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Horizontal gene transfer of a bacterial insect toxin gene into the *Epichloë* fungal symbionts of grasses

Karen V. Ambrose¹, Albrecht M. Koppenhöfer² & Faith C. Belanger¹¹Department of Plant Biology and Pathology, Rutgers University, New Brunswick, NJ 08901, ²Department of Entomology, Rutgers University, New Brunswick, NJ 08901.

Horizontal gene transfer is recognized as an important factor in genome evolution, particularly when the newly acquired gene confers a new capability to the recipient species. We identified a gene similar to the *makes caterpillars floppy* (*mcf1* and *mcf2*) insect toxin genes in *Photorhabdus*, bacterial symbionts of nematodes, in the genomes of the *Epichloë* fungi, which are intercellular symbionts of grasses. Infection by *Epichloë* spp. often confers insect resistance to the grass hosts, largely due to the production of fungal alkaloids. A *mcf-like* gene is present in all of the *Epichloë* genome sequences currently available but in no other fungal genomes. This suggests the *Epichloë* genes were derived from a single lineage-specific HGT event. Molecular dating was used to estimate the time of the HGT event at between 7.2 and 58.8 million years ago. The *mcf-like* coding sequence from *Epichloë typhina* subsp. *poae* was cloned and expressed in *Escherichia coli*. *E. coli* cells expressing the Mcf protein were toxic to black cutworms (*Agrotis ipsilon*), whereas *E. coli* cells containing the vector only were non-toxic. These results suggest that the *Epichloë mcf-like* genes may be a component, in addition to the fungal alkaloids, of the insect resistance observed in *Epichloë*-infected grasses.

Horizontal gene transfer (HGT) between species is well established as an important factor in genome evolution^{1,2}. Most examples of HGT involve transfers between prokaryotic species, but the number of examples of HGT into eukaryotes is increasing³. Among the eukaryotes, the chitin cell wall of fungi is considered a structural barrier, likely reducing the frequency of HGT, but even so there are numerous instances of HGT in fungi³. Most such instances are from prokaryotes, but there are some documented fungus-to-fungus transfers. In a systematic examination of 60 fungal genomes, representing four phyla, a total of 713 genes of prokaryotic origin were detected in 53 genomes⁴. The Ascomycota subphylum Pezizomycotina stood out as having a large number of HGT events. Richards et al.³ analyzed the functions of 323 genes confirmed as originating from HGT into fungi, and concluded that HGT played a role in expanding the nutrient acquisition and environmental colonization capacities of many fungi.

The amount of transferred DNA ranges from single genes to secondary metabolite gene clusters and even whole chromosomes⁵. The age of HGT events into fungi is also quite variable. For example, the transfer of a bacterial urea amidolyase gene into fungi was proposed to occur before the divergence of the subphyla Pezizomycotina and Saccharomycotina, approximately 400 million years ago (Mya)⁶. In contrast, transfer of the virulence gene *ToxA* from a fungal pathogen of wheat, *Stagonospora nodorum*, into the fungus *Pyrenophora tritici-repentis*, which resulted in the creation of a new wheat pathogen, was proposed to have occurred less than 75 years ago⁷.

Here we report the presence of an insect toxin gene similar to the bacterial *makes caterpillars floppy* genes (*mcf1* and *mcf2*) of *Photorhabdus luminescens*^{8–10} in the *Epichloë* fungal endophytes of grasses. *Epichloë* are ecologically and commercially important intercellular fungal symbionts of many cool season grass species^{11,12}. The fungi grow in between the plant cells in the above ground parts of the plant (Fig. 1). Infection by the *Epichloë* fungal endophytes often confers resistance to abiotic and biotic stresses¹³. The best understood effect of endophyte infection on the grass hosts is insect resistance, which is largely attributable to the production of toxic alkaloids by the fungi^{14,15}. Of the hundreds of fungal genome sequences available at NCBI, only the *Epichloë* have a *mcf-like* gene, suggesting it is the result of a lineage-specific HGT event from bacteria into a predecessor of the modern *Epichloë*. The gene is expressed and is functional in *E. typhina* subsp. *poae* Ps1, suggesting it may be a component, in addition to the fungal alkaloids, in the observed insect resistance of some endophyte-infected grasses.

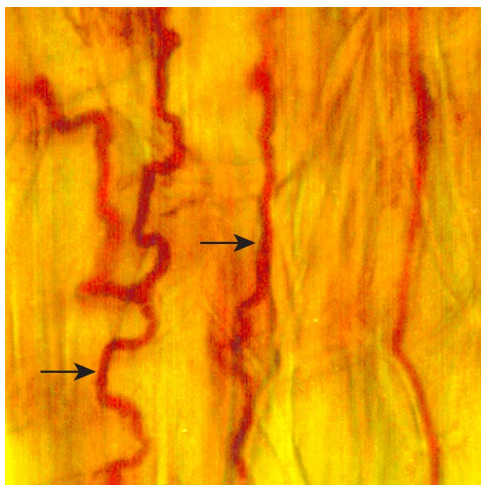


Figure 1 | Leaf sheath epidermal peel of *Poa secunda* subsp. *juncifolia* infected with *Epichloë typhina* subsp. *poae* Ps1 stained with Rose Bengal. Fungal hyphae growing in between the plant cells are indicated by arrows.

