Uridine monophosphate synthetase enables eukaryotic de novo NAD\(^+\) biosynthesis from quinolinic acid

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NAD\(^+\) biosynthesis is an attractive and promising therapeutic target for influencing health span and obesity-related phenotypes as well as tumor growth. Full and effective use of this target for therapeutic benefit requires a complete understanding of NAD\(^+\) biosynthetic pathways. Here, we report a previously unrecognized role for a conserved phosphoribosyltransferase in NAD\(^+\) biosynthesis. Because a required quinolinic acid phosphoribosyltransferase (QPRTase) is not encoded in its genome, Caenorhabditis elegans is reported to lack a de novo NAD\(^+\) biosynthetic pathway. However, all the genes of the kynurenine pathway required for quinolinic acid (QA) production from tryptophan are present. Thus, we investigated the presence of de novo NAD\(^+\) synthesis in this organism. By combining isotope-tracing and genetic experiments, we have demonstrated the presence of an intact de novo biosynthesis pathway for NAD\(^+\) from tryptophan via QA, highlighting the functional conservation of this important biosynthetic activity. Supplementation with kynurenine pathway intermediates also boosted NAD\(^+\) levels and partially reversed NAD\(^+\)-dependent phenotypes caused by mutation of pnc-1, which encodes a nicotinamide-dase required for NAD\(^+\) salvage biosynthesis, demonstrating contribution of de novo synthesis to NAD\(^+\) homeostasis. By investigating candidate phosphoribosyltransferase genes in the genome, we determined that the conserved uridine monophosphate phosphoribosyltransferase (UMPS), which acts in pyrimidine biosynthesis, is required for NAD\(^+\) biosynthesis in place of the missing QPRTase. We suggest that similar underground metabolic activity of UMPs may function in other organisms. This mechanism for NAD\(^+\) biosynthesis creates novel possibilities for manipulating NAD\(^+\) biosynthetic pathways, which is key for the future of therapeutics.

NAD\(^+\) is found in all living cells and is an essential coenzyme that impacts the entire metabolome (1). NAD\(^+\) is involved in redox reactions where it carries electrons from one reaction to another and serves as a substrate for a group of enzymes called NAD\(^+\) consumers that regulate a variety of key biological processes (2, 3). Thus, NAD\(^+\) biosynthesis has proven to be an attractive and promising therapeutic target. However, exactly how NAD\(^+\) homeostasis is maintained and the biological impact of manipulating NAD\(^+\) biosynthetic pathways remain understudied. A complete understanding of the full complement of NAD\(^+\) biosynthetic capacity as well as the consequences to the cell and the organism of manipulation of NAD\(^+\) synthetic pathways is necessary to fully maximize the effectiveness of this target for therapeutic benefit.

A wide range of animals and yeasts synthesize NAD\(^+\) via de novo synthesis from the degradation of tryptophan via the kynurenine pathway (4). Tryptophan degradation typically occurs in the nervous system and liver of most mammals (5, 6). The derivatives of the kynurenine pathway have been commonly linked to both the progression and protection of neurological disorders and neurodegenerative diseases. Quinolinic acid (QA)\(^3\) acts as an agonist to the N-methyl-D-aspartate (NMDA) glutamate receptors (7) and is considered as a neurotoxin (8). In contrast, kynurenic acid (KYNA) acts as an antagonist to a spectrum of amino acid receptors and is considered to be a neuroprotective agent (7). The contrast between QA and KYNA and their roles in neurological disease states indicates the importance of maintaining kynurenine pathway homeostasis for healthy brain function (9). The relationship between NAD\(^+\) de novo synthesis and the kynurenine pathway is important for normal metabolic function and homeostasis.

