Interdomain lateral gene transfer of an essential ferrochelatase gene in human parasitic nematodes

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Lateral gene transfer events between bacteria and animals highlight an avenue for evolutionary genomic loss/gain of function. Herein, we report functional lateral gene transfer in animal parasitic nematodes. Members of the Nematoda are heme auxotrophs, lacking the ability to synthesize heme; however, the human filarial parasite Brugia malayi has acquired a bacterial gene encoding ferrochelatase (BmFeCH), the terminal step in heme biosynthesis. BmFeCH, encoded by a 9-exon gene, is a mitochondrial-targeted, functional ferrochelatase based on enzyme assays, complementation, and inhibitor studies. Homologs have been identified in several filariae and a nonfilarial nematode. RNAi and ex vivo inhibitor experiments indicate that BmFeCH is essential for viability, validating it as a potential target for filariasis control.

The phylum Nematoda is estimated to comprise up to one hundred million species (1), including free-living worms (such as the widely used experimental organism, Caenorhabditis elegans) as well as parasitic species that threaten the health of agriculturally important plants, wildlife, domesticated animals, and humans. Comparative genomics suggests that lateral gene transfer (LGT) may have played an important role in the evolution of plant parasitism by facilitating adoption of a parasitic lifestyle (2, 3). For animal parasitic nematodes, such as the human filarial nematode Brugia malayi, several key metabolic processes have been lost (4). For example, the absence of heme biosynthesis appears to be a peculiar feature of the Phylum Nematoda (5). In filarial nematodes such as B. malayi, essential compounds may be provided by the obligate endosymbiotic bacteria Wolbachia (6). LGT events have been documented between the Wolbachia endosymbiont and its nematode hosts, but to date no functionality has been determined for the transferred DNA (7–9). Here, we describe both the functionality and essentiality of a bacterial gene acquired by filarial nematodes via LGT: that of the last step of heme biosynthesis [ferrochelatase (FeCH)] (Fig. S1). Because of its essentially, FeCH represents a potential drug target against the causative agents of lymphatic filariasis and onchocerciasis (river blindness). These diseases affect more than 150 million individuals with over 1 billion people at risk for infection (10, 11). No new classes of drugs directly targeting filariae with good efficacy and safety profiles have been developed in over 20 y, and suboptimal response/emerging resistance to currently used antifilarial drugs is apparent (12, 13).

Results

BmFeCH Origin and Features. The draft genomic sequence of B. malayi (4) revealed two adjacent ORFs (Bm1_14315 and Bm1_14320) that appeared to be components of the same ferrochelatase (FeCH) gene. We established that these ORFs are, in fact, part of a single transcript with strong similarity to FeCH genes in the GenBank database. Genome annotation did not identify any additional heme biosynthesis genes, and, even when using low stringency search/match parameters, we were unable to identify any other genes in this pathway. PCR using generic primers based on conserved regions of other heme biosynthesis pathway genes also failed to give any specific products.

The B. malayi ferrochelatase protein (BmFeCH) clusters with a subset of α-proteobacterial sequences within the Orders Rhizobiales and Rhodobacterales with high probability but is more distantly related to orthologs in Wolbachia or mammals, as shown by a consensus tree produced from Bayesian phylogenetic analysis of aligned FeCH protein sequences (Fig. 1, Fig. S2). BmFeCH also clusters closely with other nematode FeCH sequences. Evolutionarily, FeCH proteins are poorly conserved across diverse taxa (14); however, all, including the predicted BmFeCH, have invariant residues in the active site pocket. BmFeCH does not possess the C-terminal extension found in eukaryotic counterparts or the signature for the [Fe₂-2S] cluster found in Actinobacteria and metazoan FeCHs (Fig. S2). It does possess the proposed membrane-binding loop absent in Actinobacteria and Firmicutes (14).

Despite its apparent bacterial origin, sequence of the ~4.5-kb BmFeCH gene revealed a eukaryotic gene organization, containing nine putative exons and eight introns. The deduced protein includes an N-terminal extension (encoded by part of the first exon), predicted to include a mitochondrial targeting domain (Fig. S2). PCR was used to amplify full-length FeCH cDNAs from various filarial species: B. malayi, Dirofilaria immitis, Onchocerca volvulus, and Acanthocheilonema viteae. The latter lacks the Wolbachia endosymbiont, but harbored it in its evolutionary past (9). Genomic databases support the occurrence of FeCH in other filariae: Wuchereria bancrofti, Loa loa and Litomosoides sigmodontis, as well as the nonfilarial nematodes, Strongyloides ratti and Strongyloides venezuelensis (this study and refs. 15 and 16). All nematode homologs have an N-terminal extension that is not present in bacterial FeCH enzymes. The presence of FeCH homologs in filarial nematodes (Spurunira, clade III) as well as in nonfilarial worms of the genus Strongyloides (Tylencehina, clade IV) suggests that acquisition of FeCH predated the divergence of clades III and IV or that independent acquisitions occurred in the lineages leading to the Onchocercidae and Strongyloidae (15). We note that the nine-exon gene structure is conserved in all six filarial genera for which draft/on-going genomic sequence data are available (4) (www.nematodes.org/nematodegenomes/index.php/Main Page). Although S. ratti (www.sanger.ac.uk/resources/software/blast/).