Results

Detection of a makes caterpillars floppy-like Gene in *Epichloë* Genomes. Recently we reported a quantitative comparative transcriptome analysis of *E. festucae*-infected versus endophyte-free *Festuca rubra* L. subsp. *rubra* (strong creeping red fescue) using the high throughput sequencing approach of SOLiD-SAGE¹⁶. Strong creeping red fescue is a commercially important grass species and is often naturally infected with the fungal endophyte *E. festucae*^{17,18}. One SOLiD-SAGE tag recovered was from an *E. festucae* gene similar to the insect toxin genes *makes caterpillars floppy 1* and *2* (*mcf1* and *mcf2*) of *Photorhabdus luminescens*^{8,9} and the *fitD* toxin genes of some *Pseudomonas* spp.¹⁹. *Ph. luminescens* is a symbiotic bacterium of entomopathogenic nematodes. When the nematodes infect their insect host, the bacteria are regurgitated into the insect hemolymph where they release several toxins that kill the insect host^{10,20}. The *mcf1* gene was discovered in a screen of *Ph. luminescens* cosmid clones for insect toxicity and the name comes from the effect seen after normally non-toxic *Escherichia coli* expressing the gene were injected into caterpillars⁸. *mcf2*, also from *Ph. luminescens*, and *fitD*, from *Pseudomonas protegens* Pf-5 and CHA0 isolates (previously called *Pseudomonas fluorescens*)²¹ were discovered by their sequence similarity to *mcf1* and have been shown to have a similar effect on caterpillars^{9,22}. *mcf1* has a 1,623 bp N-terminal extension relative to *mcf2* (Fig. 2a). All of these genes encode large proteins (2,388 to 2,996 amino acids) with a TcdA/TcdB pore-forming domain similar to that of *Clostridium difficile* toxin B²³. *Mcf1* was found to kill insects by promoting apoptosis via the mitochondrial pathway^{20,24}. The *Ps. protegens* *FitD* toxin contributes to the oral insecticidal activity of the bacteria²⁵. The mode of action of *Mcf2* is unknown⁹.

The genome sequences of 13 *Epichloë* spp. are publicly available (15; <http://www.endophyte.uky.edu/>) and all contain sequences similar to the *E. festucae* *mcf-like* gene. However, similar sequences are not found in any of the hundreds of non-*Epichloë* fungal genome sequences currently available in NCBI. The uniqueness of the *mcf-like* gene to the *Epichloë* lineage suggests it likely arose from a HGT event from a bacterial donor. The alternative scenario in which the *Epichloë* *mcf-like* genes originated from vertical gene transmission can be rejected since this would imply that the *mcf-like* gene was lost in all fungal lineages except the *Epichloë*. Diagrams of the *Epichloë* *mcf-like* genes are shown in Fig. 2a. The annotated position of the N-terminus of the proteins was variable, but most contained genes with long open-reading frames interrupted by 2 introns. The gene structure of *E. amarillans*, *E. baconii*, *E. brachyelytri*, *E. glyceriae*, *E. mollis*,

and *E. aotearoae* is considered the normal structure, based on our expression analysis of the gene from *E. typhina* subsp. *poae* infecting *Poa secunda* subsp. *juncifolia* (big blue grass), designated *E. typhina* subsp. *poae* Ps1^{26,27} (discussed below). The *E. elymi*, *E. festucae* RC (the Rose City isolate of *E. festucae*), *E. typhina* subsp. *poae* 5819, and *E. gansuensis* isolates had mutations that resulted in frame shifts and early termination codons and so would not be expected to produce an intact *Mcf*-like protein. The *E. gansuensis* var. *inebrians* and *E. typhina* E8 isolates were annotated as having N-terminal extensions relative to most of the other species. In both cases the annotated extension can be attributed to a deletion of the presumed normal coding sequence start site and recognition of upstream open reading frames as exons. The *E. festucae* 2368 and Fl1 isolates had deletions of the 5' region of the gene and were annotated as beginning at a downstream region of the gene. The remarkable amino acid similarity of the predicted *Mcf*-like protein from *E. typhina* subsp. *poae* Ps1 to the bacterial *Mcf1* and *Mcf2* proteins is illustrated in Fig. 2b. The similarity extends throughout the 1,997 amino acid long protein and is particularly striking in the TcdA/TcdB pore forming domain.

The *Epichloë* *mcf-like* coding sequences were more similar in size to those of *Ph. luminescens* *mcf2*, but the overall number of amino acid identities in the region of overlap was similar between the *Epichloë* sequences and *mcf1*, *mcf2*, and *fitD* (Supplementary Table 1). The nearly identical similarity of the *Epichloë* sequences to all three bacterial genes, as well as a preliminary phylogenetic analysis precluded identification of the likely donor in the HGT event (discussed more below).

As is typical of bacterial genes, there are no introns in the *mcf1*, *mcf2*, or *fitD* genes, but there are presumed introns in the *Epichloë* *mcf-like* genes. Since it appears the *Epichloë* *mcf-like* genes originated from a HGT event, the introns must have been acquired subsequent to the HGT. Such intron acquisition has been found in other eukaryotic genes of bacterial origin^{4,28}. Identical intron positions in genes from different recipient species suggest a single HGT event in the oldest member of the lineage, with subsequent vertical inheritance in the more recently diverged species²⁸. All of the *Epichloë* *mcf-like* genes had an intron in the 3' region of the gene at an identical insertion site, although there is sequence variation within the intron sequence itself. Most also had a conserved 5' intron, except the *E. gansuensis* var. *inebrians* and *E. typhina* E8 isolates, both of which had deletions in the 5' sequence, which resulted in annotated N-terminal extensions. The *E. festucae* 2368 and Fl1 isolates, which had truncated 5' regions, also lacked the 5' intron. The two identical intron positions suggest there was a single HGT insertion as well as single intron acquisition events in the original horizontally transferred *mcf-like* gene, which was subsequently vertically inherited throughout the speciation of the *Epichloë*.

Estimation of the Age of the *mcf-like* Gene HGT Event into *Epichloë*. There have been numerous phylogenetic studies of *Epichloë*, but none have directly addressed the order and age of species divergence. To aid in estimation of the age of the *mcf-like* gene HGT event into *Epichloë*, we undertook such an analysis. Most phylogenetic studies of *Epichloë* have been based on β -tubulin (*tubB*), elongation factor EF-1 α (*tefA*), and γ -actin DNA sequence comparisons. In a comprehensive survey that identified optimal fungal genes for use in phylogenetic analyses, these commonly used genes were not among the best²⁹. The two most useful genes were MCM7, a subunit of the minichromosome maintenance complex³⁰, and TSR1, a gene required for synthesis of 40S ribosomal subunits³¹. MCM7 and TSR1 have been found useful in fungal phylogenies that include a wide range of taxa³²⁻³⁴. Other high-performing genes identified were NAD-dependent glutamate dehydrogenase and isoleucine tRNA synthase²⁹.

