The Wild Worm Codon Adapter: a web tool for automated codon adaptation of transgenes for expression in non-Caenorhabditis nematodes

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Abstract

Advances in genomics techniques are expanding the range of nematode species that are amenable to transgenesis. Due to divergent codon usage biases across species, codon optimization is often a critical step for the successful expression of exogenous transgenes in nematodes. Platforms for generating DNA sequences codon-optimized for the free-living model nematode Caenorhabditis elegans are broadly available. However, until now such tools did not exist for non-Caenorhabditis nematodes. We therefore developed the Wild Worm Codon Adapter, a tool for rapid transgene codon optimization for expression in non-Caenorhabditis nematodes. The app includes built-in optimization for parasitic nematodes in the Strongyloides, Nippostrongylus, and Brugia genera as well as the predatory nematode Pristionchus pacificus. The app also supports custom optimization for any species using user-provided optimization rules. In addition, the app supports automated insertion of synthetic or native introns, as well as the analysis of codon bias in transgene and native sequences. Here, we describe this web-based tool and demonstrate how it may be used to analyze genome-wide codon bias in Strongyloides species.

Keywords: Strongyloides; Brugia; Pristionchus; Nippostrongylus; nematodes; codon optimization; introns; transgenesis

Introduction

Parasitic nematodes, including soil-transmitted gastrointestinal parasites in the genus Strongyloides and filarial nematodes such as Brugia malayi, are a major source of disease and economic burden (Lustigman et al. 2012). While Caenorhabditis elegans is often used as a model system for the study of parasitic nematodes, parasitic nematodes are behaviorally and genetically divergent from C. elegans; for example, they engage in a number of parasite-specific behaviors such as host seeking, host invasion, and intra-host migration (Haas 2003; Gang and Hallem 2016). Establishing methods that allow researchers to genetically manipulate parasitic nematodes directly is critical for understanding the genetic and cellular basis of parasitism in these species.

Historically, the application of functional genomics techniques to parasitic nematodes has lagged behind their use in the free-living model nematode C. elegans, due in part to the limited availability of parasite genomic information (Castelletto et al. 2020). High-quality reference genomes for many parasitic nematode species are now available (Hunt et al. 2016; Howe et al. 2017; International Helminth Genomes Consortium 2019), and are a critical resource for parasitic nematode functional genomics techniques such as transgenesis and CRISPR/Cas9-mediated mutagenesis (Lok et al. 2017; Castelletto et al. 2020; Liu et al. 2020). Transgenesis in parasitic nematodes is an essential tool for mechanistic studies of parasite development and behavior (Bryant et al. 2018; Gang et al. 2020). As our technical understanding of nematode genomics continues to develop beyond C. elegans, establishing accessible tools that automate the transgene design process for a broad selection of nematode species will greatly facilitate the application of genomics techniques in these species.

Transgenesis protocols are increasingly well-established in non-Caenorhabditis nematode species, including three soil-transmitted gastrointestinal parasites in the Strongyloidesidae family—the human parasite Strongyloides stercoralis, the rodent parasite Strongyloides ratti, and the Australian brushtail possum parasite Parastrongyloides trichosuri—as well as the rodent gastrointestinal parasite Nippostrongylus brasiliensis, the human-parasitic filarial nematode B. malayi, the predatory nematode Pristionchus pacificus, and the free-living nematodes Auanema rhodensis and Auanema freiburgensis (Grant et al. 2006; Lok et al. 2017; Adams et al. 2019; Castelletto et al. 2021; Han et al. 2020). Intragonadal microinjection, in which exogenous DNA is injected directly into the gonad, has been used to generate progeny expressing a range of transgenes (Schlager et al. 2009; Lok et al. 2017; Adams et al. 2019; Hong et al. 2019; Carstensen et al. 2021; Castelletto et al. 2020; Han et al. 2020). Intragonadal microinjection has also been used to achieve CRISPR/Cas9-mediated mutagenesis in S. stercoralis, S. ratti, P. pacificus, and Auanema species (Witte et al. 2015; Gang et al. 2017; Lok et al. 2017; Bryant et al. 2018; Adams et al. 2019; Han et al. 2020; Castelletto et al. 2021).
In B. malayi, transfection of infective larvae has been used to deliver reporter plasmids and CRISPR constructs (Liu et al. 2018, 2020). Most recently, lentiviral transduction of infective larvae was used to deliver RNA interference molecules and drive expression of fluorescent reporters in N. brasiliensis (Hagen et al. 2021).