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Data deposition: The FeCH cDNA sequences reported in this paper have been deposited in the GenBank database (accession nos. GQ895739 (Brogia malyai), GQ895740 (B. malyai variant FeCH), GQ895742 (Onchocerca volvulus), GQ895743 (Dirofilaria immitis), and GQ895741 (Acanthocheilonema vitaezel)).

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Bovary (Fig. 3) worms, hybridization signals were detected in oocytes within the (Fig. 3) nuclei. These results confirm the function of the BmFeCH. The gene was cloned, and recombinant protein was expressed and purified. The clone produces active ferrochelatase with a specific activity of 30.5 nmol/mg per h. Zinc (H) protoporphyrin IX formation (Zn-PPIX). For comparative purposes, recombinant FeCHs from A. vitae (AvFeCH; 25 nmol/mg per h), the Wolbachia endosymbiont of B. malayi (wBmFeCH; 43.5 nmol/mg per h), and human (HsFeCH; 90.6 nmol/mg per h) were produced. Functionality was also assayed by addition of the competitive FeCH inhibitor N-methyl mesoporphyrin (NNMP) (18), which reduced or eliminated FeCH activity (Fig. 4). The IC50 values for NNMP inhibition of BmFeCH and AvFeCH were ~4.5 μM whereas wBmFeCH and HsFeCH were ~1.6 μM and 0.16 μM, respectively.

The in vivo functionality of BmFeCH was assessed by using genetic complementation of an Escherichia coli FeCH deletion mutant, ΔhemH (FeCH), mutant that is unable to grow unless hemin is supplied to the media (19). Supplementation with PPIX, the substrate for FeCH, failed to support growth of the mutant strain. However, the E. coli ΔhemH strain containing the BmFeCH cDNA grew in the absence of supplemented hemin. This result occurred with full-length BmFeCH and an N-terminally truncated version, as well as with wBmFeCH and HsFeCH.

To evaluate functionality in intact nematodes, we generated transgenic C. elegans lines that carried the full-length BmFeCH cDNA fused to the green fluorescent protein (GFP) DNA sequence (BmFeCH::GFP) integrated into the genome. One selected line, IP637, is similar to wild-type (WT) with respect to brood size and viability, suggesting that expression of BmFeCH::GFP has no major detrimental effect under normal culture conditions.

Control (WT) C. elegans grow on chemically defined axenic basal medium (ABM) if supplemented with hemin (refs. 5 and 20) and this study) but not PPIX (Fig. S3). Worms grown in ABM or PPIX-containing ABM do not die but arrest after one or two molts and fail to reach adulthood. Transgenic IP637 worms were also assayed for growth on axenic medium and were able to grow to adulthood when supplemented with either PPIX or to a lesser extent hemin (Fig. S3). FeCH inhibitor studies were performed with IP637 worms. In the presence of PPIX, their growth was inhibited when NNMP was added to the media. This effect could be partially rescued by addition of hemin to the media (Fig. S3). Taken together, these results indicate that BmFeCH is functionally expressed in C. elegans.

**BmFeCH Mitochondrial Targeting.** To confirm the function of the predicted BmFeCH mitochondrial-targeting domain, we compared localization of full-length BmFeCH::GFP to a version lacking 28 amino acids at the N terminus (IP633; BmFeCHΔ28::GFP) in transgenic C. elegans lines. Using MitoTracker Red CMXRos (which specifically stains mitochondria), we determined that, in lines carrying a full-length BmFeCH::GFP translational fusion (IP637 and IP634), the GFP signal localizes to mitochondria (Fig. 2). In contrast, in the transgenic C. elegans line carrying BmFeCHΔ28::GFP (IP633), the mitochondrial localization pattern is lost and the GFP signal is detected in the cytoplasm and nuclei. These results confirm our in silico localization prediction as well as previous studies in mammalian cells (17).

**In Vivo Localization.** To localize BmFeCH in B. malayi, a digoxigenin-labeled 467-bp BmFeCH RNA probe was used for in situ hybridization (ISH). BmFeCH appears to be ubiquitously expressed in both male and female tissues, except for female late-stage embryos and male late-stage sperm cells (Fig. 3). In adult female worms, hybridization signals were detected in oocytes within the ovary (Fig. S4) and early stage embryos, such as morula stage (Fig. 3B and C), but not in late-stage embryos, like curved microfilariae (Fig. 3D) and stretched microfilariae (Fig. 3E and F). However, the uterine epithelial cells that contain late-stage embryos showed strong hybridization signals (Fig. 3D–F). Hybridization signals were also observed in somatic tissues such as lateral cords, intestine, and hypodermis (Fig. 3A). In adult male worms, BmFeCH transcripts were mainly detected in early stages of developing sperm such as spermatocytes (Fig. 3G and H), but not in late stages such as spermatids (Fig. S3). BmFeCH transcripts were also detected in lateral cords, muscles (Fig. 3H and I), intestine, and vas deferens (Fig. 3J).