Maximum parsimony phylogenetic analyses of MCM7, TSR1, NAD-dependent glutamate dehydrogenase, and isoleucine tRNA

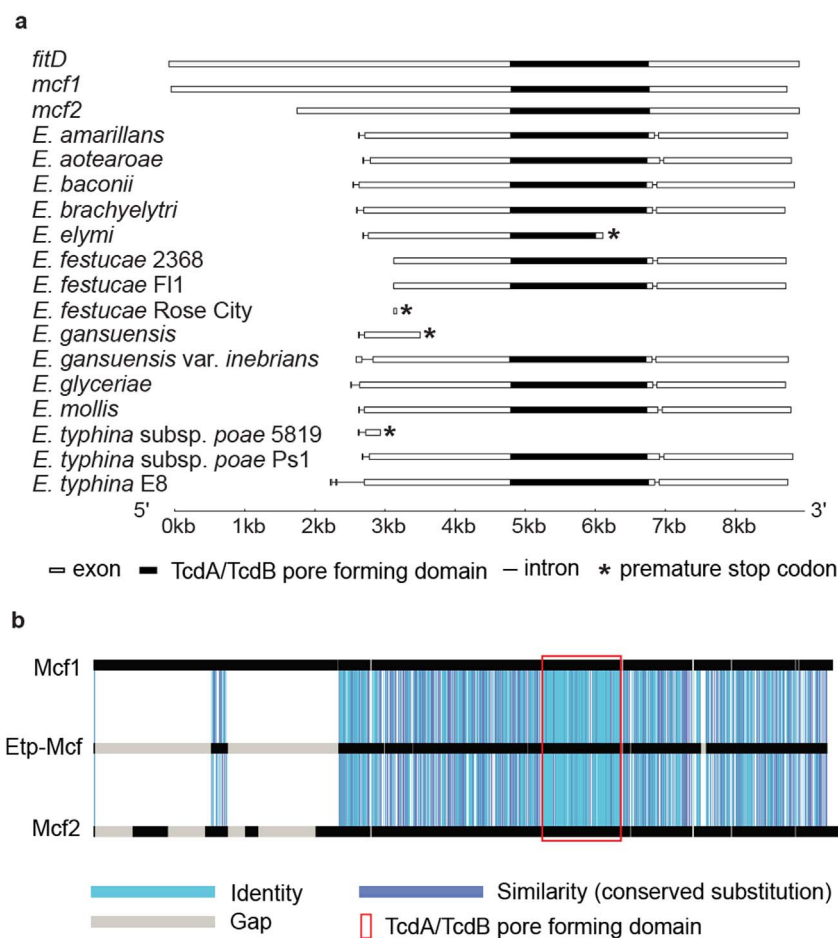


Figure 2 | Gene structure of the *Epichloë mcf-like* genes and amino acid similarity of *E. typhina* subsp. *poae* Ps1-Mcf protein with the bacterial Mcf proteins. (a) Diagrams of gene structure of the bacterial *mcf1*, *mcf2*, *fitD*, and *Epichloë mcf-like* genes. The exons are indicated by boxes and the introns by lines. The conserved TcdA/TcdB pore-forming region is indicated by filled in boxes. The positions of the premature stop codons in some of the *Epichloë* genes are indicated by *. The end of intron 1 of the *E. typhina* subsp. *poae* 5819 sequence was modified from the annotated version to that of the experimentally determined position in *E. typhina* subsp. *poae* Ps1. (b) GECA analysis illustrating the amino acid sequence similarity of the *E. typhina* subsp. *poae* Ps1-Mcf protein (1,997 amino acids; accession KJ502561) to Mcf1 (2,929 amino acids; accession AF503504.2) and Mcf2 (2,388 amino acids; accession AY445665) proteins from *Ph. luminescens*.

synthase sequences from 13 *Epichloë*, as well as some related Clavicipitaceae, are shown in Fig. 3 and Supplementary Fig. 1. Accession numbers of the sequences are given in Supplementary Table 2. To facilitate the alignments for these analyses the intron sequences were removed and only the DNA coding sequences were used. The *Metarhizium anisopliae* sequences were designated as the outgroups for rooting the trees since *M. anisopliae* is in a sister clade, which diverged prior to that of *Epichloë*³⁵. In all of the trees, the two *E. gansuensis* sequences were in a sister clade to the rest of the *Epichloë* spp., which had *E. glyceriae* at its base. Since both *E. gansuensis* and *E. glyceriae* have the *mcf-like* gene in their genomes, the HGT event must therefore have occurred in the common ancestor of these two basal *Epichloë* spp.

To estimate the age of the HGT event we used a molecular clock approach³⁶ similar to that used to estimate the age of polyploidy in *Gossypium* (cotton)³⁷, the divergence times of maize LTR-retrotransposons³⁸, and the age of a segmental duplication in maize³⁹. First we estimated the divergence rate for each gene by calibration to *Atkinsonella hypoxylon* using the formula $T = K_s/2r$. *A. hypoxylon* (synonym: *Balansia hypoxylon*)⁴⁰ is closely related to *B. epichloë*⁴¹. *B. epichloë* was placed in a sister clade to the clade containing *E. typhina* and the *Claviceps* spp. and the divergence of the two clades was estimated to have occurred 81 Mya³⁵. The K_s (substitutions at synonymous sites) and K_a (substitutions at nonsynonymous sites) for

the coding sequence of each gene relative to the *A. hypoxylon* ortholog was determined (Supplementary Table 3). In this analysis glutamate dehydrogenase from *E. gansuensis* var *inebrians* was not included, since the gene sequence was split between two contigs, which introduced a gap into the alignment. The low K_a/K_s ratios for these four genes indicate they are under purifying selection. From the K_s for each gene, the rate of divergence of synonymous sites (r) for each gene was calculated (Supplementary Table 3). The calculated rates varied from 5.93×10^{-9} to 7.72×10^{-9} substitutions per synonymous site per year, which are similar to reported fungal gene divergence rates⁴².

