought to enrich the symbiotic bacteria for DNA isolation. Initially, physical disruption of the host’s mycelium resulted in high levels of contamination with fungal DNA, which complicated the assembly of the endosymbiont’s genome. Eventually, we succeeded in retrieving a bacterial cell pellet by filtration and centrifugation of the turbid supernatant of shaking cultures in baffled flasks and isolated the genomic DNA from resuspended bacteria.

The genome of the bacterial endosymbiont was sequenced using a combination of Oxford Nanopore MiniION and Illumina NextSeq sequencing, and both data sets were used to generate a hybrid genome assembly. Of the 118 contigs, a single 2.4 Mb contig of putative bacterial origin was identified through homology searches using the *Mycoavidus*-like 16S rDNA sequence previously amplified from *M. verticillata* NRRL 6337. Following trimming of overlapping ends (suggesting a circular chromosome) the final 2.2 Mb contig was found to contain 1,768 CDS, 6 rRNAs, 42 tRNAs, and a GC content of 50.6% (genome accession number: PRJNA733818). The 16S rDNA sequence of the new strain has 98.82% nucleotide identity to *M. cystinexigenes* B1-EB1 (33). Even so, genomic comparisons showed an average nucleotide identity of only 81.85% across the two genomes. By current standards for molecular species discrimination, the newly identified *Mortierella* endosymbiont should be considered a new species (*Ca. M. necroximicus*) (38, 39).

By comparative genomic analyses, we noted that the genomes of the two endofungal strains AG77 and B2-EB1 isolated from *M. elongata* (33, 35) are 400 to 500 kb larger than the genome of *Ca. M. necroximicus*. Only the genome of strain B2-EB isolated from *M. parvispora* (34) is smaller (~500 kb) than the genome of *Ca. M. necroximicus* (2.2 Mb). When investigating shared protein orthologs, we noted that a core genome encoding 1,164 proteins exists among the four genomes at the 70% identity level (Fig. 2A). However, a further all-versus-all comparison showed B1-EB1 and AG77 to be the most closely related as they share ~75% of their deduced proteome. The B2-EB and *Ca. M. necroximicus* strains are more distantly related to B1-EB1 and AG77, as well as each other, with only a small number of proteins shared exclusively with either B1-EB1 (20 and 17 proteins, respectively) or AG77 (17 and 22 proteins, respectively) (Fig. 2B).

Whereas biosynthetic gene clusters (BGCs) are present in the genomes of all four studied *Mortierella* symbionts, antiSMASH
analysis (40) revealed that the biosynthetic potential for secondary metabolites is by far the greatest in Ca. M. necroximicus (Fig. 2C). Despite the relatively small genome for Mycoavidus standards, ~12% of its protein-encoding capacity is dedicated to natural product biosynthesis. We identified nine nonribosomal peptide synthetase (NRPS) gene clusters, two polyketide synthase (PKS) gene clusters, two hybrid PKS/NRPS gene clusters, and five other BGCs (Fig. 2D). Notably, several large PKS and NRPS gene loci present in the Ca. M. necroximicus genome are absent in the genomes of strains B1-EBT, B2-EB, and AG77 (Fig. 2B). This BGC list includes a cryptic BGC (Mcyst_0009–0017) encoding a PKS/NRPS hybrid that shows high similarity to the necroxime assembly line from *Burkholderia* sp. strain B8 (97% coverage, ~70% amino acid identity), which has been unequivocally
Fig. 2. Comparative genomic analyses of *Mycoavidus* spp. (A) Number of orthologous proteins among the four *Mycoavidus* strains at 70% identity. (B) Circos plot of shared protein orthologs, and secondary metabolite loci (detected by antiSMASH v5) in *Mycoavidus* genomes. Outer blocks (orange, brown, yellow, green) represent genome sizes, while the inner blocks represent genomic positions of secondary metabolite loci. Lines linking the three genomes show position of genes whose proteins are orthologous at 70% identity. Depicted are the genome sequences of *M. cysteinexigens* strains AG77, B1-EB, B2-EB, and Ca. *M. necroximicus* (Ca. M. nec.). (C) Number of gene clusters putatively coding for natural products in *Mycoavidus* spp. detected by antiSMASH and by manual assignment. (D) BGCs and their encoded assembly lines identified from the endofungal Ca. *M. necroximicus* are displayed. A, adenylation; AT, acyltransferase; C, condensation; DH, dehydratase; E, epimerization; Gnat, GCN5-related N-acetyltransferase; KR, ketoreductase; KS, ketosynthase; MT, methyltransferase; OX, oxygenase; TE, thioesterase domains. Acyl carrier (light blue) and peptidyl carrier proteins (dark blue) are shown as circles without designators. (E) Homologous benzolactone BGCs in the genome of *Burkholderia* strain B8 and Ca. *M. necroximicus*. 