In non-Caenorhabditis nematodes and other species, successful transgene expression often requires the use of species-specific preferred codons. Codon usage bias is pervasive in species from all taxa, including nematodes: most amino acids may be encoded by multiple synonymous codons, and individual species tend to favor a specific set of codons, particularly for encoding highly expressed genes (Sharp and Li 1987; Cutter et al. 2006; Mitreva et al. 2006). The use of preferred codons is thought to promote efficient translation, although the exact mechanisms are not clear (Plotkin and Kudla 2011). Codon usage bias is believed to regulate the expression of exogenous transgenes as well as endogenous genes (Redemann et al. 2011). In non-Caenorhabditis nematodes, expression of exogenous genes from transgenes is promoted by the use of species-specific codon usage patterns (Han et al. 2020; Hagen et al. 2021).

Although the process of codon-adapting transgenes for C. elegans is simplified by web-based platforms (Grote et al. 2005; Redemann et al. 2011), transgenes codon-adapted for other nematode species were previously designed by hand. Therefore, we created a web-based application, the Wild Worm Codon Adapter, that automates the process of codon optimization for transgene expression in non-Caenorhabditis nematode species. Furthermore, the application permits automated insertion of synthetic or native introns into codon-optimized cDNA sequences, inclusion of which can significantly increase gene expression (Junio et al. 2008; Li et al. 2011; Crane et al. 2019; Han et al. 2020). Finally, the app enables users to rapidly assess relative codon bias of transgene sequences and native genes using genus-specific codon adaptation indices.

**Materials and methods**

**Data source and preferred codon selection**

Codon usage rules for Strongyloides, Brugia, Pristionchus, and C. elegans were calculated based on previously published codon count and frequency data (Supplementary File S1). For Strongyloides, codon frequency data are from 11,458 codons from the 50 most abundant S. ratti expressed sequence tag (EST) sequences (Mitreva et al. 2006). For C. elegans, codon count data are from 178 genes (73,164 codons) with the highest bias toward translationally optimal codons (Sharp and Bradnam 1997). For Brugia and Pristionchus, codon frequency data are from the ~10% of genes (bin 7) with the highest expression (Han et al. 2020). For Nippostrongylus, RNA-seq expression data across three N. brasiliensis life stages [infective 3rd-stage larvae (iL3s), activated iL3s, red-blood-cell-feeding iL3s] was retrieved from WormBase ParaSite (Eccles et al. 2018). The 10% of genes with the highest median expression across all life stages were identified (2279 genes). Coding sequences for these highly expressed genes were retrieved from WormBase ParaSite with the biomart package v2.42.1 (Durinc et al. 2005, 2009), and total codon usage counts (Supplementary File S1) were calculated using the uco function in the seqinr package v3.6.1.

Count and frequency data were used to quantify the relative adaptiveness of individual codons: the frequency that codon “i” encodes amino acid “AA” ÷ the frequency of the codon most often used for encoding amino acid “AA” (Sharp and Li 1987; Jansen et al. 2003). Preferred codons were defined as the codons with the highest relative adaptiveness value for each amino acid (Supplementary File S1).

**Codon adaptation index**

The codon biases of individual sequences are quantified by calculating a Codon Adaptation Index (CAI), defined as the geometric average of relative adaptiveness of all codons in the sequence (Sharp and Li 1987; Jansen et al. 2003). CAI values relative to species-specific relative adaptiveness values are calculated using the seqinr package v3.6.1.

**Fractional GC content**

The fraction of G+C bases in a sequence is calculated using the seqinr package v3.6.1.