The calculated rates of divergence of each gene were then used to calculate the time of divergence (T) between the *Claviceps* clade and the *Epichloë* clade as well as between *E. glyceriae* and *E. gansuensis* sequences (Supplementary Tables 4 and 5). The mean calculated divergence time between the *Claviceps* clade and the *Epichloë* clade was 58.8 Mya and the mean divergence time between *E. gansuensis* and *E. glyceriae* was 7.2 Mya. These estimates indicate that the single introduction of the *mcf-like* gene into *Epichloë* was sometime after the divergence of the *Claviceps* and *Epichloë* clades at 58.8 Mya and before the divergence of *E. gansuensis* and *E. glyceriae* at 7.2 Mya.

The *Epichloë mcf-like* Gene Donor is Likely a Bacterium Related to *Xenorhabdus* or *Photorhabdus*. Phylogenetic analysis of genes



MCM7

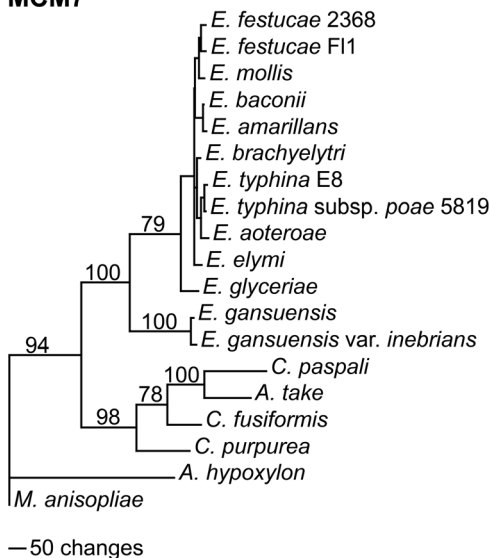


Figure 3 | Rooted 50% majority rule maximum parsimony phylogenetic tree of the MCM7 DNA coding sequences. The *Metarhizium anisopliae* sequences were designated as the outgroup for rooting the tree. The numbers at the nodes are the bootstrap percentages based on 1,000 replications. The tree was based upon 2,472 total characters, of which 1,549 were constant, 343 variable characters were parsimony uninformative, and 580 variable characters were parsimony informative. The unmarked nodes all had bootstrap support of 69 or higher.

derived from a HGT event can often be used to identify the donor^{43,44}. The best matches to the *Epichloë* Mcf-like amino acid sequences were Mcf1 and Mcf2 from *Photorhabdus luminescens*, and FitD from *Pseudomonas* spp. As discussed above, the overall similarity of the *Epichloë* Mcf-like sequences to the bacterial proteins was nearly the same (Supplementary Table 1). Similar Mcf2 proteins were also found in *Xenorhabdus* spp., also bacterial symbionts of entomopathogenic nematodes¹⁰, but have not been functionally characterized. Maximum parsimony phylogenetic analysis of the bacterial Mcf proteins and *Epichloë* mcf-like proteins is shown in Fig. 4. The *Vibrio tubiashii* sequence was chosen to root the tree since its similarity to the *Epichloë* sequences is considerably lower than the *Xenorhabdus*, *Photorhabdus*, and *Pseudomonas* sequences. Similar trees were obtained when using the the neighbor joining and branch and bound methods (Supplementary Fig. 2). The sequence alignment used to generate the tree is shown in Supplementary Fig. 3. Accession numbers of the sequences used are given in Supplementary Table 2.

In the phylogenetic tree the Mcf-like sequences from *E. gansuensis* var. *inebrians* and *E. typhina* subsp. *poae* were placed as basal to the *Xenorhabdus* spp. Mcf2 and *Photorhabdus* spp. Mcf proteins, making it impossible to hypothesize as to a specific bacterial donor. Such a phylogenetic placement suggests the bacterial donor may be an unknown or extinct bacterial species related to extant *Xenorhabdus* or *Photorhabdus* spp. In a phylogenetic comparison of many *Pseudomonas* spp., only one clade had *fitD* genes, and the authors suggested they were acquired through horizontal gene transfer¹⁹. The phylogenetic placement of the *Pseudomonas* *FitD* sequences as well as the overall length similarity to Mcf1 of *Photorhabdus* spp., are consistent with a *Photorhabdus* sp. as the donor to *Pseudomonas*.

Expression and Activity of the *Epichloë* mcf-like Gene. The recovery of a SOLiD-SAGE tag from *Festuca rubra* infected with *E. festucae*¹⁶ supports the expression of the *E. festucae* mcf-like gene. To confirm expression, we amplified and sequenced the mcf-like cDNA from *Festuca rubra* infected with the Rose City isolate of *E. festucae*,