**BmFeCH Functionality.** We used a variety of approaches to verify the functionality of BmFeCH. The gene was cloned, and recombinant protein was expressed and purified. The clone produces active ferrochelatase with a specific activity of 30.5 nmol/mg per h. Zinc (H) protoporphyrin IX formation (Zn-PPIX). For comparative purposes, recombinant FeCHs from A. vitae (AvFeCH; 25 nmol/mg per h), the Wolbachia endosymbiont of B. malayi (wBmFeCH; 43.5 nmol/mg per h), and human (HsFeCH; 90.6 nmol/mg per h) were produced. Functionality was also assayed by addition of the competitive FeCH inhibitor N-methyl mesoporphyrin (NNMP) (18), which reduced or eliminated FeCH activity (Fig. 4). The IC50 values for NNMP inhibition of BmFeCH and AvFeCH were ~4.5 μM whereas wBmFeCH and HsFeCH were ~1.6 μM and 0.16 μM, respectively.

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**BmFeCH Essentiality.** Because *B. malayi* contains the *Wolbachia* endosymbiont that encodes a functional FeCH gene and is sensitive to antibiotics (such as the tetracyclines), we were able to assess the relative contributions of wBmFeCH and BmFeCH by comparing the effect of NMMP on adult *B. malayi* and tetracycline-treated *B. malayi* adults cleared of their *Wolbachia*. We measured *Wolbachia* levels by quantitative PCR (qPCR) and measured worm motility in both groups. Motility in adult *B. malayi* females and males was significantly reduced by exposure to NMMP relative to untreated controls (Fig. 5A). Viability in both adult *B. malayi* and tetracycline-treated *B. malayi* was also significantly decreased in all NMMP treatment groups compared with the untreated controls (Fig. 5B). These results suggest that the inhibition was not *Wolbachia* specific, as similar results were obtained in antibiotic- and nonantibiotic-treated groups. Presumably, both wBmFeCH and BmFeCH are affected by NMMP, further suggesting the worm’s requirement for heme biosynthesis for viability. The sensitivity of filarial nematode FeCH to NMMP was also demonstrated in studies using adult *A. viteae*, a species lacking *Wolbachia*. Male and female worms showed reduced motility by approximately day 4 of exposure to 100 μM NMMP and appeared moribund by day 7 (Fig. 5C).

RNA interference (RNAi) was used to investigate the effect of FeCH depletion upon the *B. malayi* female germ line and during embryonic development. Unlike *C. elegans*, embryonic development in *B. malayi* takes place entirely within the uterus, so RNAi in *B. malayi* potentially affects every developmental stage. We found that FeCH RNAi led to consistent abnormalities in the nuclei in the germ line as well as in embryos. Untreated WT ovaries displayed spheroid nuclei surrounded by numerous *Wolbachia*, and organized around a central rachis in a syncytium (Fig. 6A) whereas FeCH RNAi-treated females exhibited misshapen germ-line nuclei, often lenticular or polygonal, and also occasionally characterized by an uneven *Wolbachia* distribution and abnormal rachis (Fig. 6B). The same misshapen nuclei were observed in all developing embryos (*n > 500*), associated with embryonic elongation and actin cytoskeleton defects (Fig. 6C and E). The majority of elongated embryos exposed to FeCH heterogeneous short interfering RNA (hsiRNA) eventually ruptured, likely due to cytoskeleton defects compromising overall hypodermal integrity (Fig. 6F and H). The observation of similar phenotypes when using hsiRNA mixtures derived from different regions of *BmFeCH* indicates that the defects are specific to FeCH transcript reduction rather than nonspecific off-target effects. By contrast, mature microfilariae did not display any phenotype, perhaps because of reduced FeCH metabolism or insensitivity to RNAi at this stage. Control worms incubated in the absence of RNA (Fig. 6A, D, and G) or in the presence of hsiRNA derived from a plasmid vector (Fig. 6I) did not show any defects.

**Discussion**

We report a functional LGT in a nematode species parasitic in the Metazoa. LGT is widespread among certain nematodes and may have played a role in the remarkable evolutionary success of this phylum and in the adoption of parasitic lifestyles. Genomes of several plant-parasitic nematodes contain genes acquired by LGT that encode enzymes involved in plant cell wall degradation (2, 3). LGT has previously been noted in animal parasitic filarial nematodes (7–9), but thus far none have been shown to be functional. BmFeCH is functional and essential, as indicated by multiple lines of evidence, including *E. coli* complementation, heme auxotrophy complementation, ISH experiments showing ubiquitous expression in males and females, inhibitor studies, and the lethal effect of RNAi silencing of BmFeCH in *B. malayi* embryos.