The genome of Caenorhabditis elegans encodes all the enzymes involved in the kynurenine pathway (Fig. 1A). However, the genome lacks the critical quinolinic acid phosphoribosyltransferase (QPRTase) ortholog that would convert QA into nicotinic acid mononucleotide (NaMN) for biosynthesis of NAD\(^+\). Because the C. elegans lack an apparent QPRTase homolog, it has been assumed that this species lacks active NAD\(^+\) de novo synthesis (2, 11, 12). We predict that organisms lacking a de novo NAD\(^+\) synthesis pathway would be unable to clear the end product of tryptophan degradation, QA, a neurotoxin (7, 8). Thus, we examined the role of de novo synthesis in C. elegans. We discovered that the isotopic label supplied via Trp is incorporated into NAD\(^+\). Blocking the kynurenine pathway lowers global NAD\(^+\) levels compared with controls and prevents incorporation of the isotopic label, supplied via Trp, into NAD\(^+\). We found that phenotypes that are dependent on NAD\(^+\) levels could be reversed upon supplementation with QA

\(^3\)The abbreviations used are: QA, quinolinic acid; 3HAA, 3-hydroxyanthranilate; Kyn, kynurenine; KYNU-1, kynureninase; NA, nicotinic acid; QPR-Tase, quinolinic acid phosphoribosyltransferase; UMPS-1, uridine monophosphate synthetase; F, forward; R, reverse.
and other kynurenine metabolites. Furthermore, we identified a candidate enzyme that is required for incorporation of isotopic label supplied via QA into NAD⁺. Finally, we connected de novo NAD⁺ synthesis to fecundity. This evidence supports the hypothesis that NAD⁺ de novo synthesis is active and contributes to NAD⁺ biosynthetic capacity and homeostasis of C. elegans.

Results

NAD⁺ de novo synthesis contributes to biosynthetic capacity and reproduction

To investigate whether the kynurenine pathway contributes to NAD⁺ de novo biosynthesis in C. elegans, we examined the effect of loss of the pathway on global NAD⁺ levels. kynu-1 encodes kynureninase and is required for conversion of 3-hydroxy-L-kynurenine to 3-hydroxyanthranilic acid (3HAA) and indirectly for formation of QA (13). The kynu-1(tm4924) mutants, which have a deletion of three internal exons of the kynu-1 gene (14), have decreased global NAD⁺ levels compared with controls (Fig. 1B). We conclude that the kynurenine pathway contributes to NAD⁺ homeostasis in C. elegans perhaps by contributing to biosynthetic capacity.

To directly test whether de novo NAD⁺ synthesis from tryptophan via QA occurs, we used isotopically labeled metabolic tracers. After short-term supplementation of cultures with deuterium-labeled Trp, we successfully detected isotope label in the tryptophan pool (Fig. 1C). Furthermore, we detected isotope label in 15% of the QA pool and in an average of 9% of the

