**CHES-1-like, the ortholog of a non-obstructive azoospermia-associated gene, blocks germline stem cell differentiation by upregulating Dpp expression in *Drosophila* testis**

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**ABSTRACT**

Azoospermia is a high risk factor for testicular germ cell tumors, whose underlying molecular mechanisms remain unknown. In a genome-wide association study to identify novel loci associated with human non-obstructive azoospermia (NOA), we uncovered a single nucleotide polymorphism (rs1887102, \( P=2.60 \times 10^{-7} \)) in a human gene FOXN3. FOXN3 is an evolutionarily conserved gene. We used *Drosophila melanogaster* as a model system to test whether CHES-1-like, the *Drosophila* FOXN3 ortholog, is required for male fertility. CHES-1-like knockout flies are viable and fertile, and show no defects in spermatogenesis. However, ectopic expression of CHES-1-like in germ cells significantly reduced male fertility. With CHES-1-like overexpression, spermatogonia fail to differentiate after four rounds of mitotic division, but continue to divide to form tumor like structures. In these testes, expression levels of differentiation factor, Bam, were reduced, but the expression region of Bam was expanded. Further reduced Bam expression in CHES-1-like expressing testes exhibited enhanced tumor-like structure formation. The expression of daughters against dpp (*dad*), a downstream gene of dpp signaling, was upregulated by CHES-1-like expression in testes. We found that CHES-1-like could directly bind to the dpp promoter. We propose a model that CHES-1-like overexpression in germ cells activates dpp expression, inhibits spermatocyte differentiation, and finally leads to germ cell tumors.

**INTRODUCTION**

Testicular germ cell tumors (TGCTs) are the most common cancer among young men in industrialized countries [1]. TGCTs are thought to be derived from germ cell lineage cells that are blocked in differentiation and maturation [1]. The causative genetic aberrations were rarely identified. Genome-wide association study (GWAS) studies revealed polymorphic loci linked to the KIT/KITLG [2, 3], RAS [4], and steroid signaling pathways [5, 6]. However, the molecular mechanisms underlying TGCTs remain poorly understood.

*Drosophila* and human testes share many features of spermatogenesis [7, 8]. Many mutants of homologous human and *Drosophila* genes exhibit similar testicular phenotypes. The adult fly testis is a blind tube that opens into the seminal vesicle and ejaculatory duct [8]. The apical tip of the tube is a cluster of somatic cells called hub cells. Eight to ten germ line stem cells (GSCs) are tightly associated with the hub cells, and each is enveloped by two cyst-stem cells (CySCs). Each GSC divides asymmetrically to maintain one cell associated with the hub as a GSC, and another to leave the niche and become a primary spermatogonial cell. Spermatogonial cells...
undergo four rounds mitosis before further differentiation, and then enter meiosis and mature into spermatids [9].

The self-renewal and differentiation of early germ cells in flies are tightly controlled [9]. Similar to humans, flies also develop testis tumors when germ cells fail to differentiate and over-proliferate [10]. Janus kinase–signal transducer and activator of transcription (JAK–STAT) and bone morphogenetic protein (BMP) signaling are critical for GSC maintenance [8, 9]. Malfunction of these two pathways could lead to testis tumors in flies. Hub cells secrete Unpaired (Upd) to bind receptor Dormless on GSCs and CySCs, which activates JAK–STAT signaling and maintains germline and somatic stem cell self-renewal [11, 12]. The ectopic expression of Upd in GSCs results in testis tumors with a massive accumulation of undifferentiated GSC-like cells [11, 12].

Two BMP-like molecules, Dpp and Gbb, expressed in hub and cyst cells are required for GSC maintenance [13–15]. Dpp and Gbb are received by GSCs, where they repress the expression of the differentiation factor, Bag-of-marbles (Bam) [13–15]. Bam and its regulator, Benign gonadal cell neoplasm (Bgen), are required for restricting proliferation of mitotically amplifying spermatogonia [16, 17]. Mutations in bam or bgen lead to testis tumors with extensive proliferation of undifferentiated germ cells [18, 19]. Since BMP signaling could repress bam expression, ectopic expression of dpp in germ cells leads to reduced bam expression and the formation of tumor-like structures in testis [13, 15]. Despite its important functions in fly spermatogenesis, BMP signaling is also required in testis development and spermatogenesis in mammalian systems [20]. Aberrant BMP signaling was reported in human samples with TGCTs [21]. Therefore, investigation of germ cell differentiation in flies might provide insight into potential mechanisms for human TGCTs.