Makes caterpillars floppy-like protein

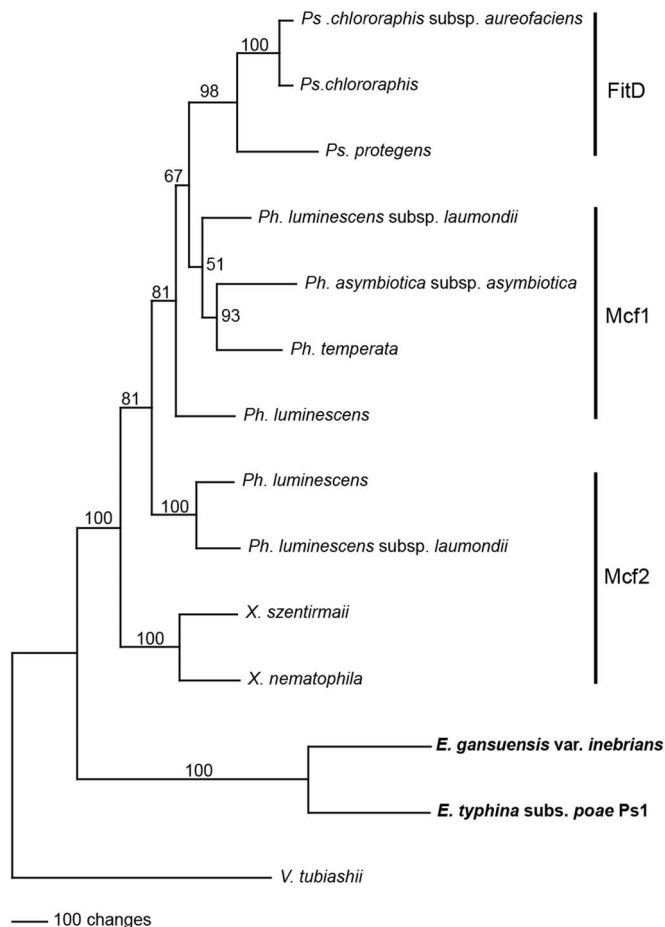


Figure 4 | Rooted 50% majority rule maximum parsimony phylogenetic tree of bacterial Mcf proteins and the Mcf-like protein sequences from *E. gansuensis* var. *inebrians* and *E. typhina* subsp. *poae*. The *Vibrio tubiashii* sequence was designated as the outgroup for rooting the tree. The numbers at the nodes are the bootstrap percentages based on 1,000 replications. The tree was based upon 1,983 total characters, of which 391 were constant, 404 variable characters were parsimony uninformative, and 1,188 variable characters were parsimony informative.

designated *E. festucae* RC. The expression level was extremely low and required three rounds of PCR to obtain enough sample for sequencing. Sequence analysis of the *E. festucae* RC mcf-like cDNA revealed a single base insertion resulting in a frame-shift and an early termination codon, relative to the *E. festucae* 2368 and F11 isolates. The single base insertion was confirmed by sequencing that region of the *E. festucae* RC genome. Although the mcf-like gene is expressed in the *E. festucae* RC isolate, it could not produce a full-length protein.

We also analyzed expression of the mcf-like gene in *E. typhina* subsp. *poae* Ps1, the fungal endophyte infecting *Poa secunda* subsp. *juncifolia*²⁶. The *E. typhina* subsp. *poae* Ps1-mcf cDNA generated from infected plant leaf sheath tissue was easily amplified with one round of PCR (Fig. 5a). Sequence analysis of the amplified cDNA revealed it could produce a full-length protein. The cDNA sequence also confirmed the positions of the two annotated introns. The intron sequences were determined by sequencing of genomic DNA.

Since the mcf-like gene in *E. typhina* subsp. *poae* Ps1 (henceforth designated as *E. typhina* subsp. *poae* Ps1-mcf) was expressed and was predicted to produce a full-length protein, its activity against insects was assessed. The toxicity of the bacterial Mcf proteins was determined by injection of *E. coli* cells expressing the protein into insect



hemocoel^{18,9,22}. The *E. typhina* subsp. *poae* Ps1-*mcf* coding sequence was expressed in *E. coli* by cloning into the expression vector pCold II. This is the same strategy that was used to determine the activity of the *Ps. protegens* FitD protein²². A band of the expected size was seen on SDS-PAGE analysis of proteins from induced *E. coli* cultures (Fig. 5b). The induced protein was found in the insoluble fraction as was reported for the *Ps. protegens* FitD protein²². The toxicity of the induced protein was assayed by injecting black cutworms (*Agrotis ipsilon*) with 50 million *E. coli* cells harboring either the pCold II vector only or cells expressing the *E. typhina* subsp. *poae* Ps1-Mcf protein. The *E. coli* cells containing the vector only were not toxic to the cutworms, whereas injection of the cells expressing the Mcf protein resulted in statistically significant levels of cutworm deaths when compared to the results for the vector only controls (Table 1; Fig. 6). Lower levels of *E. coli* cells (20 million) expressing the *E. typhina* subsp. *poae* Ps1-Mcf protein were not toxic to the cutworms.

Discussion

Here we have reported the characterization of a *mcf-like* gene in the *Epichloë* fungal endophytes of grasses that is the result of a HGT event from a bacterial donor. The gene is present in the genomes of all 13 of the *Epichloë* spp. for which whole genome sequence data is available, as well as 2 additional isolates used in this study, but in no other fungal species. In 11 of the 15 *Epichloë* sequences a long protein, containing the TcdA/TcdB pore-forming region, is predicted and in 4 of the *Epichloë* sequences there was a mutation leading to an early termination codon. In 3 out of these 4 cases, multiple isolates of the same species provided examples of early termination of the *mcf-like* gene and also of a gene capable of producing a potentially functional protein. A similar isolate specific breakdown of some alkaloid biosynthetic genes was seen among *Epichloë* spp.¹⁵. Phylogenetic analysis suggests the donor bacterium may have been an as yet unknown predecessor of extant *Xenorhabdus* or *Photorhabdus* spp. The mechanisms underlying HGT events are not known, but clearly the donor and recipient species must be in close contact. It is easy to imagine that the recipient *Epichloë* sp. could have come in contact

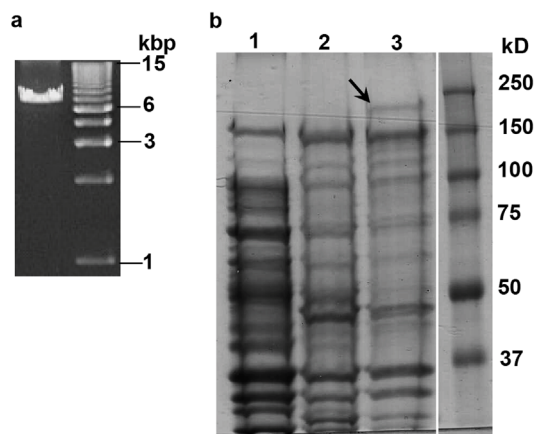


Figure 5 | Expression of *E. typhina* subsp. *poae* Ps1-*mcf* in vivo and in *E. coli*. (a) PCR product of the *E. typhina* subsp. *poae* Ps1-*mcf* transcript using infected *Poa secunda* subsp. *juncifolia* plant cDNA generated from oligo(dT) as template. (b) SDS-PAGE analysis of insoluble *E. coli* proteins. Lane 1: *E. coli* containing the pCold II vector only subjected to induction conditions. Lane 2: *E. coli* containing the pCold II::*E. typhina* subsp. *poae* Ps1-Mcf with no induction of protein expression. Lane 3: *E. coli* containing the pCold II::*E. typhina* subsp. *poae* Ps1-Mcf subjected to induction conditions. Arrow indicates presence of the induced protein at the expected size of 223 kD.