**Intron insertion**

Intron sequences are either: the three canonical artificial intron sequences used for C. elegans (Fire et al. 1995), P. pacificus native introns (Han et al. 2020), Periodic A<sub>r</sub>/T<sub>r</sub>, Cluster (PATC)-rich introns from the C. elegans gene smu-2 (introns 3–5) (Aljohni et al. 2020), or custom introns provided by users via FASTA file upload. Built-in intron sequences are flanked by canonical 5′-GT…AG-3′ splice recognition sequences (Shapiro and Senapathy 1987; Blumenthal and Steward, 1997; Wheeler et al. 2020). For intron placement, the optimized cDNA sequence is divided at three predicted intron insertion sites spaced approximately equidistantly. Users may choose to insert introns at the equidistant sites, or may further refine insertion site locations by identifying the closest conserved invertebrate exon splice sites (5′-AG+G-3′, 5′-AG+AA-3′; the “+” symbol indicates the exact insertion site) (Shapiro and Senapathy 1987; Blumenthal and Steward, 1997). The user-specified number of introns (up to a maximum of three) are inserted into the sequence using the 5′ insertion site first and continuing in the 3′ direction (Crane et al. 2019; Han et al. 2020). When inserting introns using conserved invertebrate exon splice sites, if there are fewer insertion sites than the user-requested number of introns, the program will insert as many introns as there are available insertion sites.

**Coding sequence lookup**

In “Analyze Sequences” mode, users may submit search terms including stable gene IDs, C. elegans gene names, or matched gene IDs and cDNA sequences as either a two-column CSV file or a FASTA file. If users supply only gene IDs, either via a text box or file upload, the app first fetches the coding sequences from WormBase ParaSite via the biomart package v2.42.1 (Durinc et al. 2005, 2009). The following types of gene IDs may be used: stable gene or transcript IDs with prefixes “SSTP,” “SRAE,” “SPAL,” “SVE,” “Ppa,” “Bma,” “NBR,” or “WB”; C. elegans stable transcript IDs; or C. elegans gene names prefaced with the string “Ce-” (e.g., Ce-tau-4).

**Genome-wide codon bias and gene ontology (GO) analysis**

FASTA files containing all coding sequences (CDS) for S. stercoralis, S. ratti, S. papillosus, S. venezuelensis, N. brasiliensis, B. malayi, P. pacificus, and C. elegans were downloaded from WormBase ParaSite (WBP515) and analyzed using the app. For C. elegans and Strongyloides species, results were filtered to identify six functional subsets: the 2% of genes with highest and lowest S. ratti CAI values, the 2% of genes with the highest and lowest C. elegans CAI values, and the 2% of genes with highest and lowest RNA-seq
expression in free-living females. Log_{10} counts per million (CPM) expression in free-living adult females was downloaded from the Strongyloides RNA-seq Browser (Bryant et al. 2021). For statistical comparisons of the expression of the highest and lowest Strongyloides-codon-adapted genes, relative to all genes, a 2-way ANOVA (type III) with Tukey post-hoc tests was performed in R using the car package v3.0-8. GO analyses of functional subsets were performed using the gprofiler2 package v0.1.9, with a false discovery rate (FDR)-corrected p-value of <0.05. Commonly enriched GO terms in each subset were defined as GO terms that were enriched in all four of the Strongyloides species (or at least three Strongyloides species in the case of gene-expression-based subsets) with an FDR-corrected p-value of ≤0.001. For statistical comparisons of genome-wide CAI values and fractional GC content between species, Kruskal-Wallis tests with post-hoc Dunn’s tests were performed in R using the dunn.test package v1.3.5. For post-hoc Dunn’s tests, p-values were corrected using the Bonferroni method.

Data availability
Preprocessing and analysis source code, plus codon frequency data, intron sequences, and supplementary files are available at: https://github.com/HallemLab/Bryant-and-Hallem-2021. The following supplementary files have been uploaded to figshare: https://doi.org/10.25387/g3.14462481. Supplementary File S1 contains codon usage frequencies and optimal codons for: highly abundant S. ratti EST transcripts (Mitreva et al. 2006); highly expressed C. elegans, P. pacificus, B. malayi, and N. brasiliensis genes (Sharp and Bradnam 1997; Eccles et al. 2018; Han et al. 2020); and all S. ratti ESTs (Mitreva et al. 2006). Supplementary File S2 contains a code freeze for the Wild Worm Codon Adapter. Supplementary File S3 contains gene IDs, CAI values, GC ratios, and GO term accession numbers of the 2% of genes with the highest and lowest CAI values for each Strongyloides species and C. elegans. Supplementary File S4 contains GO analysis results for the 2% of genes with the highest and lowest CAI values for each Strongyloides species and C. elegans. Supplementary File S5 contains GO terms significantly enriched in all four Strongyloides species for the highest (top 2%) and lowest (bottom 2%) Strongyloides codon-adapted sequences, as well as GO terms significantly enriched in at least three Strongyloides species for genes with the highest (top 2%) and lowest (bottom 2%) expression in free-living females.

