Disruption of INOS, a Gene Encoding myo-Inositol Phosphate Synthase, Causes Male Sterility in Drosophila melanogaster

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ABSTRACT Inositol is a precursor for the phospholipid membrane component phosphatidylinositol (PI), involved in signal transduction pathways, endoplasmic reticulum stress, and osmoregulation. Alterations of inositol metabolism have been implicated in human reproductive issues, the therapeutic effects of drugs used to treat epilepsy and bipolar disorder, spinal cord defects, and diseases including diabetes and Alzheimer’s. The sole known inositol synthetic enzyme is myo-inositol synthase (MIPS), and the homolog in Drosophila melanogaster is encoded by the Inos gene. Three identical deletion strains (inosΔDF/CyO) were constructed, confirmed by PCR and sequencing, and homozygotes (inosΔDF/inosΔDF) were shown to lack the transcript encoding the MIPS enzyme. Without inositol, homozygous inosΔDF deletion fertilized eggs develop only to the first-instar larval stage. When transferred as pupae to food without inositol, however, inosΔDF homozygotes die significantly sooner than wild-type flies. Even with dietary inositol the homozygous inosΔDF males are sterile. An inos allele, with a P-element inserted into the first intron, fails to complement this male sterile phenotype. An additional copy of the Inos gene inserted into another chromosome rescues all the phenotypes. These genetic and phenotypic analyses establish D. melanogaster as an excellent model organism in which to examine the role of inositol synthesis in development and reproduction.

The metabolism of myo-inositol plays a role in fertilization and early embryonic development, and alterations have been implicated in reproductive issues, cancer, neurodegenerative disorders, spinal cord defects, epilepsy, and bipolar disorder (Croze and Soulage 2013; Bevilacqua et al. 2015; Koguchi et al. 2016; Ye and Greenberg 2015; Yu et al. 2017). Myo-inositol is a six-carbon sugar alcohol found in eukaryotic and many prokaryotic cells. It is the precursor for the phospholipid membrane component phosphatidylinositol (PI), and has important roles in signal transduction, endoplasmic reticulum stress, and osmoregulation (Henry et al. 2014; Chow et al. 2015).

There are three ways an organism can acquire inositol. The first is through inositol transport from the extracellular environment (Spector and Lorenzo 1975). The second is via recycling by dephosphorylation of inositol phosphates (Berridge and Irvine 1984). The third is synthesis via a two-step process from glucose-6-phosphate, for which the first step is catalyzed by myo-inositol-3-phosphate synthase (MIPS) (Loewus and Kelly 1962; Eisenberg et al. 1964). The properties and catalytic mechanisms of MIPS are similar in animals, plants and yeast (Chen and Charalampous 1964; Loewus et al. 1983; Klig et al. 1994; Molina et al. 1999). Currently the genomes of more than one hundred organisms, ranging from microbes to man, contain annotated orthologs which encode MIPS (NCBI). In most organisms it is a homo-tetramer (approximately 62 kD per subunit) (Henry et al. 2014).

High levels (millimolar concentrations) of myo-inositol have been detected in seminiferous tubule fluid of several mammalian species. To establish and maintain this high level, one hundred times more than in plasma, there must be some combination of extremely high levels of synthesis, transport against a large concentration gradient, and a barrier
to loss of inositol (cell-cell contact and unidirectional active transport) (Lewin et al. 1976). In mice higher levels of MIPS mRNA have been demonstrated in the Sertoli cells, pachytene spermatocytes, and the round spermatids of the testis (Chauvin and Griswold 2004). The testicular barrier isolates these cells from exogenous molecules (Lefever et al. 2015, Mruk and Cheng 2015). Similarly, the somatic cyst cells form the barrier in D. melanogaster that tightly encapsulates developing spermatids in later stages of spermatogenesis, preventing transport of extracellular compounds (Fairchild et al. 2015).