Table 1 | Cumulative numbers of black cutworm larvae deaths after each treatment. Insect assays were performed by injecting either sterile water, 5×10^7 total induced *E. coli* cells containing the pCold II vector only as a negative control or the Etp-Mcf::pCold II plasmid in a total volume of 5 μ L. The experiments also included larvae that were untreated

Black cutworm mortality assay				
Experiment 1 (n = 6; P = 0.001)				
Treatment/Time	24 h	48 h		
Untreated	0	0		
Water	0	0		
pCold II vector only:: <i>E. coli</i>	0	0		
Etp-Mcf::pCold II:: <i>E. coli</i>	3	6		
Experiment 2 (n = 10; P = 0.029)				
Treatment/Time	24 h	48 h	72 h	120 h
Untreated	0	0	0	0
Water	0	0	0	0
pCold II vector only:: <i>E. coli</i>	0	0	1	1
Etp-Mcf::pCold II:: <i>E. coli</i>	1	1	5	6

with the bacterial donor either in soil or in association with a plant host.

HGT events are of particular significance when the newly acquired gene confers a new capability to the recipient species. A recent example is the transfer of a bacterial mannanase gene to the insect *Hypothenemus hampei*, the coffee berry borer beetle⁴³. The acquisition of the mannanase gene was considered to have led to the adaptation of the insect to a new host, the coffee berry, which is rich in mannans. Here we have demonstrated that injection of *E. coli* expressing the *E. typhina* subsp. *poae* Ps1-*mcf* gene into black cutworms (*Agrotis ipsilon*), a pest of various crops, results in death of the insect, similar to the insecticidal activities of the bacterial *mcf1*, *mcf2*, and *fitD* genes^{8,9,22}. The bacterial Mcf proteins are considered critical components of the overall insect toxicity of the bacteria. The *E. typhina* subsp. *poae* Ps1-Mcf protein has been maintained as an active protein over millions of years since its acquisition by an ancestral *Epichloë* sp. This suggests it may also be a component, in addition to alkaloids that are well-established as having insecticidal activity^{15,45}, in the insect resistance observed in plants containing this endophyte isolate⁴⁶. Other *Epichloë* spp. have genes apparently capable of producing Mcf-like proteins that may also have activity against insects, as shown here for the *E. typhina* subsp. *poae* Ps1 isolate.

Bacterial toxin genes of the aerolysin family (unrelated to the *mcf* toxin genes) were found to have undergone recurrent HGT to many diverse eukaryotes, possibly due to their adaptive value⁴⁷. Moran et al.⁴⁷ proposed that genes derived from HGT are more likely to be maintained if they can function alone and are of immediate benefit

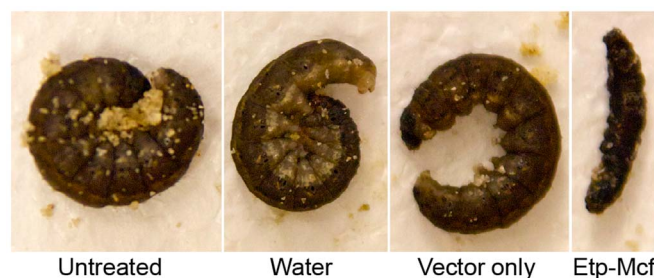


Figure 6 | Toxicity of the *E. typhina* subsp. *poae* Ps1-Mcf protein to black cutworms. Fourth instar cutworm larvae were injected with water, 50 million *E. coli* cells containing the pCold II vector only, or 50 million *E. coli* cells expressing the *E. typhina* subsp. *poae* Ps1-Mcf protein. The live cutworms curl in response to prodding.



to the organism. These characteristics apply to the *Epichloë mcf* genes, also derived from HGT, since the genes may contribute to the insect toxicity of *Epichloë*-infected plants.

Based on phylogenetic analyses and divergence time estimates, the single HGT event into *Epichloë* was estimated to have occurred sometime between 7.2 and 58.8 Mya. Dating of fungal evolution is problematic due to the minimal fossil record, so as more fungal calibration points become available it will be possible to refine the divergence time estimates presented here. The rooted phylogenetic trees generated in these analyses, based on genes established as phylogenetically high-performing genes²⁹, are also revealing regarding the evolutionary history of *Epichloë*. In all four phylogenetic trees the *Epichloë* spp. separated into two clades, one comprised of the two *E. gansuensis* isolates and the other comprised of the rest of the *Epichloë* spp., with *E. glyceriae* at its base. The phylogenetic analyses presented here suggest that an as yet unknown (or possibly extinct) species was the predecessor of *E. gansuensis* and *E. glyceriae* and that the rest of the currently known *Epichloë* spp. were derived from *E. glyceriae*.

It has been proposed that *Epichloë* spp. evolved through coevolution with their grass hosts⁴⁸. The two most basal members of this genus, *E. gansuensis* (host grass *Achnatherum inebrians*, tribe Stipeae) and *E. glyceriae* (host grass *Glyceria striata*, tribe Meliceae) are symbionts of ancient tribes of the grass subfamily Pooideae. However, *E. brachyelytri*, the symbiont of *Brachyelytrum erectum*, which is in the most ancient Pooideae tribe Brachyelytreae^{49,50}, was phylogenetically closer to the other *Epichloë* spp. than to *E. gansuensis* or *E. glyceriae*. These results indicate that the speciation of *Epichloë* was not exclusively through coevolution with their hosts since, in an exclusive model of coevolution, *E. brachyelytri* would be expected to be the most basal *Epichloë* sp. The phylogenetic trees presented here suggest that *E. brachyelytri* evolved via cross species transfer of an *Epichloë* sp. infecting a more recently evolved host grass to the ancient plant species *B. erectum*. Jackson⁵¹ has also concluded that the *Epichloë*-host phylogenies are incongruent and do not support codivergence as a characteristic of the association. Lemaire et al.⁵², proposed host specificity without co-speciation for the *Burkholderia* bacterial leaf endosymbionts. The *Epichloë* – grass symbiosis may be a similar case.