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linked to necroxime biosynthesis by targeted gene knockouts (Fig. 2E) (27). The only major difference between the two BGCs is the NRPS gene necA, which is missing in the genome of Ca. M. necrocinus. This finding is in full agreement with the current biosynthetic model, since NecA is responsible for the attachment of the peptide side chain in 1 (Fig. 1A) (27), which is absent in 3 and 4. Furthermore, the architecture of the encoded PKS/NRPS modules is perfectly in line with the biosynthesis of the benzolactone enamide backbone of 3 and 4. Based on these in silico predictions, we inferred that this PKS/NRPS hybrid gene cluster codes for the biosynthesis of 3 and 4 (SI Appendix, Figs. S6 and S8). Together with the metabolic profiling of the cured fungal strain, these data indicate that the bacterial endosymbionts, not the fungus, are the true producers of the benzolactones 3 and 4. CJ-12,950 and CJ-13,357 are thus important additions to the small group of natural products that were believed to be fungal metabolites but are actually produced by bacterial endosymbionts; rhizoxins (41) and rhizoxins (42) from symbionts of Rhizopus microsporus (43), and endolides from Stachylidium bicolor (44). From an ecological viewpoint, it is remarkable that endosymbiotic bacteria were identified as the true producers of the virulence factor of the rice-seedling blight fungus, Mortierella verticillata (36, 37, 45). Given the different ecological context of Mortierella, however, we assumed that the necroximes may have another function in microbial interactions.

**Necroximes Protect the Fungal Host from Nematode Attacks.** To learn more about the potential role of necroximes (3 and 4) in the ecological context of the Mortierella–Mycoavidus symbiosis, we investigated whether these toxins could impair the growth of, or even kill, competitors. Therefore, we considered that the common natural habitat of Mortierella species, including M. verticillata NRRL 6337, is soil, and that microbial survival in the soil environment is not only determined by the capacity to grow under harsh conditions but also by the ability to defend oneself from (micro)predators (46). Among the most abundant fungal predators are nematodes, which share the same soil habitat as Mortierella (47).

To determine if 3 and 4 or any other endobacteria-derived substance have anthelmintic activity, we first performed a viability assay against the model organism, Caenorhabditis elegans (48). We cultured both cured (Mycosividus-free) and symbiotic M. verticillata NRRL 6337 on potato dextrose agar (PDA agar). Cultures were extracted, and each extract was fractionated by preparative HPLC. The individual fractions (F1 to F9) were subsequently tested against C. elegans. Anthelmintic activity in this assay was determined by the ability of C. elegans to feed on a supplied Escherichia coli food source in the presence of the different fractions. Consumption of bacteria indicates unimpeded nematode growth, whereas growth of E. coli indicates that the nematodes are negatively affected by the added substances (Fig. 3A). Notably, all fractions of the extract obtained from the cured strain culture were found to be inactive in the C. elegans assay. In contrast, we observed a marked nematocidal activity of fraction 6 from the extract of the symbiotic fungus. By LC/MS measurements we confirmed the presence of 3 and 4 in the active fraction. In order to determine the anthelmintic potency of the major metabolite 4, we performed the viability assay against C. elegans using increasing concentrations of the pure substance and determined an inhibitory concentration at 50% (IC50) value of 11.3 μg · mL⁻¹ (24.66 μM) (Fig. 3B). Interestingly, the amount of isolated necroximes from fungal cultures grown on agar plates is ∼11 μg · mL⁻¹. Assuming that the actual concentration in fungal hyphae is slightly higher due to an uneven diffusion into the agar and some loss during the purification steps, we conclude that the concentrations inside and around the fungal mycelium are sufficiently high to fully protect it from mycophagous nematodes.