D. melanogaster development has been extensively studied. After embryogenesis first-instar larvae emerge and feed nearly continuously for one day, then molt, becoming larger second-instar larvae, which continue to feed for another day. After molting again, they become third-instar larvae, which feed for three days before pupating. Pupal development proceeds for three to five days after which the adults eclose. The typical lifespan of an adult is 20–60 days. The D. melanogaster ovary begins forming during embryogenesis, and the ovary structure develops during pupal stages. In the adult ovary there are typically sixteen to twenty ovarioles. Fourteen identifiable stages of oocyte development are evident in linear order in ovarioles. Upon eclosion, the most mature oocytes of D. melanogaster females are in stage seven (previtellogenic). Approximately one and a half days later the ovarioles of these females will have all fourteen stages present, including mature eggs (Spradling 1993; McLaughlin and Bratu 2015). Male spermatogenesis also begins during embryogenesis. Testes of third-instar larvae are ovoid containing pre-meiotic stage germ-line cells. In pupae, testes elongate, coil, and the first spermatids may also begin to appear. In the adult male, the germline stem cells are located at the apical end of the testes. These cells undergo mitotic division, producing a stem cell and a primary spermatogonial cell, which becomes enclosed in two somatically derived cyst cells. Four mitotic divisions yield sixteen spermatocytes, these then undergo meiosis producing 64 haploid spermatids. Later steps of spermatid maturation include nuclear shaping and individualization. Actin has a major role in the individualization process. Coiling occurs, the mature sperm are released into the testis lumen, and then transferred to the seminal vesicle for storage until they are used for fertilization (Fuller 1993, White-Cooper 2009).

High throughput expression analyses reveal that Inos, the gene encoding D. melanogaster myo-inositol-3-phosphate (MIPS), is highly expressed in testes and the head (Chen et al. 2014). The D. melanogaster Inos gene is located on chromosome 2 at band 43C3. A D. melanogaster Inos cDNA was cloned and expressed yielding a 565 amino acid MIPS protein with a molecular weight of 62.3 kD (Park and Kim 2004).

In the current study, myo-inositol synthesis and its role in growth, development, and reproduction, were explored in the model organism D. melanogaster. A precise deletion of the sole inositol biosynthetic gene, Inos, was generated. MIPS mutants have been reported in plants (Donahue et al. 2010) and many unicellular organisms (Klig et al. 1994; Molina et al. 1999); this may be the first report examining the phenotype of a MIPS gene deletion in animals. Homozygous inos<sup>df</sup> deletion embryos hatch, but without dietary inositol, they die within a few hours as first-instar larvae. If pupae grown with inositol are transferred to food without inositol, homozygous inos<sup>df</sup> deletion adults have a significantly shorter lifespan than wild-type flies. Surprisingly, even on rich food with dietary inositol the deletion homozygotes (inos<sup>df</sup>) are sterile. No sperm were observed in their seminal vesicles. The physiological basis of this sterility was further examined. A rescue construct containing an additional copy of the Inos gene on another chromosome restores the growth, development, and viability of homozygous deletion larvae and adults on food without inositol. The male sterile phenotype was also complemented by this construct. These studies contribute to the understanding of the role of inositol synthesis in growth, development, and reproduction.

**MATERIALS AND METHODS**

**Fly stocks and maintenance**

Flies were maintained in standard laboratory conditions at 18° or 25° and 70–80% humidity on a 12hr:12hr light-dark cycle. Stocks from the Bloomington Drosophila Stock Center include Canton-S (#1, CS), Oregon R (#5, OR), P[hsFLP]y w; snal<sup>sn</sup> /CyO (#1929), y<sup>2</sup> w; P[Su/Pol-P] Inos<sup>KG10769</sup>CyO, S<sup>bw</sup> y<sup>1</sup>; ryp<sup>y</sup> #14921, hereafter identified as P-inos<sup>KG10769</sup>, w<sup>+</sup> snal<sup>sn</sup> /CyO, S<sup>bw</sup> y<sup>1</sup> #13198, and w<sup>+</sup> snal<sup>sn</sup> /CyO, S<sup>bw</sup> y<sup>1</sup> P[ActGFP w]<sup>CC2</sup> #9325, hereafter identified as CyO<sup>GFP</sup>. Stocks from Exelixis Harvard Medical School include w<sup>+</sup>; P[XP]d00881 (#D00881, hereafter identified as D[XP+]) and w<sup>+</sup>; P[WH]#00895 (#F00895, hereafter identified as F[WH-]).