![Figure 1. NAD⁺ de novo synthesis contributes to NAD⁺ biocapacity and reproduction. A, schematic of NAD⁺ de novo synthesis in C. elegans. NFK is N-formylkynurenine; 3HK is hydroxy-L-kynurenine. B, LC-MS measurements of relative levels of NAD⁺ in N2 and kynu-1(tm4924) mutant animals. Boxes show the upper and lower quartile values; + indicates the mean value, and line indicates the median. Error bars indicate the maximum and minimum of the population distribution. *, 0.01 < p < 0.05; calculated with Welch’s two sample t test. C–E, percent incorporation of isotope label from d⁵-Trp supplementation into the Trp pool (C), the QA pool (D), and NAD⁺ pool (E) in N2 and kynu-1(tm4924) mutant animals. Dots represent each biological replicate of N2 and kynu-1(tm4924) mutant animals sampled after 4 h of exposure to d⁵-Trp. + indicates the mean value, and error bars are S.D. The peak area for total QA was undetectable in kynu-1 mutants, likely as a result of blocking the biosynthesis pathway. F, ratio of percent isotope label incorporated into Trp, QA, and NAD⁺ in kynu-1(tm4924) mutants compared with N2. Error bars indicate S.D. ***, p < 0.001; calculated with Welch’s two sample t test. G, progeny production is reported for N2 control and kynu-1(tm4924) mutant animals and kynu-1 mutant animals supplemented with 20 mM QA. Error bars indicate S.D. *, 0.01 < p < 0.05, calculated with Student’s t test.](https://example.com/figure1.png)
NAD$^+$ pool (Fig. 1, D and E). This demonstrates the flow of label from Trp to NAD$^+$ via a de novo synthesis pathway. Next, we investigated whether this flow was dependent on an active kynurenine pathway. We predicted that loss of kynu-1 would block incorporation of Trp-supplied label into NAD$^+$. The Trp pool was isotonically labeled in kynu-1 mutants at a level similar to the control (Fig. 1C). As expected, mutation of kynu-1 decreased the efficiency of label incorporation into NAD$^+$ more than 2-fold relative to controls (Fig. 1, E and F). Labeled QA was below the detection limit upon removal of kynu-1 activity (Fig. 1D). This supports our hypothesis that NAD$^+$ de novo synthesis from tryptophan is actively contributing to NAD$^+$ biosynthetic capacity in C. elegans, although the organism lacks a functional QPRTase homolog in genome.

We next asked whether NAD$^+$ de novo synthesis from tryptophan was functionally important by examining fecundity. We observed a decrease in progeny production in kynu-1 mutants compared with controls (Fig. 1G). However, supplementation with QA, the kynurenine pathway end product and the NAD$^+$ de novo synthesis precursor, restored the brood size in kynu-1 mutants, supporting a functional role for de novo NAD$^+$ biosynthesis in fecundity (Fig. 1G).

**Supplementation with de novo precursors reverses NAD$^+$-dependent phenotypes**

We previously reported that reduced levels of NAD$^+$ caused by the lack of salvage synthesis in pnc-1 mutants impair reproductive development in C. elegans (15). Although the levels of nicotinic acid (NA), the product of PNC-1, are reduced almost 20-fold in the mutant, NAD$^+$ levels are only reduced by about 30% (15). Therefore, we hypothesized that alternative NAD$^+$ biosynthetic pathways could respond to the loss of salvage synthesis to maintain global NAD$^+$ levels. To investigate this hypothesis, we examined expression of tdo-2, which encodes the enzyme that catalyzes the rate-limiting step of the kynurenine pathway. We detected a greater than 4-fold increase in tdo-2 transcript levels in pnc-1 mutants (Fig. 2A), supporting an active and functional role for the de novo pathway in NAD$^+$ homeostasis in C. elegans. We further reasoned that an increase in available de novo precursors in combination with the detected up-regulation of tdo-2 might ameliorate pnc-1 phenotype. To test this hypothesis, we supplemented pnc-1 mutant animals with QA, Kyn, and 3HAA. All three supplements improved the reproductive gonad delay phenotype of pnc-1 mutants (Fig. 2B). Supplementation with QA also boosted the global levels of NAD$^+$ in pnc-1 mutants (Fig. 2C). We conclude that boosting de novo synthesis can reverse NAD$^+$-dependent phenotypes in pnc-1 mutants and that de novo synthesis functionally contributes to NAD$^+$ homeostasis.