We suggest that heme biosynthesis in the Nematoda was completely lost (or alternatively, it was never present) but that the last step, catalyzing porphyrin metallation, was acquired by LGT in an ancestor of modern nematodes where it (re)assumed function in the same compartment as in other heme-synthesizing eukaryotes. Although BmFeCH is most closely related to α-proteobacterial FeCH, during its evolution within metazoan organisms, the gene has evolved into a typical eukaryotic gene by acquiring introns and exons. BmFeCH also encodes an additional N-terminal mitochondrial targeting sequence that is not found in α-proteobacterial FeCHs but is conserved across filarial FeCHs. The mitochondrial localization, revealed by transgene expression in *C. elegans*, is
consistent with localization in other eukaryotic heme prototrophs, where FeCH is targeted to the inner surface of the inner mitochondrial membrane and is involved in porphyrin metallation (21).

A. viteae maintained for 7 d in vitro. (A) Motility of tetracycline-treated B. malayi and untreated control worms cultured in the presence of 100 μM (red), 10 μM (blue) and 0 μM (green) NMMP. (Dose response of control male worms to 10 μM NMMP not performed). Motility was measured on a scale of 0 to 4 and the data plotted as a percentage of the control worms (untreated with NMMP) at each time point set at 100%. (B) Viability of tetracycline-treated B. malayi (maroon bars) and untreated control worms (green bars) cultured in the presence of 100 μM, 10 μM, or 0 μM NMMP for 7 days. Viability was determined by the MTT assay and the value of the control group was set to 100. Viability of control male worms after exposure to 10 μM NMMP not performed. (C) Motility of A. viteae male and female worms cultured in the presence of 100 μM NMMP (red), 100 μM ivermectin (green), or no inhibitor (blue). Motility was measured on a scale of 0 to 4 and the data plotted as a percentage of control worms (untreated), set at 100%.

It is not obvious why some filarial nematodes have functional FeCH genes when they also have Wolbachia endosymbionts that contain a complete heme biosynthetic pathway. Possibly, the FeCH gene enables nematode survival on PPIX in life-cycle stages where exogenous heme is scarce or Wolbachia titer is low. Because the FeCH gene does not appear to have originated from the Wolbachia endosymbiont and has been retained in filarial species not harboring Wolbachia, it seems unlikely that this gene plays a role in Wolbachia–host interactions. Rather, the presence of LGT FeCH in other nematode clades (such as in Strongyloides spp; clade IV) and the essentiality in filarial nematode embryos point to a model whereby a FeCH may help meet high heme demands at critical stages of the parasite life cycle. However, it is possible that, despite retaining FeCH activity, the protein has assumed an alternative essential cellular function distinct from porphyrin metallation.

The finding of FeCH only in filarial nematodes and members of the genus Strongyloides is suggestive of a role in adaptations toward animal parasitism in these groups (16). FeCH has assumed a critical role in filarial nematode species where it is essential. The bacterial ancestry of BmFeCH results in considerable divergence between these nematode enzymes and their mammalian counterparts. Such divergence lends an opportunity for targeting FeCH using drugs specific to bacterial enzymes for filariasis control. It was suggested that the presence of BmFeCH, along with the FeCH-like protein and the wBmFeCH (from Wolbachia) in the same parasitic organism, might represent a serious challenge to efforts aimed at developing FeCH as a drug target (15). However, our findings indicate that nematode FeCH-like proteins do not function as FeCHs. Conversely, RNAi
indicates that BmFeCH is essential for B. malayi and that the FeCH inhibitor NMMP impairs both filarial motility and viability. Accordingly, drug screens targeting filarial FeCHs are warranted.

**Materials and Methods**

Additional information is provided in SI Materials and Methods.

**Filarial Nematode Material.** B. malayi worms were purchased from TRS Labs. *O. volvulus* worms were generously provided by Sara Lustigman (New York Blood Center, New York, NY). *A. viteae* worms were a gift from Kenneth Pfarr (University Hospital Bonn, Bonn, Germany) and Peter Fischer (Washington University, St Louis, MO). *M. immitis* RNA samples were kindly provided by Michael Crawford (Divergence, St. Louis, MO).

**FeCH Cloning into E. coli and Expression of FeCH Genes.** Worm RNA extractions were performed using the RNeasy mini kit (Qiagen). Rapid amplification of 5′ and 3′ cDNA ends (RACE) were carried out with the SMARTer RACE cDNA amplification kit (Clontech) from RNA preparations of *B. malayi*, *O. volvulus*, *M. immitis*, and *A. viteae* worms. Subsequent PCR amplification and DNA sequencing between the 3′ and 5′ ends enabled the completion of accurate coding sequences. The newly acquired *B. malayi* FeCH (BmFeCH) and *A. viteae* FeCH (AvFeCH) coding sequences were cloned into pET21a+ vector (Novagen) for production of recombinant proteins with C-terminal 6XHis-tags in *E. coli* using methods established previously (22). For controls and for comparative purposes, we produced recombinant FeCH from human (HsFeCH), Wolbachia from *B. malayi* (wBmFeCH) and *E. coli* (EcFeCH) as described previously (22).