All fly stocks were grown on either rich food (https://bdsc.indiana.edu/information/recipes/bloomfood.html) or chemically defined food. Inositol-free chemically defined food was prepared by merging the protocols of Sang (1955), Falk and Nash (1974), and Grandison et al. (2009). Vitamin mix and a stock solution of amino acids were used (Grandison et al. 2009) as follows. Various concentrations of inositol (0, 0.5mM, 170mM) were added as indicated in the text and figures. One hundred milliliters (mls) of defined food were prepared by dissolving 0.613g vitamin mix in 45ml of autoclaved agar solution (3g/100ml), adding 10ml of 10x amino acids stock, choline to 1mM and sucrose to 0.2M. 300μL of 30% Tegosept was added to the 100mL of food.

**Fertility and Development Tests**

Fertility tests were performed with 2-5 day old flies, transferred to the food indicated and allowed to mate for twenty-four hours. GFP-marked CyO balancers were used to distinguish heterozygotes (inos<sup>df</sup>/CyO<sup>GFP</sup>) from deletion homozygotes (inos<sup>df</sup>/inos<sup>df</sup>, GFP-negative) through-out development. The number of eggs laid within the next twenty-four hours, and the number of eggs hatched within subsequent forty-eight hours, were recorded. Progeny were monitored until adulthood. Larval stages were determined using mouthhook morphology. GFP was visualized using a Nikon SMZ1500 dissecting microscope with epifluorescence using a GFP-B filter cube. A fixed number of males and females (4:6) were maintained in each vial for the survivorship experiment.

**Generation of Inos Gene Deletion**

MIPS deletion flies were generated through the use of the FLP-FRT system. Fly strains with FRT containing inserts that flank the MIPS gene were purchased from the Exelixis Collection at the Harvard Medical School. The deletion strains (inos<sup>df</sup>) were generated by excising the DNA between the elements (Parks et al. 2004). The genetic selection of the deletion strains is described in the results section.

**Creating a Rescue Strain With a Second Copy of the Inos Gene**

A genomic BAC clone (CH322-61014) CHORI (Children’s Hospital Oakland Research Institute), harboring a section of chromosome 2 from 596 base pairs upstream of the Inos gene to 13,275 base pairs downstream (P[acman] Resources Genome Browser, http://flypush.imagen.tmc.edu/cgi-bin/gb2/gbrowse/genesac; Venken et al. 2009), was microinjected and inserted at an attP site on chromosome 3 (Genetics Services Inc.). A homozygous rescue strain (w<sup>+</sup>; inos<sup>df</sup>/inos<sup>df</sup>; res/res) was created by crossing the Inos rescue chromosome into the inos<sup>df</sup> deletion strain.
DNA was extracted (Huang et al. 2009) and PCR amplified using primers for the WH- (CCTCGATATAACGAGCTAAAAAC) and XP+ (TACTATTCCTTTCACTCGCACTTATTG) elements (Parks et al. 2004). Primers were designed for the Inos upstream flanking genomic DNA (GAGCTAGTGGGAAATGCAAGG) and downstream genomic DNA (ATTCGGTTAGTTCCCGCCAG). The bands were excised using QIAquick Gel Extraction Kit (Qiagen, Germantown, MD), and sequenced (Eurofins Operon).

**RT-PCR**

Total RNA was extracted from 2-5 day old flies grown on rich food using Trizol (Life Technologies) (Green 2012). Total RNA (5 μg) was DNase treated using the DNA-free Kit (Ambion) and then reverse transcribed with Moloney Murine Leukemia Virus Reverse Transcriptase (M-MLV RT) (Promega), and in other experiments with Avian Myeloblastosis Virus Reverse Transcriptase (AMV-RT) (Promega). The following primers were used: forward Inos 5' UTR TTCCAGAAGCAGACCGAAATA and reverse Inos 3' UTR TTTGATGTCTTTATCAAGTCAAG, forward Rp49 GGATCCAGCTTACGCTTAC and reverse Rp49 CGGAACTTCTTTGAAATCG. Size standards used were 1kb (Promega) or 100bp (Fisher). RT-PCR products were visualized with agarose gels (0.5–2%). The bands were excised using QIAquick Gel Extraction Kit (Qiagen, Germantown, MD), and sequenced (Eurofins Operon).