**UMPS-1 contributes to NAD$^+$ biosynthesis**

If de novo synthesis is indeed active in C. elegans, then how is QA used as a substrate without the QPRTase enzyme? We hypothesized that another phosphoribosyltransferase encoded in the genome would be required to synthesize NAD$^+$ from QA. The C. elegans genome contains seven annotated phosphoribosyltransferases. Interestingly, only UMQS-1 (uridine monophosphate synthetase) has a domain structure similar to QPRTase, with both the phosphoribosyltransferase domain and a carboxylase domain. We specifically asked whether umqs-1 is functionally involved in NAD$^+$ biosynthesis by determining whether it is required for QA supplementation to rescue pnc-1 phenotypes or for incorporation of isotope label supplied via Trp into NAD$^+$. As noted above, QA supplementation rescues gonad developmental delay in pnc-1 mutants. We predicted that if umqs-1 played a role in de novo NAD$^+$ synthesis, then loss of umqs-1 would block the ability of QA but not NA to restore normal gonad development to a proportion of pnc-1 mutants. To test this prediction, we used the deletion allele ok2703, which removes a portion of the 3rd as well as the entire final (4th) exon of the umqs-1 gene. As expected, whereas NA increases the proportion of normal gonad development for both pnc-1 mutants and umqs-1(ok2703);pnc-1 double mutants, QA affects only the pnc-1 mutant but not the umqs-1(ok2703);pnc-1 double mutant (Fig. 3A). Next, we predicted that loss of umqs-1 would lower NAD$^+$ steady-state levels and increase QA steady-state levels if it were participating in de novo NAD$^+$ biosynthesis at the step typically catalyzed by QPRTase. Indeed, global NAD$^+$ levels are decreased and QA levels are increased in umqs-1(ok2703) mutants compared with controls (Fig. 3, B and C). The decrease in NAD$^+$ levels is similar to that observed in kynu-1 mutants (Fig. 1B).

**Figure 2. Supplementation with NAD$^+$ de novo precursors reverses NAD$^+$-dependent phenotypes.** A, ratio of tdo-2 mRNA levels in pnc-1(pk9605) mutants and N2 controls. tdo-2 encodes the enzyme that catalyzes the rate-limiting step in the kynurenine pathway, *, 0.01 < p < 0.05, calculated with Student’s t test. B, supplementation of 20 mM QA, Kyn, and 3HAA to pnc-1(pk9605) mutants restores the gonadal developmental delay. For histograms, error bars are S.D. ***, 0.001 < p < 0.01; ***, p < 0.001; calculated with Fisher’s exact test. C, supplementation of 20 mM QA to pnc-1(pk9605) mutants restores NAD$^+$ levels. LC-MS measurements of relative levels of NAD$^+$ and in N2 and pnc-1(pk9605) mutant. Plots are described in legend to Fig. 1. *p < 0.05; calculated with Welch’s two sample t test.
If UMPS-1 substitutes for the missing QPRTase, we predict that loss of umps-1 would block incorporation of isotope label provided via Trp into NAD\(^+\). We exposed control animals and umps-1\((ok2703)\) mutants to deuterium-labeled tryptophan for 4 h (Fig. 3D), and we measured incorporation of deuterium label into the pool of Trp, QA, and NAD\(^+\). In contrast to loss of kynu-1, umps-1 does not block label incorporation into QA (Fig. 3E), as expected for a gene acting downstream of QA in the pathway. However, umps-1\((ok2703)\) decreased the proportion of the NAD\(^+\) pool that became labeled (Fig. 3F). To ensure that the observed effects associated with the ok2703 deletion were specifically attributable to the deletion within the umps-1 gene, we also examined three additional umps-1 alleles, zw456, tm6379, and mn160, for their effect on isotope label incorporation into NAD\(^+\) and found consistent results (Fig. 3, F and G). This metabolic tracing analysis provides key evidence that de novo NAD\(^+\) synthesis from tryptophan contributes to NAD\(^+\) biosynthesis and that UMPS-1 is required for de novo synthesis. In support of this observation, loss of umps-1\((ok2703)\) disrupts fecundity (Fig. 3H). This further suggests that UMPS-1 can substitute as the QPRTase missing from the C. elegans genome. These data highlight functional conservation of de novo NAD\(^+\) synthesis.
biosynthesis and demonstrate an unexpected flexibility for application of a pyrimidine biosynthesis enzyme in contributing to NAD⁺ biosynthesis.