In order to corroborate a potential host-protective role of the symbiont-derived toxin, we next focused on a fungivorous nematode. Therefore, we selected Aphelenchus avenae, a predator using a stylet to feed on fungi, which pierces the fungal cell wall and allows the fungivore to ingest the fungal cytoplasm (49, 50). Sharing the same soil habitat, A. avenae represents a realistic predator of Mortierella spp. (51). To investigate the effects of symbiotic and cured M. verticillata strains on the feeding behavior and survival of A. avenae, we determined the number of animals that were harvested from fungal–nematodal cocultures. In addition, we compared the mobility ratios of the nematodes in correlation to the presence or absence of the bacterial symbiont. As a control, we employed the symbiont-bearing, but necroxime-negative, M. verticillata strain CSB 225.35 (Fig. 1E), thus ruling out an influence solely based on the presence of bacterial symbionts.

To determine nematodal propagation rates, we inoculated plate cultures of symbiotic and cured fungi with A. avenae and cocultivated both organisms for 17 to 24 d (three biological triplicates). Subsequently, nematodes were isolated from the cocultivation plates by Baermann funneling (52), transferred onto water-agar plates, and counted by stereomicroscopic visualization. We found that significantly fewer nematodes are able to grow in the presence of the necroxime-producing endosymbionts (Fig. 3C and SI Appendix, Fig. S9 and Tables S5 and S6).

To scrutinize the effect of the toxin on the fitness of the fungivorous nematodes, we harvested the animals from cocultures and determined their movement—and thus the mobility ratio—by image analysis and mathematical quantification (Fig. 3D). Using stereoscopic time series to track their movement, we compared the area covered by each moving nematode during the time series to the area covered solely by its body without movement, allowing us to differentiate active (living) from inactive (dead or paralyzed) animals. Analyzing a minimum of 176 nematodes from three independent experiments, we observed a significant decrease in the mobility ratio of nematodes grown on symbiotic M. verticillata NRRL 6337 compared to cured NRRL 6337 cultures and to necroxime-negative CSB 225.35 cultures (Fig. 3E and SI Appendix, Fig. S10 and Tables S7 and S8).

HPLC analyses of the plate extracts detected necroximes only in symbiotic cultures of NRRL 6337 but not in cured strains or CBS 225.35, correlating once again the toxins with reduced numbers and lower fitness of the nematodes. To unambiguously assign the nematocidal activity in the propagation assay to the necroximes, we repeated the A. avenae assay with the cured Mortierella strain and chemically complemented the major toxin. Specifically, we overlaid the cured strain NRRL 6337 with solutions of 4 in increasing concentrations (25 μM [IC50], 50 μM, 109 μM, and 219 μM). We then compared A. avenae propagation in necroxime-complemented cultures to untreated cured as well as symbiotic fungi by microscopic examination after two weeks of coincubation. For cultures supplemented with 25 μM or 50 μM of 4, we noted a moderate reduction of nematode propagation, whereas in cultures supplemented with 109 μM or 219 μM of 4 the presence of nematodes in the fungus was abolished (Fig. 3F and SI Appendix, Figs. S11 and S12). The elevated concentrations compared to the IC50 value can be explained due to an uneven distribution of 4 into deeper layers of the hydrophobic fungal colony and the ongoing growth of fungal hyphae, which were not wetted with toxin solution. Nonetheless, these experiments unambiguously verified that the chemical complementation restores the anthelmintic effect. Thus, we uncovered an important role of a natural product in the complex tripartite interplay of symbiont, host, and (micro)predator (Fig. 3G).