**Drosophila melanogaster ovary and seminal receptacle dissection and mounting**

All of the females were six to nine days old when dissected. Non-virgins were allowed to mate from eclosion for nine days. Ovaries and seminal receptacles were dissected and washed in Ringer’s Solution (128mM NaCl, 5mM KCl, and 2mM CaCl2). Seminal receptacles were examined, using differential interference contrast microscopy with a 40X objective, for 2-4 females per strain or condition. Approximately twenty ovarioles per ovary were examined from 3-8 females per strain or condition. Ovaries and seminal receptacles were viewed on a Nikon E600 microscope and images were captured using a Retiga 2000R camera (Q imaging). Entire ovaries were scanned and the number of egg chambers at each stage was tabulated (Spradling 1993).

**Drosophila melanogaster Testes Dissections, Mounting, and Staining**

For phase contrast imaging, testes from a minimum of five males per strain were dissected in Hoyle’s medium (Baccetti et al., 1979), mounted in 1X PBS, and viewed using an EVOS Fl Auto Imaging System. For confocal microscopy, testes were dissected, fixed with 4% formaldehyde in 1X PBS, and stained with rhodamine phalloidin (Cytoskeleton, Inc.) at 1:1000 in 1X PBS (Bonaccorsi et al. 2011). At least thirteen males per strain were dissected, typically both testes were examined. The testes were placed in depression slides with approximately 20μl of 2μg/ml DAPI in 1X PBS. Samples were viewed using an Olympus FV 1000 fluoroview confocal laser scanning biological microscope.

**Data Availability**

Strains available upon request. All the relevant data are within the paper.
Statistical analyses

Standard error was calculated for all experiments. The p-values were determined using student’s t-test for the data in figures 3 and 4, the Mantel-Cox log-rank test was used to analyze the data in figure 5, and chi-square analyses were used to analyze the data in figure 6.

RESULTS AND DISCUSSION

High levels of inositol synthesis, via myo-inositol-3-phosphate synthase (MIPS), have been demonstrated in the male reproductive organs of D. melanogaster and many mammals (Lewin et al. 1976; Loewus et al. 1983; Chauvin and Griswold 2004; Chintapalli et al. 2007). Inositol supplementation is known to reduce sterility, enhance early embryonic development, and reduce the incidence of neural tube defects in mammals (Ting et al. 2003; Alebousi et al. 2009; Colazingari et al. 2014). Together this suggests that MIPS may play a role in reproduction and development.

Inos gene deletions were constructed and confirmed

To directly test the involvement of MIPS in reproduction and development, the FLP-FRT system was used to generate a deletion of the Inos gene (which encodes MIPS) of Drosophila melanogaster. Initial evidence that recombination occurred, and that the Inos gene was deleted, was the appearance of dark red-eyed and white-eyed progeny when heat shocked hs-FLP males containing both the FRT-bearing insertion elements D{XP+} and F{WH-} were crossed to white-eyed (w1118) females. The insertion elements, initially in trans in males prior to recombination, are oriented so that recombination at the FRTs will generate chromosomes with two copies of the mini-white gene (deleting Inos) or no copies of the mini-white gene (with two copies of Inos). Three independently isolated putative deletion strains were recovered and outcrossed twice to white-eyed flies (w1118/w1118, snpCyO), and the dark red eyed curly-winged progeny were retained. PCR and sequencing results confirmed the generation of three identical inos4DF deletion strains. The 4,663 bp deletion extends from 40 bases upstream of the beginning of the 5’ UTR to 670 bases downstream of the end of the 3’ UTR (Figure 1).