**Ferrochelatase Enzyme Assays.** Enzyme activities were assayed using purified recombinant N-terminal 6XHis-tagged BmFeCH, AvFeCH, wBmFeCH, and HsFeCH proteins at 37 °C for 10–20 min, essentially as previously described (23). For NMMP (N-methyl mesoporphyrin; Frontier Sciences), inhibitor studies, assays were initiated by the addition of the substrate protoporphyrin IX (PPX) after a 10-min enzyme preincubation with different concentrations of inhibitor.

**Sequence and Phylogenetic Analysis.** For alignments, putative FeCH protein sequences obtained from the different filarial species, along with FeCH orthologs retrieved from National Center for Biotechnology Information GenBank (ncbi.nlm.nih.gov) following BLASTX similarity searches, were aligned using CLUSTALX 1.83 (24). After manual refinement, the sequences in the alignment were further analyzed by TargetP 1.1 (25) for predicting subcellular targeting signals. BmFeCH cDNA sequences were compared with their genomic locus sequence by LFasta (pbil.univ-lyon1.fr/lfasta.php).

For the consensus tree construction, sequences were aligned and curated using the phylogeny.fr web service at phylogeny.fr (26). Sequence alignments were performed with T-Coffee (v6.85) (27). After alignment, the sequences were curated, and ambiguous regions (i.e., containing gaps and/or poorly aligned) were removed with Gblocks (v0.91B) (28). The curated alignment containing 126 sites was analyzed locally using MrBayes (v3.2.1 x64) (29). Trees were visualized with FigTree v1.4.0 (tree.bio.ed.ac.uk), and alignment containing 126 sites was analyzed locally using MrBayes (v3.2.1 x64) (29). Trees were visualized with FigTree v1.4.0 (tree.bio.ed.ac.uk), and final figures were created in Inkscape (inkscape.org).

**E. coli ΔhemH Complementation with Heterologous FeCH Genes.** The *E. coli ΔhemH* deletion strain VS200 has been described previously (19). The pET21a vectors containing the corresponding human, *B. malayi*, Wolbachia, and *E. coli* putative FeCH genes were transformed into the *E. coli ΔhemH* strain, both with and without RIL plasmid cotransformation. Transformants were selected on 20 μM hemin-containing LB plates with appropriate antibiotics and incubated at 37 °C overnight. Hemin was freshly prepared as a 5-mM stock solution in 50% (vol/vol) ethanol containing 0.02 M NaOH. The selected transgenic clones were further tested on LB plates with no hemin addition.

**E.coli-Based FeCH Inhibitor Assays.** The *E. coli hemH* mutant strain VS200, containing separately HisFeCH, BmFeCH, AvFeCH, wBmFeCH, or EcFeCH genes, was used in inhibitor assays. Transgenic *E. coli* clones from 0.6–0.8 OD₆₀₀ (in LB medium) were diluted to 0.01 OD₆₀₀ before initiating growth assays in the presence or absence of different concentrations of NMMP. The *E. coli* cells were grown at 30 °C for 3 h before determination of cell growth by measurement of the final OD₆₀₀ values. The average cell growth ratio (final OD₆₀₀/0.01) for untreated control was set at 1.0.

**Ex Vivo B. malayi and A. viteae Motility and Viability Tests.** *B. malayi* adults were isolated from jirds. For *Wolbachia*-depleted *B. malayi*, jirds were treated at Liverpool School of Tropical Medicine (Liverpool, United Kingdom) with 2.5 mg/ml tetracycline (Sigma) in drinking water for 6 wk (with tetracycline/water prepared fresh daily) plus 2 wk of water alone before recovery of adult worms. The reduction in *Wolbachia* load following tetracycline treatment was determined by quantitative PCR (qPCR) to determine the copy number of the genes encoding *Wolbachia* surface protein ( wsp) and *B. malayi* GST (gst) as previously described (30). Following treatment, *Wolbachia* loads were reduced by 99.24% in female worms and 99.91% in male worms. Worms were cultured in the presence of different concentrations of NMMP for 7 d. Motility was scored daily using a scoring system described previously (31), with 0 being immotile and 4 being highly active and motile and the results for each group were presented as a percentage of the control group. Viability was evaluated at day 7 using the MTT assay (32, 33). The reduction of MTT for each group is presented as a percentage of the control group.

Adult male and female *A. viteae* were incubated with different concentrations of the FeCH inhibitor NMMP for 7 d (three replicates per experiment). The anthelmintic ivermectin (100 μM; Sigma) was used as a positive control. Motility was measured daily (as described above and in ref. 31).