**Conclusions.**

In this study, we uncovered a previously overlooked bacterial endosymbiont that protects the important soil-dwelling fungus M. verticillata from a fungivorous nematode. Comparative genomics indicate that the yet unculturable bacterial symbionts belong to a new species that is endowed with a high biosynthetic potential. Through metabolic profiling of the symbiotic wild type
Fig. 3. Nematocidal activity of symbiont-derived toxins. (A) Viability assay of *C. elegans* in presence of extract fractions of symbiotic and cured *M. verticillata* NRRL 6337. HPLC profiles of extracts are shown with corresponding effect on nematodes, measured as effect on the *E. coli* optical density (OD). When nematode growth is impaired by the fraction, *E. coli* cells are not consumed, and thus the OD_{600} is not altered (error bars represent mean of three biological replicates). The red asterisk represents 4. (B) Toxicity screening of 4 against *C. elegans*. The red line marks IC_{50} at 11.3 μg · mL^{-1} (24.66 μM; 95% CI, 21.45 to 28.37 μM; error bars as mean of five biological replicates). (C) Nematode counts from propagation assay of *M. verticillata* and *A. avenae* cocultures. Bars represent relative nematode numbers compared to the mean of the nematode count from cured *M. verticillata* NRRL 6337 cultures. cur., cured; sym., symbiotic. *P < 0.02; ***P < 0.001; ****P < 0.0001. Data represent three biological replicates with three technical replicates each. (D) Workflow of image analysis and mathematical evaluation of *A. avenae* mobility in fungal–nematodal coincubations. Processing of time series is demonstrated by one time frame. Exemplary images of nematodes from two time frames (frame 1 and 26) are shown to illustrate differences in motility. Results of calculated mobility ratios (MR) were used for live or paralyzed/dead categorization. (E) Results of image analysis and mathematical quantification of nematode movement. Bars show ratio between moving/living nematodes and paralyzed/dead nematodes, which were harvested from cocultures of *A. avenae* with symbiotic *M. verticillata* NRRL 6337 cultures, cured NRRL 6337 cultures, or CSB 225.35 cultures. Numbers and error bars were calculated from minimal 176 worms from three biological replicates. (F) Stereomicroscopic images and schematic picture of chemical complementation assay with a magnitude of 25×. Sample of nematodes harvested from plates containing symbiotic, cured, or cured and with 4 chemically complemented *M. verticillata* NRRL 6337 cultures. (G) Schematic summary of tripartite interaction between fungal host, bacterial endosymbiont, and mycophagous nematodes.
and cured aposymbiotic fungi, we provide evidence that the endofungal bacteria are the true producers of highly toxic macro-

lides that were previously believed to be fungal metabolites. Im-

portantly, these compounds (necroximes) efficiently protect the host from nematode attack, as demonstrated by coculture exper-

iments, chemical complementation, and image analyses. Thus, this

work not only reveals an ecological role of endofungal bacteria but also introduces a strategy to ward off micropredators. Conse-

quently, the bacterial biosynthesis of necroximes provides an ad-

vantage of the fungal–bacterial alliance over other aposymbiotic or

necroxime-negative symbiotic M. verticillata strains in the soil niche.

Beyond inspiring the discovery of related tactics in symbioses, our

findings may set the basis for new biocontrol agents, with the

prospect of shielding plant hosts from plant-pathogenic nematodes.

Materials and Methods

Isolation of Natural Products. For 3 and 4 isolation, M. verticillata NRRL 6337 was cultivated on PDA plates (Bacto, BD) at 26 °C. The culture was extracted twice with 1:1 volume of ethyl acetate overnight. The organic phase was concentrated under reduced pressure and the residue was dissolved in methanol. The extracts were prefractionated on an open Sephadex LH-20 column with methanol as eluent. Necroxxime-containing fractions were further purified with a preparative HPLC under following conditions: A, H₂O + 0.01% TFA; B, methanol; and 15 to 100% B in 35 min, 15 mL·min⁻¹ (Phenomenex, Luna, 10 μm, C18(2)), 100 A, 250 x 21.2 mm). NMR analysis was carried out on a 600 Mhz Avance Ultra Shield Bruker), and signals were referenced to the residual solvent signal (DMSO-d₆).