Our previous work has successfully used Drosophila testis as a model system to evaluate the possible loci associated with a severe symptom of male infertility: non-obstructive azoospermia (NOA) [22, 23]. We found two loci near DMRT2 and DMRT3, two genes encoding a DM domain containing a transcription factor, were associated with NOA [22]. Interestingly, DMRT1, the paralog of DMRT2/3, was found to be associated with human TGCTs [24, 25]. Studies showed that the risk of TGCTs was increased in azoospermia or subfertile patients [1, 26–28]. Therefore, genes associated with azoospermia might also modulate the risk of TGCTs.

In the same GWAS study [23, 29], we uncovered that a single nucleotide polymorphism (SNP) in the human FOXN3 gene is associated with NOA. FOXN3 is evolutionary conserved. As indicated in Ensembl database, fly gene CHES-1-like is the ortholog of both human FOXN3 and FOXN2, which is a one to multiple orthologous relationship. To evaluate the functional relevance of the GWAS study, we assessed the function of CHES-1-like in fly spermatogenesis. CHES-1-like mutant male flies were viable and fertile. We found that CHES-1-like is not required for GSC maintenance or other spermatogenesis processes in fly testes. However, ectopic expression of CHES-1-like in germ cells significantly reduced male fertility. When CHES-1-like was overexpressed, spermatogonia failed to differentiate after four rounds of mitotic division, but continued to divide to form tumor-like structures. We found that CHES-1-like could activate dpp expression and block spermatocyte differentiation. Our results suggest that NOA-associated SNPs could be a potential modulator of testis tumor development.

RESULTS

Loss of CHES-1-like does not cause spermatogenesis defects

In our previous NOA GWAS screen [23, 29], one SNP (rs1887102, P=2.60 ×10^-7) in the human gene, FOXN3, was found to be associated with NOA (Figure 1A). FOXN3 is an evolutionarily conserved gene. In Drosophila, CHES-1-like is the ortholog of FOXN3. To evaluate whether this loci is functionally relevant to spermatogenesis, we tested the function of CHES-1-like in fly testes.

We knockdown CHES-1-like expression in germ cells of fly testes (Nos>CHES-1-like RNAi) and did not observe obvious defects (Supplementary Figure 1). We generated CHES-1-like deletion mutants using Cas9-mediated mutagenesis. We recovered multiple lines with different indels confirmed by PCR and sequencing (Figure 1B, 1C). Both the hemizygous mutant male flies and homozygous mutant female flies were viable and fertile (Figure 1D). We further examined the testes of CHES-1-like mutants by immunostaining with antibodies recognizing hub cells, germ cells, and cyst cells (Figure 1E, 1F). The patterns of all cell types were identical to the wild type controls, indicating that loss of CHES-1-like does not affect spermatogenesis in flies.

Ectopic expression of CHES-1-like in germ cells induces testis tumor formation

Since CHES-1-like loss of function did not result in spermatogenesis defects, we decided to examine whether ectopic expression of CHES-1-like could lead to testis malfunction. We generated UAS-CHES-1-like transgenic flies, and crossed these flies with different Gal4 lines expressing specifically in germ cells (Nos-Gal4 and Bam-Gal4; Figure 2A), cyst cells (Tj-Gal4; Supplementary Figure 2), and hub cells (Upd-Gal4; Supplementary Figure 3) in fly testes.