**In Situ Hybridization.** Parasite material and slide preparation, RNA probe synthesis, and ISH procedures were performed as described previously (34). For these experiments, 10-μm frozen sections cut from adult *B. malayi* worms were used. Digoxygenin-labeled RNA probes were synthesized from a 467-bp fragment of *BmFeCH* that was subcloned by the following primer pair: forward primer, 5′-CGAATCTCTACCCGCT-3′; reverse primer, 5′-TCAAGAACATTGCAGAAGT-3′. Sense and antisense RNA probes were used for ISH, and the sense probe served as a negative control.

**B. malayi RNAi.** *B. malayi* females were soaked in 1 μM BmFeCH hsiRNA for 2 d (changing the medium and RNA every 12 h). The hsiRNA mixtures were prepared from (i) a 514-bp fragment corresponding to nucleotide positions 170–683 of the *BmFeCH* ORF, and (ii) a 393-bp fragment corresponding to positions 663–1055 of the ORF. Methods for preparing hsiRNA mixtures for use in RNAi and the subsequent fixation and staining protocols have been described previously (35). Two different preparations of hsiRNA derived from the 5′ end of *BmFeCH* were used in separate experiments whereas hsiRNA derived from the 3′ region was used once. Controls included worms similarly cultured both in the absence of any RNA or in the presence of 1 μM control hsiRNA (Lit28i Polylinker ShortCut sRNA Mix; New England Biolabs).

**C. elegans Strains, Constructs, and Transgenesis.** *C. elegans* N2 strain var. Bristol was obtained from the *C. elegans* stock-center (Caenorhabditis Genetics Center, University of Minnesota) and cultured by standard methods unless otherwise stated. unc-119(ed3) mutants were used in this work (36). The following transgenic *C. elegans* strains were also generated and used: iP633: unc-119(ed3), nEx172 [pDP#MM016b; psur5SNTER-BmFeCH-GFP] iP634: unc-119(ed3), nEx173 [pDP#MM016b; psur5BmFeCH-GFP] iP637: nbs171 [pDP#MM016b; psur5BmFeCH-GFP].

Integrand line iP637 was generated from strain iP634 using the standard UV light array integration procedure (37).

A general backbone for all *C. elegans* transgenic constructs was prepared by modifying the GFP expression vector pPD95.75 (gift of Andrew Fire Department of Genetics, Stanford University, Palo Alto, CA). Constructs were injected into unc-119(ed3) worms according to standard protocols (38). Plasmid pDP#MM016b, carrying a unc-119(ed3) rescue construct (36), was used as a cotransformation marker. For each construct, at least two independent transgenic lines were established and inspected for consistency in expression patterns; a single line was used in further analysis as a reference strain.

**C. elegans Microscopy.** Mitochondrial in vivo staining for colocalization experiments was performed by growing worms in the dark on nematode growth medium (NGM) standard plates containing Mitotracker Red CMXRos (2 μM; Molecular Probes). Differential interference contrast (DIC) micrographs were acquired with a Zeiss Axiovert 200M microscope (Zeiss) equipped with epifluorescence and processed using the Axiosvision package release 4.5. Confocal pictures were acquired with a LSM 510 META laser scanning microscope (Zeiss) and processed with the LSM Zeiss package release 4.2.

**Axenic Growth of *C. elegans*.** Axenic growth of *C. elegans* was by established procedure (39) with some minor modifications. Axenic cultures were produced using the standard sodium hypochlorite treatment (40). Eggs were allowed to hatch overnight in axenic basal medium (ABM) in the absence of a home environment. The concentration of live L1 larvae was adjusted to obtain an average concentration of 15–30 animals per 100 μL. A porphyrin source was prepared with T-Coffee (v6.85) (27). After alignment, the sequences were curated, and ambiguous regions (i.e., containing gaps and/or poorly aligned) were removed with Gblocks (v0.91B) (28). The curated alignment containing 126 sites was analyzed locally using MrBayes (v3.2.1 x64) (29). Trees were visualized with FigTree v1.4.0 (tree.bio.ed.ac.uk), and final figures were created in Inkscape (inkscape.org).

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added in the form of hemin chloride or PPIX at a final concentration of 10 mg/L, equivalent to 15.3 μM hemin chloride and 17.8 μM PPIX. A control lacking any porphyrin was also included. Worms were cultured in 96-well plates in a 100 µL volume for 11 d at 25 °C. Axenic medium supplemented with hemin supports slow growth of *C. elegans*, allowing for the completion of only one generation cycle in this period. Twenty-four wells were cultured for a given strain and condition. The number of parental animals reaching adulthood was scored and averaged. NMMP inhibitor assays were performed by supplementing the cultures containing hemin or PPIX with NMMP at 10 μM and assaying growth of one generation of worms as described above. A solvent-control was included in the NMMP test.