Inos deletion strains lack Inos transcript

RT-PCR was performed using primers designed to the 5’ and 3’ UTRs of the Inos transcript with poly(A)+ RNA from wild-type (OR), Inos rescue strain (homozygous deletion with an extra Inos gene on chromosome 3 (inos4DF/inos4DF; res/res)), homozygous deletion (inos4DF/inos4DF), or homozygous P-inoSK07679/P-inoSK07679 flies (Figure 2). Primers to reverse transcribe and amplify Rp49 (ribosomal protein L32) poly(A)+ RNA were used simultaneously to serve as internal experimental and loading controls. The wild-type flies express at least three different Inos transcripts, one major and multiple minor (Figure 2). All of the Inos transcripts evident in wild-type flies were absent from the homozygous deletion (inos4DF/inos4DF) or homozygous P-inoSK07679/P-inoSK07679 strains, and were present in the Inos rescue strain (inos4DF/inos4DF; res/res).

Figure 3 Lack of Dietary Inositol Does Not Affect Hatching Rate. The percentage of eggs hatched is indicated by the bars, and the number of eggs monitored (n) is indicated for each cross. Mean ± SE of five independent trials are represented. P > 0.6. As expected, since CyGFP/CyGFP is lethal, the heterozygote hatching rate is approximately 75%.

Figure 4 inos4DF deletion homozygotes die as first-instar larvae on chemically defined food without inositol. Percent of homozygotes (inos4DF/inos4DF, GFP-negative) of the total population at each developmental stage. Mean ± SE of three independent trials are represented. P = 0.004.
Figure 5: Homozygous inos^{ADF} deletion flies die without dietary inositol on chemically defined food. Homozygous inos^{ADF} deletion flies and wild-type (CS) flies grown on chemically defined food with 0.5mM inositol (top panel, \( P = 0.302 \)) or without inositol (bottom panel, \( P = 8.23 \times 10^{-19} \)). Mean ± SE of three independent trials are represented.

(Figure 2). One major and multiple minor Inos transcripts were observed when wild-type (CS or OR) cDNA was amplified with either MMLV-RT or AMV-RT. Although RNA (splice) isoforms detected with a single reverse transcriptase may be artifacts, Yu et al. demonstrated that transcripts detected with both MMLV-RT and AMV-RT prove to be consistently validated RNA isoforms (Yu et al. 2014). The biological role of these RNA isoforms remains to be explored. The dominant band (~1.8 kb) was sequenced and aligned to the expected Inos transcript (99.9% identical). This transcript encodes the full-length MIPS protein expressed by Park and Kim (2004).

Inos deletion affects development and longevity

Without dietary inositol

To determine if inositol is essential for development, progeny of heterozygous deletion flies were grown with or without dietary inositol. Heterozygotes (inos\(^{ADF}/\)CyO\(^{GRF}\)) with GFP-marked CyO balancers were distinguished from deletion homozygotes (inos\(^{ADF}/\)inos\(^{ADF}\), GFP-negative) throughout development. In these experiments, heterozygotes (inos\(^{ADF}/\)CyO\(^{GRF}\)) were mated on chemically defined food. Absence of dietary inositol did not affect the proportion of eggs that hatched \(( P > 0.6 \), Figure 3\). To determine if later development was affected, 237 eggs on chemically defined food with inositol and 349 eggs on chemically defined food without inositol were examined from deposition to adulthood. As expected, homozygous and heterozygous deletion progeny survived on the chemically defined food with inositol. The heterozygous deletion strain inos\(^{ADF}/\)CyO\(^{GRF}\) survived to adulthood without dietary inositol, as did the Inos rescue strain (inos\(^{ADF}/\)inos\(^{ADF}\), GFP-negative), the proportion of which are displayed at each developmental stage. The remainder of the progeny (totaling 100% at each stage), GFP-containing heterozygotes inos\(^{ADF}/\)CyO\(^{GRF}\) and CyO\(^{GRF}/\)CyO\(^{GRF}\) homozygotes, are not shown. Deletion homozygotes (inos\(^{ADF}/\)inos\(^{ADF}\)) only survived to the first-instar larval stage on chemically defined food without dietary inositol. No homozygous inos\(^{ADF}/\)inos\(^{ADF}\) deletion (GFP-negative) second-instar larvae were observed on chemically defined food without inositol \(( P = 0.004 \) ).