Results and discussion
Software functionality
The Wild Worm Codon Adapter app (https://hallemlab.shinyapps.io/Wild_Worm_Codon_Adapter/) features two usage modes: “Optimize Sequences” mode and “Analyze Sequences” mode (Figures 1 and 2, Supplementary File S2). “Optimize Sequences” mode automates the process of transgene codon adaptation and intron insertion. For codon optimization, the app features built-in preferred codons for four non-Caenorhabditis nematode genera for which transgenesis protocols are increasingly well-established: Strongyloides, Nippostrongylus, Priestionchus, and Brugia. The app also includes the option to codon-optimize transgenes for expression in C. elegans (e.g., for researchers wishing to express parasite sequences heterologously in C. elegans), similar to established platforms (Grote et al. 2005; Redemann et al. 2011). This mode also supports custom codon optimization based on a user-provided list of preferred codons.

To generate codon-optimized sequences suitable for expression in Strongyloides, Nippostrongylus, Brugia, Priestionchus, or C. elegans, users select the appropriate codon usage rule and then submit cDNA or amino acid sequences via a text window or file upload. To codon-optimize transgenes for expression in any additional organism of interest, users may prefer to supply a custom list of preferred codons. For example, users wishing to codon-optimize parasite sequences for heterologous expression in mammalian cell culture would upload a list of mammalian optimal codons. The uploaded custom list of preferred codons must use the following format: a 2-column CSV file listing single-letter amino acid symbols and corresponding 3-letter optimal codon sequence; only one optimal codon should be provided per amino acid and stop codons should be designated using the *** symbol. An example custom preferred codon table is available to download from the website.

For Strongyloides species, codon optimization is based on codon usage patterns in S. ratti (Mitreva et al. 2006). Codon optimizations for Brugia, Priestionchus, and Nippostrongylus species are based on codon bias in B. malayi, P. pacificus, and N. brasiliensis, respectively (Eccles et al. 2018; Han et al. 2020). Codon usage is often well conserved between closely related nematode species (Mitreva et al. 2006; Han et al. 2020). Thus, codon usage rules generated from individual species are likely effective across closely related species (e.g., across members of a genus).

We calculated built-in codon usage rules from the codon usage patterns observed in highly expressed genes (Sharp and Bradnam 1997; Mitreva et al. 2006; Eccles et al. 2018; Han et al. 2020), since the codon usage patterns of highly expressed genes are thought to correlate with higher protein expression (Sharp and Li 1987; Plotkin and Kudla 2011). However, the codon usage rules for highly expressed S. ratti genes are extremely similar to those observed across all S. ratti coding sequences (Supplementary File S1). In contrast, the preferred codon usage rules we implement for Strongyloides, Nippostrongylus, Priestionchus, and Brugia are distinct from the C. elegans codon usage rules (Supplementary File S1), consistent with observations that individual transgenes show limited expression across nematode species with divergent genomes (Hunt et al. 2016; Lok et al. 2017; Castelletto et al. 2020; Han et al. 2020).

Previous studies have suggested that highly divergent codon usage patterns between nematodes are driven in part by the extreme AT-richness of some nematode genomes (Cutter et al. 2006; Mitreva et al. 2006; Hunt et al. 2016; Han et al. 2020). As expected, in S. ratti and B. malayi, which have AT-rich genomes, the preferred codons that comprise the built-in usage rules are AT-biased (preferred codon fractional GC content: S. ratti = 0.33, B. malayi = 0.32, N. brasiliensis = 0.54, P. pacificus = 0.55, C. elegans = 0.51). Thus, codon optimization for expression in Strongyloides and Brugia species will likely yield AT-rich optimized sequences. Given the use of a single codon sequence per amino acid and the AT-bias in preferred codons for some species, users may want to eliminate repeated nucleotide patterns, hairpin loops, or unwanted restriction sites prior to final gene synthesis.