Fertility was significantly reduced in Nos>CHES-1-like (N=132) and Bam>CHES-1-like (N=91) male flies (Figure 2B). DNA staining in the testis tail revealed that
Figure 1: Loss of CHES-1-like does not cause spermatogenesis defects in Drosophila. A. The SNP rs1887102 at 14q32.11 was identified as a NOA associated SNP. Additive models of logistic regression analyses were used to estimate the P-value of association analyses in 981 NOA cases and 1,657 controls. The marker SNP is shown in purple and the $r^2$ value of the rest of the SNPs are indicated by different colors. The genes within the region of interest are annotated, with arrows indicating transcription direction. B. The design of knock out (KO) strategy for CHES-1-like gene using CRISPR/Cas9 technology. The scheme shows that the design of the injected construct, sequence of gRNAs, and the location of the gRNAs. C. Indels identified in Three CHES-1-like alleles. D. The fertility of controls and CHES-1-like mutants. E-F'. There is no structural defects of the CHES-1-like KO testes. FasIII labels hub cells (Blue), Zfh-1 labels cyst stem cells (Red), Vasa labels germ cells (Green). (E') and (F') are enlarged images of (E) and (F).
Figure 2: Ectopic expression of CHES-1-like in germ cells causes male fertility defect and tumor like structure formation. 

A. The patterns of Nos-Gal4 and Bam-Gal4 driving HA tagged CHES-1-like protein expression. 

B. The fertility rate of male flies with indicated genotypes. CHES-1-like overexpression in germ cells reduced male fertility. Chi-square test was used. ***: P<0.001; **: P<0.01. 

C. The quantification of the clustered elongated spermatids at the proximal ends of the testes with indicated genotypes. Nos>CHES-1-like testes have reduced numbers of elongated spermatids. Student’s t test was used. ***: P<0.001. 

D-F. Whole mount testes with indicated genotypes were stained with Hochest 33342. (D2-F2 and D3-F3) are enlarged views of (D1-F1) to show the proximal end of the testes and clustered spermatids. The boxed region of (E1) shows a cyst with over-proliferated germ cells. 

G. The quantification of testes containing cysts with over-proliferated germ cells (hyperplasia cysts). Chi-square test was used. ***: P<0.001; *: P<0.05. 

H1-J1. The phase contrast images of testes with indicated genotypes show an increasing of the small cells that resembled GSCs and spermatogonia in the apical region of both Nos>CHES-1-like and Bam>CHES-1-like testes (areas indicated by yellow double head arrows). The hyperplasia cysts were indicated by red arrows. (H2-J5) Immunostaining of testes with indicated genotypes. DNA, germ cells, hub cells, cyst cells, mature cyst cells, fusomes, and dividing cells were labeled with Hochest 33342, Vasa, FasIII, DE-Cadherin, Eya, 1B1, and pH3, respectively. Nos>CHES-1-like testes has overproliferated spermatogonial cells indicated by small branched fusomes and pH3 labeling. Scale bars: 20 μM.
clusters of elongated spermatids were greatly reduced in Nos>CHES-1-like testes compared to normal testes (Figure 2C-2F). An increase in the small cells resembling GSCs and spermatogonia in the apical region of both Nos>CHES-1-like and Bam>CHES-1-like testes was observed by phase contrast microscopy (Figure 2H1, 2I1, 2J1). Cysts with an overproliferation of small cells were also observed in the distal region of the Nos>CHES-1-like testis tips (Figure 2G, 2H1, 2I1, 2J1).

We further examined Nos>CHES-1-like and Bam>CHES-1-like testes with various cellular markers (Figure 2H-2J). DNA and Vasa staining confirmed that the region of GSCs and spermatogonial cells were expanded (Figure 2H2, 2I2, 2J2). In the Nos>CHES-1-like testes, cysts packed with small cells form tumor-like structures in which Vasa staining is lost (Figure 2I2). We used the 1B1 antibody to label fusome, a membrane structure that connects sibling germ cells. Most fusomes observed in the apical tip of Nos>CHES-1-like and Bam>CHES-1-like testes were small and branched (Figure 2H4, 2I4, 2J4), suggesting that the majority of cells were interconnected and resembled proliferating spermatogonia. This indicates that the spermatogonial cells failed to cease mitotic division after four rounds, and continually divided to form tumor-like structures. Indeed, using the cell proliferation marker phospho-Histone H3 (pH3) antibody to label dividing germ cells, pH3 could be observed only at the tip of wild type testes (Figure 2H5). However, ectopic pH3 labeling was observed in some cysts distal from the tip of the testes with ectopic CHES-1-like expression in germ cells (Figure 2I5, 2J5). We also used FastIII to label hub cells, and Eya and DE-cadherin to label cyst cells. There was no dramatic change in the number and morphology of either cell type (Figure 2H-2J and Supplemental Figure 1).