Sufficient inositol or Inos transcript may have been transferred to the eggs allowing for survival of the deletion homozygotes through embryogenesis to the first-instar larval stage. Maternal transfer of Inos transcript was ruled out by other experiments in which homozygous inos\(^{ADF}/\)inos\(^{ADF}\) deletion females were mated to heterozygous inos\(^{ADF}/\)CyO\(^{GRF}\) males on rich food supplemented with inositol. All the progeny survived on this rich food supplemented with inositol, demonstrating that Inos transcript is not essential for development. So maternally supplied inositol, without Inos transcript, is sufficient to support development to the first larval-instar stage. Although the precise cause of death during the first-instar stage is not known, inositol is likely to be needed for membrane biogenesis and/or signal transduction.

The longevity of homozygous inos\(^{ADF}\) deletion flies was compared to wild-type (CS) flies. Pupae were transferred to chemically defined food with inositol. The percentage of eggs hatched is indicated by the bars, and the number of eggs monitored \(( n \) ) is indicated for each cross. Mean ± SE of three independent trials are represented.
A total of approximately two hundred adults were examined, at least forty for each strain on each food. As expected, the life span of homozygous inosDDF deletion flies was significantly greater on chemically defined food with inositol than without inositol (Figure 5). Even without inositol most of the homozygous inosDDF deletion flies survived nearly a week. This was not surprising, as the inositol accumulated by larvae grown on rich food could support survival but would eventually be

| Cross         | Stages 1-8 | Stages 9-12 | Stage 13 | Stage 14 |
|---------------|------------|-------------|----------|----------|
| CS♀ x CS♂     | 61% (±8%)  | 34% (±6%)   | 2% (±1%) | 3% (±3%) |
| Virgin CS♀    | 45% (±6%)  | 4% (±3%)    | 1% (±0%) | 50% (±9%)|
| CS♀ x inosDDF | 47% (±7%)  | 3% (±1%)    | 0% (±0%) | 50% (±7%)|

Figure 7 Ovarioles of females mated to deletion males look virgin-like. Dissected ovarioles from wild-type (CS) females imaged with differential interference contrast microscopy. A) mated to wild-type (CS) males, B) virgin, C) mated to homozygous deletion males (inosDDF / inosDDF), D) mated to Inos rescue strain (inosDDF / inosDDF;res/res) males, and E) drawn images of oogenesis stages and table displaying percentage of each stage present in entire ovaries. The number of ovarioles per ovary is similar among the strains, fewer are shown in this figure for image clarity. Mean ± SE of three independent trials are represented. Scale bars correspond to 100 μm.

Figure 8 Testes of homozygous inosDDF deletion males are smaller than wild-type males’ testes. Phase contrast microscopy (representative images) of testes dissected from 2-day-old (A) wild-type (CS), (B) homozygous deletion (inosDDF / inosDDF), (C) Inos rescue strain (inosDDF / inosDDF;res/res), and (D) homozygous P-inosG307677/P-inosG307677 male flies. Scale bars correspond to 400 μm.
Figure 9 Homozygous inos^{ADF} deletion males lack sperm in the seminal vesicles. Microscopy of testes dissected from 2-day-old male flies. Rows 1-4, confocal imaging of samples stained with DAPI (blue) and rhodamine phalloidin (red). Top row (apical end) scale bars are 50 μm (A-D), second
male fertility (McElroy et al. 2013). Equal numbers of males and females were examined and no differences were observed. The p-value of $8.23 \times 10^{-16}$ indicates that the difference in life span between homozygous inos^{ADF} deletion and wild-type (CS) flies on chemically defined food, without inositol, is statistically significant. The rescue Inos construct restored wild-type like development, growth, and survival to homozygous inos^{ADF} deletion flies grown on chemically defined food without inositol. There is no significant difference between survival of homozygous inos^{ADF} deletion and wild-type (CS) flies on chemically defined food supplemented with inositol ($P = 0.302$, Figure 5).