To insert introns into optimized cDNA sequences, users select the type and number of introns for insertion into the optimized sequence (up to three). Users may choose between three sets of built-in unique intron sequences: canonical C. elegans artificial introns (Fire et al. 1995), P. pacificus native introns (Han et al. 2020), or Periodic A₅/T₅ Cluster (PATC)-rich introns that enhance germ-line expression of transgenes in C. elegans (Alijohani et al. 2020). Alternatively, users may upload a FASTA file containing a custom set of introns. The app identifies three putative intron insertion
Figure 1 A UI overview of the Wild Worm Codon Adapter app. The overview shows user inputs, content displayed in the browser, and downloadable files. The application features two modes: Optimize Sequences (blue panels) and Analyze Sequences (red panels) that are accessed via separate tabs in the browser window. The app also includes an About tab that presents methods information and a menu for downloading files. Available files include example input files as well as tables of species-specific optimal codons, species-specific codon frequencies, and relative adaptiveness values used to calculate codon adaptation indices.

| User selects operation mode |
|----------------------------|
| Optimize Sequences         |
| Analyze Sequences          |

| User inputs sequence and optimization parameters |
|--------------------------------------------------|
| 1. Use text window or file upload control to load a cDNA or amino acid sequence |
| 2. Use dropdown menu to select a built-in codon usage rule - or - Use file upload control to load a custom list of preferred codons |
| 3. Use dropdown menu to select an intron sequence source - or - Use file upload control to load a custom set of introns |
| 4. Use dropdown menu to select the desired number of introns (0-3) |

| Content Displayed in Browser |
|------------------------------|
| 1. Sequence information for original and optimized sequences: |
| - fractional GC ratio |
| - Codon Adaptation Index (CAI) value relative to selected codon usage rules |
| 2. Original sequence |
| 3. Optimized sequence |
| - without introns |
| - with desired number of introns |

| Downloadable Files |
|--------------------|
| Optimized sequence (.txt file) |
| - without introns |
| - with desired number of introns |

| User inputs gene(s) of interest |
|---------------------------------|
| Use textbox to input sequence - or - Use second textbox to input gene/ transcript IDs, using the formats: |
| - Stable IDs with the prefix: SSTP|SRAE|SVE|SPAL|NBR |
| Bme|Ppa|WB |
| - C. elegans gene names with ‘Ce-’ prefix |
| - C. elegans transcript IDs - or - Use the file upload control to load: |
| - One-column .csv file listing geneIDs |
| - Two-column .csv file listing names and coding sequences |
| - .fasta file containing named coding sequences |

| Content Displayed in Browser |
|------------------------------|
| Sequence information, including: |
| - GC content |
| - Codon Adaptation Index (CAI) value relative to codon usage rules for the following species: |
| - C. elegans |
| - S. ratti |
| - N. brasiiliensis |
| - B. malayi |
| - P. pacificus |

| Downloadable Files |
|--------------------|
| Sequence information (.xlsx file) |
Figure 2: Detailed graphic view of the codebase of the Wild Worm Codon Adapter app. The application features two usage modes: Optimize Sequences (upper, blue panel) and Analyze Sequences (lower, red panel). Gray boxes are preprocessed data inputs generated from published data. Red boxes are user inputs. Green boxes are output elements displayed within the browser; output values directly depend on inputs provided via red elements. Blue boxes are commands run in R. White boxes are code details or input options. Purple boxes are downloadable output file options. Dashed lines show division of code elements into the named files.
sites spaced approximately equidistantly within the optimized cDNA sequence, and inserts the user-specified type and number of introns (Fire et al. 1995; Blumenthal and Steward, 1997; Redemann et al. 2011). In C. elegans and P. pacificus, a single 5' intron is sufficient for intron-mediated enhancement of gene expression, whereas a single 3' intron is not (Crane et al. 2019; Han et al. 2020). Thus, when the user chooses to insert fewer than three introns, the three hypothetical insertion sites are filled as needed, starting from the 5' site.

The app displays the original (non-optimized) sequence and the optimized sequence with and without introns; users may download optimized sequences as plain text (.txt) files. "Optimize Sequences" mode reports the fractional GC content for the original and optimized sequences. When performing optimization using built-in cDNA usage rules, the app also provides a measure of the cDNA usage bias for both the original and optimized cDNA sequences by reporting CAI values relative to the selected usage rule.