Methods

Plant and Fungal Samples. *Epichloë* spp.-infected plants were grown by clonal propagation and maintained in 15 cm pots in the greenhouse. Each *Epichloë* sp. was isolated from its host plant by plating surface-sterilized leaf sheath tissue on potato dextrose agar (Difco Laboratories, Detroit, MI).

DNA and RNA Isolation. Genomic DNA of *Epichloë* spp. was extracted from culture grown in potato dextrose broth for 14 days on a shaker (175 rpm) at room temperature (23–25°C). The DNA was isolated as previously described⁵³. RNA was obtained from the innermost leaf sheath tissues of *Epichloë*-infected plants. RNA isolation was as previously described¹⁶. Nucleic acid concentration was measured using a Nanodrop ND-1000 Spectrophotometer (Thermo Fisher Scientific, Waltham, MA).

Amplification and Sequencing of *Epichloë mcf*-like Genes and cDNAs. Initial screening to determine the presence of the *mcf*-like gene in *E. festucae* RC and *E. typhina* subsp. *poae* Ps1 fungal isolates was done using fungal genomic DNA. Upon detection of the gene, first-strand cDNA from 5 µg *Epichloë*-infected plant total RNA was synthesized by using SuperScriptTM III Reverse Transcriptase (Life Technologies, Carlsbad, CA) and 500 ng of oligo(dT)₁₈ primer according to the manufacturer's instructions. PCR was performed in 50 µL with either 0.5 µg of fungal genomic DNA or 1 µL *Epichloë*-infected plant cDNA generated from oligo(dT)₁₈, 1.25 units of PrimeSTAR HS DNA Polymerase, 1X PrimeSTAR Buffer with Mg²⁺ (TaKaRa Bio Inc., Shiga, Japan), 200 µM each dNTP, and 0.3 µM of each forward and reverse primer (Integrated DNA Technologies, Inc., Coralville, IA). Two-step PCR was performed by template denaturation at 98°C for 10 seconds followed by 6 minutes extension at 68°C for 30 cycles in a GeneAmp 9700 thermal cycler (Applied Biosystems, Inc., Foster City, CA). The amplification products were visualized on a 1% TBE agarose gel. The amplified genomic and cDNA PCR fragments were purified by using 0.5X Agencourt AMPure XP (Beckman Coulter, Inc., Brea, CA) to remove any fragments under 1 kb and then sequenced directly (Genewiz, Inc., South Plainfield, NJ). For each sequencing reaction, about 800 ng of purified PCR product in 10 µL was treated with 2 µL ExoSAP-IT (USB Corp., Cleveland, OH) to remove

unincorporated primers and excess dNTPs. The ExoSAP-IT reaction was performed at 37°C for 15 min followed by heating at 80°C for 15 min to inactivate the enzymes.

Expression of the *E. typhina* subsp. *poae* Ps1-*mcf* Gene in *E. coli*. To test the anti-insect activity of the *E. typhina* subsp. *poae* Ps1-Mcf protein, the cDNA was cloned into the expression vector pCold II (TaKaRa Bio). The *E. typhina* subsp. *poae* Ps1 cDNA was amplified by PCR as described above with oligonucleotides that introduced a *KpnI* site at the 5' end and an *EcoRI* site at the 3' end. The sequences of the oligos are: forward 5'-GATATAACCATGGCTCACAACT-3' and reverse 5'-TCAACTGAATTCCTACTGATTTCCAGC-3'. Two hundred ng of the purified PCR product was digested with the restriction enzyme *KpnI* (TaKaRa Bio), purified with 0.35X Agencourt AMPure XP (Beckman Coulter, Inc) to remove any fragments under 1.5 kb, digested with *EcoRI*, and again purified with 0.35X AMPure XP. The expression vector pCold II was similarly digested with *KpnI* and *EcoRI*. The digested pCold II plasmid was treated with shrimp alkaline phosphatase (Affymetrix, Santa Clara, CA) to prevent vector religation. Overnight ligation of the digested *E. typhina* subsp. *poae* Ps1-*mcf* PCR fragment and pCold II plasmid was done at a 3:1 insert:vector molar ratio using T4 DNA ligase (New England Biolabs, Inc., Ipswich, MA).

Two µL of the ligation product was used to transform 20 µL *E. coli* DH5α electro-competent cells. The transformed cells were incubated in SOC medium for 1 hour at 37°C, followed by overnight growth of cells on LB medium supplemented with 100 µg/mL ampicillin. Transformed bacterial colonies were screened for plasmids containing the *E. typhina* subsp. *poae* Ps1-*mcf* gene insert by using PCR as described above. Selected clones were grown in 3 mL of LB + ampicillin broth overnight followed by plasmid purification using QIAprep Spin Miniprep Kit (Qiagen, Valencia, CA). DNA sequencing of the plasmids were done as described above. A plasmid containing the correct sequence was transformed into *E. coli* BL21-CodonPlus (DE)-RIPL competent cells (Agilent Technologies, Santa Clara, CA). For use as a control, a transformant containing the pCold II vector only was also generated.