**Discussion**

NAD⁺ homeostasis is critical for healthy metabolic function and important biological stress responses. Not surprisingly, cells use more than one biosynthetic route for the production of NAD⁺ (1). All of the pathways that contribute to NAD⁺ biosynthetic capacity are also highly conserved throughout evolution, alluding to the importance of NAD⁺ as a cellular hub for metabolism in all organisms (1–3). Our results highlight the conservation of the NAD⁺ biosynthetic pathways and expand our knowledge of comprehensive mechanisms for NAD⁺ biosynthesis.

Despite the absence of what was assumed to be a key enzyme in their genome, *C. elegans* maintain an intact pathway for NAD⁺ de novo synthesis. We used stable isotope metabolic tracer analysis to directly demonstrate metabolite flow from Trp to NAD⁺ via QA, and we demonstrated the contribution of the pathway to maintenance of global NAD⁺ levels as well as function in fecundity. Using stable isotope metabolic tracers combined with genetic analysis, we also demonstrated that umps-1 is required for this de novo synthesis of NAD⁺, suggesting that the well-conserved enzyme uridine monophosphate synthase, known for its primary role in pyrimidine biosynthesis, likely plays a dual role in *C. elegans*. This second role in NAD⁺ biosynthesis highlights the possibility for the contribution of similar underground metabolic activity (16) to NAD⁺ biosynthetic capacity in other organisms, even humans.

One of the reasons we predicted that *C. elegans* may have an active de novo NAD⁺ synthesis pathway is that a method to clear the end product of the kynurenine pathway, QA, a known neurotoxin, would be necessary. It is interesting to note that *Drosophila* also lack a QPRTase ortholog. However, the enzyme 3-hydroxyanthranilic acid dioxygenase, required for production of QA, is also missing from the *Drosophila* genome, suggesting that *Drosophila* do not produce QA and thus may actually lack this otherwise conserved de novo pathway for NAD⁺ biosynthesis. Alternatively, given the importance of de novo synthesis to most metazoans, it would not be surprising to find intact mechanisms for de novo synthesis of NAD⁺ in *Drosophila*.

We have concluded that umps-1 is required for de novo NAD⁺ biosynthesis. However, we also considered the possibility that perturbations to pyrimidine biosynthesis indirectly negatively affect NAD⁺ biosynthesis. Although we have not formally ruled this out, we favor a direct model because we have demonstrated that UMPS-1 specifically acts between QA and NAD⁺ in biosynthesis, but yet is not downstream of nicotinic acid in ameliorating the NAD-dependent gonad delay phenotype of the NAD⁺ salvage biosynthesis mutant *pnc-1*.

We previously reported that salvage NAD⁺ synthesis is critical for the normal progression of gonad development, for full fecundity, and for efficient glucose metabolism in the cytosol (12, 15, 17, 18). Interestingly, loss of NAD⁺ de novo synthesis also resulted in decreased progeny production compared with wild type. Supplementation with NAD⁺ precursors was able to reverse the brood size defect in *knu-1* mutants, supporting the importance of NAD⁺ biosynthesis for normal reproduction. Both de novo NAD⁺ synthesis and salvage NAD⁺ synthesis (17) are involved in fecundity. This highlights the role of maintaining global NAD⁺ biosynthetic capacity for reproduction.

Our results also highlight the relevance of potential homeostatic mechanisms in response to targeting NAD⁺ biosynthetic pathways for therapeutic use. For instance, inhibiting salvage NAD⁺ synthesis can result in increased de novo synthesis. Similarly, targeting the kynurenine pathway in neurological disorders could result in altered NAD⁺ synthesis. Homeostatic interactions among NAD⁺ biosynthetic pathways should be taken into consideration when manipulating NAD⁺ biosynthesis and metabolism for therapeutic benefits.