Finally, the "Analyze Sequences" mode (Figure 1) was designed to support descriptive analyses of transgene sequences as well as native Strongyloides, B. malayi, P. pacificus, N. brasiliensis, or C. elegans coding sequences. To measure how well codon-adapted a transgene is for expression in Strongyloides, Brugia, Nippostrongylus, Pristionchus, or C. elegans, users may submit the relevant coding sequence(s). For individual sequences, the following values are calculated and displayed as a downloadable table: fractional GC content; and CAI values relative to Strongyloides, Brugia, Nippostrongylus, Pristionchus, and C. elegans cDNA sequences (Figure 3A; p < 0.0001 comparing C. elegans, Brugia, and Nippostrongylus to each remaining species, Kruskal-Wallis test with Dunn’s post-hoc tests). Also consistent with previous observations that Strongyloides genomes are highly AT-rich (Mitreva et al. 2006; Cutter et al. 2006; Hunt et al. 2016), we observed that Strongyloides coding sequences displayed lower fractional GC content than C. elegans, N. brasiliensis, P. pacificus, and B. malayi coding sequences (Figure 3B; p < 0.00005 comparing C. elegans to each remaining species, 2-way ANOVA with Tukey post-hoc tests).

Benchmarking and example usage

To benchmark the use of the CAI to quantify codon adaptiveness of native genes, we used "Analyze Sequences" mode to assess and compare genome-wide codon bias patterns in Strongyloides species, B. malayi, N. brasiliensis, P. pacificus, and C. elegans. Consistent with previous findings (Mitreva et al. 2006), we found that the distribution of genome-wide codon bias varied by genus, such that the genus-specific CAI values of C. elegans coding sequences were lower than those for parasitic species (Figure 3A; p < 0.0001 for all comparisons to C. elegans, Kruskal-Wallis test with Dunn’s post-hoc tests). Also consistent with previous observations that Strongyloides species tend to be more AT-rich than the coding sequences of other species (p < 0.0001 for the grouped Strongyloides species versus each remaining species, Kruskal-Wallis test with Dunn’s post-hoc tests). For both panels, values were calculated by submitting a list of all predicted coding sequences to the Wild Worm Codon Adapter app in Analyze Sequences mode. (C) Violin plot of log2 counts per million (CPM) expression in free-living females for all genes (gray), genes in the top 2% of CAI values for each species (red), and genes in the bottom 2% of CAI values for each species (blue). Dots indicate median values. ***p < 0.00005, 2-way ANOVA with Tukey post-hoc tests. ns = not significant (p > 0.05).

Figure 3 Codon adaptiveness and GC content across species. (A) Density plot of CAI values in a species’ genome. Codon adaptiveness varies across species; C. elegans genome-wide codon adaptiveness is significantly lower than other species (p < 0.0001 comparing C. elegans to each remaining species, Kruskal-Wallis test with Dunn’s post-hoc tests). For each species, codon bias is calculated relative to the genus-level cDNA usage rules (Supplementary File S1). (B) Density plot of fractional GC content across species. Consistent with previous reports (Cutter et al. 2006; Mitreva et al. 2006; Hunt et al. 2016), Strongyloides coding sequences tend to be more AT-rich than the coding sequences of other species (p < 0.0001 for the grouped Strongyloides species versus each remaining species, Kruskal-Wallis test with Dunn’s post-hoc tests). For both panels, values were calculated by submitting a list of all predicted coding sequences to the Wild Worm Codon Adapter app in Analyze Sequences mode. (C) Violin plot of log2 counts per million (CPM) expression in free-living females for all genes (gray), genes in the top 2% of CAI values for each species (red), and genes in the bottom 2% of CAI values for each species (blue). Dots indicate median values. ***p < 0.00005, 2-way ANOVA with Tukey post-hoc tests. ns = not significant (p > 0.05).
Figure 4: Functional enrichment of highly and poorly codon-adapted genes. GO enrichment across Strongyloides species for sequences displaying high and low degrees of Strongyloides codon bias. (A, B) Manhattan plots and table showing GO enrichment for sequences in the top 2% of CAI values for each species. X-axis labels (A) and table Source values (B) indicate the three GO vocabulary categories: biological process (BP), molecular function (MF), and cellular component (CC). Numbered circles (A) and Term ID numbers (B) indicate GO terms that are significantly enriched ($p<0.001$) in all four Strongyloides species. For Manhattan plots, $-\log_{10}(p$-values) along the y-axis are capped if $\geq 16$; this threshold is indicated by a gray dashed line. All GO terms included in (B) are also significantly enriched in the top 2% of C. elegans-codon-adapted C. elegans sequences. Red terms are also significantly enriched among genes with the highest expression (top 2%) in free-living females, in at least 3 Strongyloides species. (C, D) Manhattan plots and table of GO enrichment for sequences that compose the bottom 2% of CAI values. None of the GO terms included in (D) are also significantly enriched ($p<0.001$) in poorly C. elegans-codon-adapted C. elegans sequences or in genes with the lowest expression (bottom 2%) in free-living females, in at least three Strongyloides species. For all plots, $p$-values are FDR-corrected.
grouped together and compared to each remaining species, Kruskal-Wallis test with Dunn’s post-hoc tests). Together these observations emphasize the likely benefit of transgene codon-optimization for promoting successful transgenesis in non-<i>Caenorhabditis</i> species. Finally, consistent with our use of <i>Strongyloides</i> codon usage rules based on highly abundant <i>S. ratti</i> ESTs, the highest and lowest <i>Strongyloides</i>-codon-adapted sequences generally displayed significantly higher and lower expression, respectively, in free-living female life stages relative to the expression of all genes in the genome (Figure 3C). However, the range of gene expression values between the highest and lowest codon-adapted sequences are largely overlapping; thus, the degree of codon adaptation exhibited by individual genes is not the sole determinant of mRNA expression level in these species.