For expression of the *E. typhina* subsp. *poae* Ps1-Mcf protein, the BL21-CodonPlus (DE3)-RIPL transformant was induced by cold shock and isopropyl-β-D-thiogalactopyranoside (IPTG). We detected loss of *E. typhina* subsp. *poae* Ps1-*mcf*:pCold II recombinant plasmid in *E. coli* cells during the course of long induction periods. This was not observed in *E. coli* cells containing the pCold II vector only. Previous studies have shown that carbenicillin and ampicillin, even in high concentrations, quickly lose their ability to maintain selective pressure due to saturation of the media with the antibiotic degrading enzyme β-lactamase^{54,55}. Extra metabolic burden placed on the bacterial cells to maintain the large recombinant plasmid combined with the steady decrease of antibiotic in culture may have contributed to the plasmid loss that we observed. In order to circumvent this problem, the culture supernatant was replaced with fresh medium at regular intervals.

To induce expression a 10 mL culture of the cells was grown for 15 h at 30°C in LB and 200 µg/mL carbenicillin. A 400 µL aliquot of the culture was pelleted, resuspended in 400 µL, added to 10 mL of fresh LB plus carbenicillin and incubated at 30°C to an OD of 0.4–0.5. The culture was then cold-shocked for 15 minutes on ice and subsequently pelleted in a Model J2-21 centrifuge (Beckman Coulter, Inc., Brea, CA) set to 5°C to obtain the bacterial cells. Fresh LB broth supplemented with carbenicillin was used to resuspend the bacterial pellets. To induce protein expression in the culture, IPTG was added to 0.1 mM and the culture was then incubated at 15°C with rotational shaking for 21–24 h. The bacterial cells were pelleted and resuspended in fresh LB and carbenicillin every 3 h for the initial 6 h and again after 12 h to prevent β-lactamase saturation in the culture supernatant. The transformant containing pCold II vector only was treated the same way.

For sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) analysis, 1 ml each of *E. coli* cells expressing the *E. typhina* subsp. *poae* Ps1-Mcf protein and the vector only control cultures were pelleted and lysed using 1X FastBreak Cell Lysis Reagent (Promega, Madison, WI) following the manufacturer's protocol. Following lysis, samples were centrifuged at 10,000 RPM for 15 minutes and the supernatants removed. The proteins remaining in the pellet were solubilized by adding 100 µL of 2X SDS sample buffer⁵⁶, boiled for 5 minutes, and 30 µL was then subjected to electrophoresis on a 10% gel.

Insect Assays to Determine Toxicity of *E. typhina* subsp. *poae* Ps1-Mcf Protein.

BL21-CodonPlus (DE3)-RIPL cells expressing the *E. typhina* subsp. *poae* Ps1-Mcf:pCold II recombinant insert and vector only control were prepared as described above. Cells were pelleted and resuspended in sterile water. Black cutworm (*Agrotis ipsilon* Hufnagel), eggs were obtained from Benzoin Research Inc. (Carlisle, PA) and reared on insect diet (Southland Products Inc. Lake Village, AR) at 27°C and 14:10 (L:D) hours to fourth instars. Immediately before injection the larvae were surface sterilized in 0.8% sodium hypochlorite for 30 seconds and then rinsed in sterile distilled water. Six to ten fourth instars were injected with 5 × 10⁷ *E. coli* cells containing either the *E. typhina* subsp. *poae* Ps1-Mcf:pCold II induced protein or the pCold II vector only control. Additional controls were water injections and untreated samples. The assay was conducted twice. Injections were in a total volume of 5 µL using a Hamilton microsyringe with a 30-gauge needle. The larvae were then placed individually in 30 mL plastic cups with 10 g of moist (10% w/v) pasteurized (3 h at 72°C) sand with about 1 cm² diet and incubated at room temperature (22–24°C). Mortality was assessed at 24, 48, 72, and 120 h after injection. Larvae were considered dead when they no longer reacted to being poked repeatedly with a probe. The data at



the final time points were analyzed by Fisher's exact test to compare the toxin-expressing samples to the vector only controls. Significance was evaluated at $P < 0.05$.

Schematic Representation of *mcf* Gene Structure. Schematic representation of the *mcf* gene structures was done using Gene Structure Display Server (57; <http://gsds.cbi.pku.edu.cn/index.php?input=site>). Illustrative comparison of amino acid similarity for Mcf1, Mcf2 and *E. typhina* subsp. *poae* Ps1-Mcf protein was performed using Gene Evolution and Conservation Analysis (GECA) (58; http://peroxibase.toulouse.inra.fr/tools/geca_input_demo.php).

Phylogenetic Analysis. To identify genes in addition to MCM7 and TSR1 for use in the phylogenetic analyses, the FUNYBASE genome database (<http://genome.jouy.inra.fr/funybase/>)⁵⁹ was searched for single copy orthologs identified as phylogenetically high-performing genes by Aguilera et al.²⁹. Two genes with topological scores of 96 or above and that also had significant sequence polymorphism within *Epichloë* were identified, FG570 (NAD-dependent glutamate dehydrogenase) and MS444 (isoleucine tRNA synthase). DNA sequences for these genes were obtained from the Genome Project at the University of Kentucky website (<http://www.endophyte.uky.edu/>) or from NCBI.

The Clustal-X program⁶⁰ was used to align the DNA coding sequences. The alignments generated by Clustal-X were modified manually to minimize gaps. The phylogenetic analyses were performed with the PAUP* program, version 4.0b10 for Macintosh. The phylogenetic analyses were done by using the maximum parsimony full heuristic search option set to random sequence addition, tree-bisection-reconnection (TBR) branch swapping, and Multrees on, with 1,000 bootstrap replications. Gaps were treated as missing data.

The K_a and K_d values based on pairwise comparison between species were determined by using the MEGA5.2 program with the Nei-Gojobori, Jukes-Cantor model⁶¹.

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Author contributions

K.V.A., A.M.K. and F.C.B. conceived and designed the experiments; K.V.A., A.M.K. and F.C.B. performed the experiments; A.M.K. conducted the insect injections; K.V.A., A.M.K. and F.C.B. analysed and interpreted results. All the authors discussed the results and wrote the manuscript.

Additional information

Data deposition: Sequence data for the *E. typhina* subsp. *poae* Ps1-*mc*f gene has been deposited in the GenBank database (<http://www.ncbi.nlm.nih.gov/>) under accession number KJ502561.

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