**Experimental procedures**

*C. elegans* culture and strains

*C. elegans* strains were maintained under standard conditions at 20 °C (19) with *Escherichia coli* OP50 or UV-irradiated OP50 serving as the food source. N2 is the reference control strain. UV-irradiated OP50 plates were prepared using a 999-s exposure in a GS Gene Linker UV Chamber (Bio-Rad) (12, 18). Complete killing of the *E. coli* was confirmed by absence of growth on LB agar after overnight incubation at 37 °C. The following strains and alleles were used: *pnc-1(pk9605)* (12), *knu-1(tm4924)*, and *umps-1(ok2703, mn160, tm6379, and zu456)*. ok2703 also deletes a portion of the N terminus of the neighboring gene, *spg-1*, which encodes an antibacterial caenopore (21). Strains were obtained from the CGC and Mitani Lab/ National BioResource Project, Japan.

Metabolite supplementation

NA (Alfa Aesar, Tewksbury, MA) and QA (MP Biomedicals, Santa Ana, CA) supplementations were performed on culture plates. We added filter-sterilized 25 mM stock solution of NA and QA to UV-irradiated plates and incubated plates at room temperature for 2–3 days to allow chemicals to diffuse before use.

QA, Kyn (Sigma), and HAA (Sigma) supplementations were performed in small-volume liquid culture because of limited availability of supplements. We first plated synchronized L1 animals on UV-irradiated OP50 plates for 24 h. 2–3 plates of synchronized L3 animals were then collected with M9 solution and pelleted. To the pellet we added 5 μl of concentrated heat-killed OP50 culture, supplement stock solution diluted to the experimental concentration, and M9 to a final volume of 100 μl. Stock solutions were as follows: 20 mM QA, 20 mM Kyn, or 20 mM 3HAA. Liquid culture solutions were incubated at room temperature for 48 h with gentle rocking. Finally animals were plated on UV-irradiated OP50 plates, and gonad development was scored when animals reached mid-L4 stage.

Phenotypic analysis, gonad developmental delay

Gonad developmental delay phenotype was scored as reported previously (12). Briefly, mid-L4 stage animals with an open lumen in both the vulva and the uterus were reported as normal. “Delayed” animals are those that do not yet have an
open uterine lumen when the vulva lumen achieves its characteristic mid-L4 stage morphology. We plated synchronized L1 animals on NGM plates of targeted condition, scored the gonad delay when they reached mid-L4 stage, and calculated the percentage of normal animals.

**Brood size**

Young L3 stage animals were individually plated, and production of progeny was counted for 4 days after reaching adulthood.

**Targeted metabolomics**

We performed targeted LC-MS metabolomics analysis at the Metabolomics Core Facility at Pennsylvania State University. ~50 µl of worms were collected in double distilled H2O, flash-frozen in liquid nitrogen, and stored at −80 °C. 15-µl samples were extracted in 1 ml of 3:3:2 acetonitrile/isopropyl alcohol/H2O with 1 µM chloropropamide as internal standard. Samples were homogenized using a Precellys™24 homogenizer. Extracts from samples were dried under vacuum, resuspended in HPLC Optima Water (Thermo Fisher Scientific, Waltham, MA), and divided into two fractions, one for LC-MS and one for BCA protein analysis. Samples were analyzed by LC-MS using a modified version of an ion pairing reversed phase negative ion electrospray ionization method (10). Samples were separated on a Supelco (Bellefonte, PA) Titan C18 column (100 × 2.1 mm, 1.9-µm particle size) using a water/methanol gradient with tributylamine added to the aqueous mobile phase. The LC-MS platform consisted of a Dionex Ultimate 3000 quaternary HPLC pump, a Dionex 3000 column compartment, a Dionex 3000 autosampler, and an Exactive plus orbitrap mass spectrometer controlled by Xcalibur 2.2 software (all from Thermo Fisher Scientific, San Jose, CA). The HPLC column was maintained at 30 °C at flow rate of 200 µl/min. Solvent A was 3% aqueous methanol with 10 mM tributylamine and 15 mM acetic acid; solvent B was methanol. The gradient was 0 min, 0% B; 5 min, 20% B; 7.5 min, 20% B; 13 min, 55% B; 15.5 min, 95% B; 18.5 min, 95% B; 19 min, 0% B; and 25 min, 0% B. The orbitrap was operated in negative ion mode at maximum resolution (140,000) and scanned from m/z 85 to m/z 1000. Metabolite levels were corrected to protein concentrations determined by BCA assay (Thermo Fisher Scientific).