Next, we used GO analyses to assess the putative functions of the most highly or poorly <i>Strongyloides</i>-codon-adapted coding sequences (defined as the top/bottom 2% of Sr-CAI values; Supplementary Files S3 and S4). For the highest <i>Strongyloides</i>-codon-adapted sequences, GO terms that are significantly enriched in all four <i>Strongyloides</i> species are primarily associated with structural integrity and ribosomal components (Figure 4A-B, Supplementary File S5). As expected, given that the <i>Strongyloides</i> codon usage rules are based on highly abundant sequences, there is significant overlap between GO terms enriched in the top 2% of <i>Strongyloides</i>-codon-adapted sequences and GO terms enriched in the genes that are most highly expressed in free-living females (Figure 4B, Supplementary File S5). Despite the differences between codon usage patterns in <i>Strongyloides</i> and <i>C. elegans</i>, GO terms enriched in the top 2% of <i>Strongyloides</i>-codon-adapted sequences are also significantly enriched in the top 2% of <i>C. elegans</i>-codon-adapted sequences (i.e., <i>C. elegans</i> sequences that display high codon usage bias relative to <i>C. elegans</i> codon usage rules) (Figure 4B). In contrast, for parasite sequences that are the least <i>Strongyloides</i>-codon-adapted, commonly enriched GO terms are associated with DNA metabolism and interactions (Figure 4C-D, Supplementary File S5); these terms are not enriched in low-expressing <i>Strongyloides</i> genes or poorly <i>C. elegans</i>-codon-adapted <i>C. elegans</i> sequences (Figure 4D, Supplementary File S5). Although the causes and consequences of codon bias are not fully understood (Sharp et al. 2010; Plotkin and Kudla 2011), our observations of a common set of GO terms enriched in both <i>Strongyloides</i>- and <i>C. elegans</i>-codon-adapted genes suggest that in nematodes, heighted codon bias reflects a nonrandom process that systematically imposes divergent codon usage patterns on genes associated with a common set of biological functions.

Conclusions and future directions
Codon optimization of transgenes significantly improves the likelihood of successful transgenesis in non-<i>Caenorhabditis</i> nematodes. Here, we present the Wild Worm Codon Adapter, a web-based tool designed to replicate and extend the functionality of popular codon-optimization tools to include non-<i>Caenorhabditis</i> nematode genera such as <i>Strongyloides</i>, <i>Pristionchus</i>, <i>Nippostrongylus</i>, and <i>Brugia</i> (Grote et al. 2005; Redemann et al. 2011). The open-source code is extendable; users may perform codon-optimization for an unlimited selection of species by providing a custom list of optimal codons, and additions to the list of built-in optimization rules can be implemented as requested. Going forward, we hope that the Wild Worm Codon Adapter will simplify the process of transgene design for researchers using functional genomics to study non-<i>Caenorhabditis</i> nematodes.

Web resources
A web-hosted version of the app is available at: https://hallem.shinyapps.io/Wild_Worm_Codon_Adapter/

App source code and deployment instructions are available at: https://github.com/HallemLab/Wild_Worm_Codon_Adapter

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Conflicts of interest
None declared.

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