Deletion flies fail to establish a homozygous stock even With dietary inositol

Although rich food with inositol restored viability, the homozygous inos^{ADF} deletion flies failed to establish a homozygous stock. To determine whether the males, females, or both were sterile, homozygous inos^{ADF} deletion, heterozygotes, and wild-type (CS) flies were mated on rich food supplemented with 170mM inositol. Deletion homozygous females mated to wild-type (CS) males laid eggs that hatched normally. Some other genes is known to overcome this block and, although no statistically significant difference was observed, some other genes is known to overcome this block and, although no statistically significant difference was observed.

P-element insertion into the Inos gene also causes male sterility even With dietary inositol

The phenotype of a strain containing a P-element in the first intron of the Inos gene was compared to the inos^{ADF} deletion strain. Similar to the homozygous inos^{ADF} deletion flies, the life span of homozygous P-element (P-inos^{KGO7679}/P-inos^{KGO7679}) deletion flies was significantly greater on chemically defined food with inositol than without inositol (data not shown). Moreover, both the homozygous inos^{ADF} deletion males and the homozygous P-inos^{KGO7679} males were sterile. None of the eggs hatched when homozygous P-inos^{KGO7679} females were mated to wild-type (CS) males (data not shown). Moreover, both the homozygous inos^{ADF} deletion flies and the homozygous P-inos^{KGO7679} males were sterile. None of the eggs hatched when homozygous P-inos^{KGO7679} females were mated to wild-type (CS) males 92% of the eggs hatched (Figure 6).

Wild-type females mated to deletion males have virgin-
Like ovaries and seminal receptacles

After growth on rich food, ovaries dissected from wild-type (CS) females mated to homozygous inos^{ADF} deletion males exhibited a phenotype similar to virgin wild-type (CS) females (Figure 7). In virgin wild-type females, mature stage 14 oocytes accumulate, oocyte development becomes blocked, and stages 9 through 13 of oogenesis seem to disappear (Spradling 1993). To quantify the proportion of each stage, entire ovaries were examined and the number of egg chambers was tabulated (Figure 7E). The proportion of stages 9 through 13 was greatly reduced in the ovarioles of wild-type (CS) females mated to the homozygous inos^{ADF} deletion males, similar to wild-type virgin females. In the reciprocal experiments, when homozygous inos^{ADF} deletion females, also grown with dietary inositol, were mated with wild-type (CS) males, normal ovary morphology was observed. These data suggest that mating to homozygous deletion males does not release wild-type females from the previtellogenic block observed in virgins that have accumulated mature oocytes. This is particularly interesting since other studies have shown that mating to males with sterility caused by mutations in some other genes is known to overcome this block and, although no progeny are produced, the females no longer have virgin-like ovaries (Spradling 1993).

Homozygous inos^{ADF} deletion males were observed to exhibit normal mating behavior including copulation. Since the eggs laid by wild-type (CS) females mated to these males failed to hatch (Figure 6), they were examined by confocal microscopy after staining with DAPI. These eggs did not appear to be fertilized (data not shown) so the transfer of sperm was examined next. No sperm were apparent in the seminal receptacle of wild-type (CS) females after copulation with homozygous inos^{ADF} deletion males, yet sperm were readily apparent when mated to wild-type (CS) males. In addition, deletion of Inos may alter some of the seminal fluid components (Heifetz et al. 2001; Chow et al. 2015). In many organisms the unfolded protein response pathway, which alleviates endoplasmic reticulum (ER) stress, is known to be activated by inositol deprivation. The sterility associated with ER stressed D. melanogaster males, however, is different than the phenotype described here. The ER stressed males have normal sperm production and transfer (Chow et al. 2015) yet (as described above) the homozygous inos^{ADF} deletion males in this study do not have normal sperm transfer with no sperm evident in the seminal receptacles of their mating partners.