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**Supporting Information**

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**SI Materials and Methods**

**Phylogenetic Analysis.** For consensus tree construction, sequences were aligned and curated using the phylogeny.fr Web service at [www.phylogeny.fr](http://www.phylogeny.fr) (1). Sequence alignments were performed with T-Coffee (v6.85) (2) using the following pair-wise alignment methods: the 10 best local alignments (Lalign_pair) and an accurate global alignment (slow_pair). After alignment, the sequences were curated and ambiguous regions (i.e., containing gaps and/or poorly aligned) were removed with Gblocks (v0.91b) (3) using the following parameters: minimum length of a block after gap elimination: 5; no gap positions were allowed in the final alignment; all segments with contiguous nonconserved positions bigger than 8 were rejected; minimum number of sequences for a flank position, 85%. The curated alignment containing 126 sites was analyzed locally using MrBayes (v3.2.1 x64) (4) using the following parameters: minimum length of a block after gap elimination: 5; no gap positions were allowed in the final alignment; all segments with contiguous nonconserved positions bigger than 8 were rejected; minimum number of sequences for a flank position, 85%. The curated alignment containing 126 sites was analyzed locally using MrBayes (v3.2.1 x64) (4) using the following parameters: preset aamodelpr = mixed; lset rates = gamma; mempgen = 500000, samplefreq = 100, printfreq = 1000, diagfreq = 5000. At the end of the run after 500,000 generations, the SD of split frequencies was 0.007. A relative burn in fraction of 0.25 was used when calculating the consensus tree. Trees were visualized with FigTree v1.4.0 (tree.bio.ed.ac.uk), and final figures were created in Inkscape (inkscape.org).

**Ferrochelatase Cloning into *Escherichia coli* and Expression of Ferrochelatase Genes.** For amplifications of 5′ and 3′ cDNA ends (RACE) from RNA preparations of *Brugia malayi*, *Onchocerca volvulus*, *Dicrofilaria immitis*, and *Acanthocheilonema vitellum* worms, gene-specific degenerate forward and reverse primers were designed based upon alignments of diverse ferrochelatase (*FeCH*) gene and EST sequences available in the National Center for Biotechnology Information and European Molecular Biology Laboratory–European Bioinformatics Institute databases. These primers were used in combination with the universal RACE (Rapid Amplification of cDNA Ends) primer provided by the SMARTer RACE cDNA amplification kit (Clontech) for amplification from the species-specific cDNA RACE pools (5′ or 3′). Subsequent PCR amplification and DNA sequencing between the 5′ and 3′ ends enabled acquisition of complete coding sequences. Except for the *FeCH* gene from *E. coli* (Ec*FeCH*), all cloned gene constructs were cDNA optimized to provide maximal gene expression. All *FeCH* gene constructs were transformed into *E. coli* containing the RIL plasmid (Agilent) to increase expression. The various *FeCH* genes were also cloned into the pTrcHisA vector (Invitrogen) for subsequent expression of proteins with N-terminal 6xHis-tags in *E. coli*. For recombinant FeCH expression in *E. coli*, the FeCH genes from eukaryotes were cloned with truncations at their N termini to remove the potential mitochondrial targeting domains. For FeCH from *B. malayi* (BmFeCH), the N-terminal 36 amino acids were eliminated; for FeCH from *A. vitellum* (AvFeCH), 38 amino acids; and for FeCH from *H. sapiens* (HsFeCH), 62 amino acids were removed. For comparative purposes, a full-length BmFeCH gene was also cloned and expressed.

**FeCH Enzyme Assays.** N-terminal 6xHis-tagged BmFeCH, AvFeCH, wBmFeCH, and HsFeCH recombinant proteins were purified for enzyme studies. When appropriate, the inhibitor NMMP (N-methyl mesoporphyrin; Frontier Scientific) was freshly prepared as a 5-mM stock solution in 50% ethanol containing 0.02 M NaOH. The enzyme reactions were terminated by addition of acetone. After removing the precipitated proteins by centrifugation, the formation of the end product of the reactions, Zinc (II) protoporphyrin IX (Zn-PPIX), was measured using a Perkin-Elmer LS50B spectrophotometer (excitation 420 nM; emission 587 nm).

**Ex Vivo *B. malayi* and *A. viteae* Motility and Viability Tests.** *B. malayi* adult worms isolated from control or tetracycline treated jirds were cultured in duplicate wells (two adult females or males per well, in 24-well plates, for 7 d in 2 mL of RPMI 1640 containing 10% (vol/vol) FBS (Pepbio) and penicillin-streptomycin (In-vitrogen); 200 U/mL; 200 μg/mL final concentration) with varying concentrations of NMMP. Worms not treated with NMMP were cultured in medium containing solvent (1% ethanol and 0.0004 M NaOH). Medium and inhibitor were changed every 2 d. Motility and viability were determined as described in the text.

For *A. viteae* experiments, one adult female or three adult males per replicate (experiment repeated twice) were cultured in RPMI 1640 with 2 mM glutamine, 25 mM Hepes (Gibco) with 10% (vol/vol) FCS (Gibco), 100 U/mL streptomycin, 100 μg/mL penicillin, and 0.25 μg/mL amphotericin B (Sigma). Medium was changed every 2 d. Worms not treated with NMMP or ivermectin were cultured in medium containing solvent only. Motility was determined as described in the text.