**Metabolic tracing with stable isotopes**

Stable isotope d5-trypophan (Santa Cruz Biochemicals, Dallas, TX) was used as the metabolic tracer. To collect isotopic Trp-treated C. elegans, mixed-stage worms were plated on UV-irradiated OP50 plates and incubate at 20 °C for 72 h. Worms were then collected with M9 solution and pelleted. To the pellet we added 1 ml of concentrated heat-killed OP50 culture, 100 µl of 100 mM isotopic Trp, and M9 to a final volume of 2 ml. Liquid culture solutions were incubated at room temperature for 4 h with gentle rocking. Worms and heat-killed OP50 were separately collected by centrifuging and washed with 15 ml of autoclaved water three times. Approximately 30–40 µl of worm pellets were obtained for each sample. Targeted LC-MS metabolomics analysis was performed to measure isotope incorporation.

**Quantitative real-time PCR**

RNA was extracted from N2 control and pnc-1 mutant animals cultured on OP50 plates using TRIzol Reagent (Life Technologies, Inc.). 2 µg of total RNA, quantified by Nano-Drop NA-1000 spectrophotometer (NanoDrop Technologies, Wilmington, DE), was used for reverse transcription with the High Capacity cDNA Reverse Transcription kit (Applied Biosystems, Foster City, CA). Three genes, cdc-42, pmp-3, and tba-1, were used as internal reference control (20). Real-time quantitative PCR amplifications for test and reference genes were carried out using 7.5 µl of SYBR Green (PerfeCTa SYBR Green Super Mix with ROX, Quanta Biosciences Beverly, MA), 0.6 µl of forward and reverse primers, 1.3 µl of distilled H2O, and 5 µl of diluted cDNA for each sample in a total of 15 µl. Amplification was carried out in a 7300 real-time PCR system (Applied Biosystems, Foster City, CA) with initial polymerase activation at 95 °C for 10 min, followed by 40 cycles of 95 °C for 15 s denaturation, 60 °C for 60 s for primer-specific annealing, and elongation. After 40 cycles, melting curve analysis was carried out (60–95 °C) to verify the specificity of amplicons. The following primers were used: for internal reference genes, cdc-42-F (5’-ctcggactctctgctgacaag-3’) and cdc-42-R (5’-ctcggactctctgctgacaag-3’); pmp-3-F (5’-gttcccgtgttcatcactcat-3’) and pmp-3-R (5’-acaccgtcagaagctgtaga-3’); tba-1-F (5’-gtacactcactgtacctgtctgacaag-3’) and tba-1-R (5’-ctctgttaccagactgact-3’), and for test genes, tdo-2-F (5’-gttcgatgggtttcgg-3’) and tdo-2-R (5’-accaactaacctgtatgagga-3’).

**Statistical analysis**

Fisher’s exact test was carried out to determine p values for the gonad developmental delay phenotype. For quantification of the fecundity defect and gene expression, p values were calculated using Student’s t test. Welch’s two-sample t test to calculate p values was used for LC-MS analysis. In all figures the following were used: $, 0.05 < p < 0.1; *, 0.01 < p < 0.05; **, 0.001 < p < 0.01; ***, p < 0.001.

**Author contributions**—M. R. M., W. W., and W. H. R. conceived and planned the experiments. M. R. M., W. W., and L. H. performed the experiments and analyzed the data. M. R. M. and W. H. R. wrote the manuscript.

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