Testes From deletion flies lack sperm in the seminal vesicles

To determine whether spermatogenesis was disrupted, testes of wild-type (CS), homozygous deletion (inos^{ADF}/inos^{ADF}), homozygous deletion with an extra Inos construct on chromosome 3 (inos^{ADF}/inos^{ADF};
res/res), and homozygous P-inos\textsuperscript{KCG07679} line P-inos\textsuperscript{KCG07679} males were dissected. Testes of homozygous deletion (inos\textsuperscript{ADf}/ inos\textsuperscript{ADf}) and homozygous P-inos\textsuperscript{KCG07679}P-inos\textsuperscript{KCG07679} males were overall smaller and more fragile (Figure 8). No mature sperm were detected in the seminal vesicles of homozygous inos\textsuperscript{ADf}/ inos\textsuperscript{ADf} or homozygous P-inos\textsuperscript{KCG07679}P-inos\textsuperscript{KCG07679} males (Figure 90, P, S, T). This phenotype is completely penetrant. Both homozygous deletion flies (inos\textsuperscript{ADf}/ inos\textsuperscript{ADf}) and homozygous P-inos\textsuperscript{KCG07679}P-inos\textsuperscript{KCG07679} flies have irregularly shaped spermatid bundles (Figure 9K, L), indicating a disruption of individualization. In both homozygous mutant stai ns, the first visible defect is in the alignment of the nuclei and the failure of the actin cones to form (Figure 9G, H). The actin cones are extremely rarely visible, and never seen in a bundle. Three of eleven homozygous inos\textsuperscript{ADf} deletion males show only a single cone in one of the testes, the rest show no actin cones. In contrast, in every wild-type (CS) control testis many bundles of approximately 30-64 actin cones are evident in the plane of the confocal images. Actin cones are usually found in tight parallel bundles, which promote the maturation of spermatids. During this process, actin cones remove excess cytoplasm and organelles while enclosing each spermatid in its own membrane (Couderc et al. 2017, Isaji et al. 2011, Steinhauser 2015). Disruption of myo-inositol synthesis results in nearly no actin cone formation. Consistent with the lack of actin cones, no waste bags were ever observed in tests of either homozygous mutant strain. The tests of homozygous deletion male flies with the Inos rescue construct on chromosome 3 (inos\textsuperscript{ADf}/inos\textsuperscript{ADf}; res/res) (Figure 9B, F, I, N, R) appear similar to wild-type (CS) (Figure 9A, E, I, M, Q), with sperm evident in the seminal vesicles.

The lack of sperm in the seminal vesicles in homozygous inos\textsuperscript{ADf} deletion males is consistent with failure at individualization. Many male sterile mutants of Drosophila melanogaster exhibit a “classical” phenotype of abnormalities at individualization, which reflects the arrest of spermatogenesis at a checkpoint (Wakimoto et al. 2004). So, the defect due to lack of inositol might be at an earlier stage of spermatogenesis.

In D. melanogaster spermatogenesis, primary spermatogonial cells begin their descent from the tip of the testis and are enclosed in a pair of cyst cells that serve as somatic support cells and restrictive permeability barriers. Testicular barriers are highly selective to prevent pathogens or possible mutagens from harming developing sperm. The soma-germ-line barrier provided by this cyst cell encapsulation becomes increasingly stringent as the sperm progress through development (Fairchild et al. 2015). It appears that while encapsulated, the developing spermatogonial cells cannot use nutritionally acquired myo-inositol, forcing dependence on MIPS. Without inositol synthesis, and unable to access ingested inositol due to a testicular-like barrier, the homozygous deletion flies had no sperm in their seminal vesicles and were sterile.

**BROADER IMPLICATIONS**

In humans, two common causes of infertility are oligoasthenoteratozoospermia and polycystic ovary syndrome (PCOS). The addition of myo-inositol to oligoasthenoteratozoospermia sperm has been reported to correct their morphological abnormalities and to increase their ability to fertilize eggs in vitro (Bevilacqua et al. 2015; Simi et al. 2017). Folic acid from women with PCOS has been shown to have a significantly reduced level of myo-inositol. Supplementation of these women with myo-inositol enhances their egg quality and fertility (Unfer et al. 2011; Colazinari et al. 2013; Nas and Tuu 2017). Clearly these human fertility conditions are related to inositol metabolism. This study demonstrates the requirement for inositol in reproduction in Drosophila melanogaster, and validates this organism as a model for the further examination of the role myo-inositol synthesis in reproduction.

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