**C. elegans Strains, Constructs, and Transgenesis.** A general backbone for all *C. elegans* transgenic constructs was prepared by modifying the GFP expression vector pPD95.75 (gift of Andrew Fire Department of Genetics, Stanford University, Palo Alto, CA). A 3.5-kb sequence of the sur-5 promoter contained in plasmid pTg696 (5) was cloned into the multiple cloning site of pPD95.75 containing the SpeI site at the 5′ end of the sequence and by inserting an NheI site at the 3′ end of the sur-5 promoter sequence upstream of the endogenous XmaI site of pPD95.75. psur5BmFeCH-GFP was generated by cloning the BmFeCH cDNA between the sur-5 promoter and in frame with the C-terminal GFP sequence using the NheI and XmaI sites. psur5NTER-BmFeCH-GFP was generated using PCR by removing 84 bp of the 5′ sequence of psur5BmFeCH-GFP starting from the ATG initiation codon. An ATG codon located downstream of the deleted sequence served as initiation codon. This construct resulted in an N-terminally truncated version of BmFeCH (BmFeCHΔ28; GFP) lacking the first 28 amino acids encoded by exon 1 and encompassing the predicted mitochondrial targeting domain. All constructs were verified by restriction digestion and DNA sequencing before injection.

Constructs were injected into *unc-119(ed3)* worms according to standard protocols (6). Plasmid pDP#MM016b, carrying a *unc-119(+) rescue construct* (7) was used as a cotransformation marker. Injection mixtures had the following composition: plasmid of interest 10 μg; Low Molecular Weight DNA Ladder (New England Biolabs) as a filler to give a final 100 ng DNA/μL concentration.

**Axenic Growth of *C. elegans.*** Worms were grown in axenic basal medium [ABM; 3% (wt/vol) dry yeast extract, 3% (wt/vol) soy peptone (Sigma; cat. no. P0521), 1% (wt/vol) dextrose, cholesterol 10 mg/L] in the absence of a heme source. A porphyrin source was added in the form of hemin chloride or PPIX at a final concentration of 10 mg/L, equivalent to 15.3 μM hemin chloride and 17.8 μM PPIX (hemin stock solution: 1 mg/mL in 0.1 M NaOH; PPIX stock solution: 1 mg/mL in 0.3 M ammonium acetate; the latter was added after chilling the culture media to minimize precipitation).

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Heme biosynthetic pathway in animals, fungi, and α-proteobacteria (plants, algae, archaeabacteria, and other eubacteria initiate from glutamate). Enzyme names are boxed and the FeCH inhibitor NMMP is shown on the left. ALAS, δ-aminolevulinic acid synthase; CPO, coproporphyrinogen oxidase; FeCH, ferrochelatase; HMBS, hydroxymethylbilane synthase; PBGS, porphobilinogen synthase; PPO, protoporphyrinogen oxidase; UROD, uroporphyrinogen de-carboxylase; UROS, uroporphyrinogen III synthase. The first step and last three steps occur in mitochondria.

Fig. S1. Heme biosynthetic pathway in animals, fungi, and α-proteobacteria (plants, algae, archaeabacteria, and other eubacteria initiate from glutamate). Enzyme names are boxed and the FeCH inhibitor NMMP is shown on the left. ALAS, δ-aminolevulinic acid synthase; CPO, coproporphyrinogen oxidase; FeCH, ferrochelatase; HMBS, hydroxymethylbilane synthase; PBGS, porphobilinogen synthase; PPO, protoporphyrinogen oxidase; UROD, uroporphyrinogen de-carboxylase; UROS, uroporphyrinogen III synthase. The first step and last three steps occur in mitochondria.

ClustalX sequence alignment of BmFeCH with FeCH orthologs from various eukaryotic and prokaryotic species. Positions with identical residues are shown in black, whereas positions with similar residues are shown in blue. Underneath the alignments, region involved in dimerization is found only in FeCH proteins of eukaryotic origin. The four cysteine residues involved in \([\text{2Fe-2S}]\) cluster formation are shown in bold typeface with the predicted mitochondrial targeting domains underlined. The invariant active site residues are shown in red. A C-terminus extension (green) involved in dimerization is found only in FeCH proteins of eukaryotic origin.

**Fig. S2.** ClustalX sequence alignment of BmFeCH with FeCH orthologs from Sinorhizobium meliloti (SmFeCH), Wolbachia from Brugia malayi (wBmFeCH), and Homo sapiens (HsFeCH).
Fig. S3. Growth of WT *Caenorhabditis elegans* (blue bars) and transgenic *C. elegans* IP637 carrying full-length *BmFeCH* (red bars) on axenic basal medium (ABM) and ABM supplemented with either PPIX (∼18 µM) or hemin (∼15 µM) in the presence or absence of the FeCH inhibitor NMMP (1 µM and 10 µM).