Identification of Endosymbionts in M. verticillata. For the preparation of cultured fungal strains, fungi were continuously subcultivated at 24 °C on PDA plates containing 40 g · mL⁻¹ ciprofloxacin or 50 μg · mL⁻¹ kanamycin for several months. After phenotypic changes were observed by eye, an agar plate of each fungal culture was incubated with 20 mL ethyl acetate and controlled for the presence of ≥ 3 colonies by LC/MS. Final verification of the cured fungal strains was performed by fluorescence staining (Calcofluor White Stain [Sigma] and SYTO 9 Green [Invitrogen]).

Genome Assembly for Ca. M. necrocinus. M. verticillata NRRL 6337 was grown in MM9 medium (53) and orbitally shaken at 160 rpm and 26 °C. The turbid supernatant, containing bacteria from disrupted hyphae, was twice filtered through a membrane (pore diameter, 40 μm) and centrifuged (12,000 x g, 25 °C, 10 min) until a stable pellet occurred. The genomic DNA was extracted with the MasterPure DNA Purification Kit (Epicentre). For long-read sequencing on the MinION platform, DNA quality was evaluated by pulsed-field gel electrophoresis and prepared for sequencing according to the protocol of the Ligation Sequencing Kit (Oxford Nanopore). DNA was loaded onto a single MinION flow cell, and data were collected over a 72-h period. DNA was prepared for sequencing on the Illumina NextSeq platform using the Nextera XT DNA preparation kit (Illumina) with end chemistry and with a targeted sequencing depth of 10. L. Ellegaard-Jensen, J. Aamand, B. B. Kragelund, A. H. Johnsen, S. Rosendahl, Strains

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13. M. J. DiLegge, D. K. Manter, J. M. Vivanco, A novel approach to determine generalist nematode Assays. Liquid assays for active-fraction determination and potency assessment against C. elegans were conducted as previously described (48). For A. avenae coinoculation assay, an aliquot of hyphae of each tested Mortierella strain was transferred to a PDA plate and incubated at 24 °C overnight. Nematodes were sterilized and starved. After one washing step with K-medium, nematodes were resuspended in 300 μL K-medium and aliquots of 50 μL were distributed onto the fresh fungal cultures. Plates were dried and controlled for living nematodes before they were incubated for 17 to 24 d at 20 °C. For the evaluation, nematodes were harvested via Baermann funneling (56). Funneled A. avenae were transferred on 1.5% water-agar plates containing 200 mM genitin and 50 μg · mL⁻¹ kanamycin overnight and subsequently monitored with a Zeiss Axios Zoom.V16 Stereomicroscope for worm count and bioinformatics (https://www.jipipe.org/). Remaining plates were extracted with ethyl acetate to control the metabolite production and processed as described before. For the A. avenae chemical complementation assay, an aliquot of hyphae of the respective fungus was transferred into 12-well plates filled with 1 mL PDA and incubated overnight. The 4 dissolved in 200 μL 50% MeOH was applied and evaporated at room temperature. Nematode suspensions of 50 μL were distributed onto the fungi, dried, and incubated for 14 d at 20 °C. For evaluation, the coculture was removed from the well and washed in 5 mL K-medium overnight. The mixture was filtered through miracloth (Merck) to avoid agar carryover and left at 4 °C for 1 h. The remaining worms were transferred onto 6-well plates containing 5 mL 1.5% water-agar with 200 mM genitin and 50 μg · mL⁻¹ kanamycin. After the plates were dried, the worm count from each plate was assessed with a Zeiss Axios Zoom.V16 Stereomicroscope.

Data Availability. Genome sequence data have been deposited in GenBank (PRUNA733818). The 16S rDNA sequences of the Mortierella endosymbionts were deposited at the NCBI database (BRE_MvertCBS_346.66: MZ330684; BRE_MvertCBS_220.58: MZ330685; BRE_MvertCBS_225.35: MZ330686; BRE_MvertCBS_315.52: MZ330687; BRE_MvertCBS_100561: MZ330688).

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