We further investigated Tj-Gal4-driven CHES-1-like expression in cyst cells and Upd-Gal4-driven CHES-1-like expression in hub cells, and examined the resulting testes with various markers. There was no obvious difference between wild type testes and the testes with ectopic CHES-1-like expression in germ cells (Figure 2I5, 2J5). In testis, Bam expression is repressed by BMP signaling. Therefore, CHES-1-like-mediated down regulation of Bam expression might reflect the activation of BMP signaling. Indeed, the BMP signaling downstream gene daughters against dpp (dad) was expressed at low levels in GSCs and spermatogonial cells in wild type testes (Figure 5A1-5A6). Expression levels of Dad-lacZ dramatically increased in the germ cells expressing CHES-1-like (Figure 5B1-5B6). However, ectopic expression of CHES-1-like in early cyst cells with Tj-Gal4 did not increase Dad-lacZ expression in cyst cells (Figure 5C1-5C6). The upregulation of Dad expression and downregulation of Bam expression indicates that CHES-1-like may ectopically activate BMP signaling.

**CHES-1-like inhibits spermatocyte differentiation through suppressing Bam expression**

The expression of Bam is required for spermatogonial cells to exit the mitotic cell cycle and begin differentiation [19]. It is likely due to the disruption of Bam signaling that spermatogonia were unable to cease mitosis in CHES-1-like germ cell ectopically expressing testes. To test this, we analyzed Bam expression patterns in these testes. In wild type animals, Bam is expressed primarily in the transient amplifying (TA) spermatogonia, a strip of cells near the apical tip of the testes (Figure 3A). The expression level of Bam was greatly reduced in both Nos>CHES-1-like (Figure 3B) and Bam>CHES-1-like testes (Figure 3C). However, the expression region of Bam was expanded (Figure 3F1-3J1, 3F2-3J2). We also used Bam-GFP reporter to analyze bam expression patterns. In Nos>CHES-1-like testes, GFP expression levels were reduced and the expression regions were expanded (Supplementary Figure 4). The reduction of GFP staining is obvious but less dramatic than the reduction of Bam staining in Nos>CHES-1-like testis (Figure 3B, Supplementary Figure 4), which is likely due to that not only the transcriptional levels but also the protein levels of Bam is affected by CHES-1-like expression.

It has been reported that around half of male germ cells lacking one copy of bam will complete one or more extra TA divisions before differentiation [30]. Removing one copy of bam further reduced the fertility of male flies with CHES-1-like overexpressing in germ cells (Figure 4A). DNA staining in the testis tail revealed that clusters of elongated spermatids were further reduced in these testes compared to the testes with CHES-1-like overexpressing alone (Figure 4B and 4C1-4E3). Bam expression in these testes was reduced to an undetectable level (Figure 3D, 3E). There was an increase in aberrant tumor-like cysts packed with over-proliferated small cells (Figure 4F-4I). These results suggest that CHES-1-like likely suppresses spermatocyte differentiation by down-regulating Bam expression.
Gal4-driven \( CHES-1\)-like expression in the wing imaginal disc porches and examined Dpp-lacZ expression patterns (Figure 5D-5G). In the wild type wing discs, Dpp-lacZ appeared as a thin strip at the anterior/posterior (A/P) boundary (Figure 5D1, 5D2). When CHES-1-like was overexpressed, the expression region of Dpp-lacZ was greatly expanded (Figure 5E1-5E3). As a result, the adult wings of the \( MS1096>CHES-1\)-like flies were deformed (Figure 5F1-5G2). However, another gene \( ptc\) does not change its expression patterns in the wing porches when CHES-1-like is overexpressed (Supplementary Figure 6), indicating that the changes of Dpp-LacZ expression patterns is not due to the morphological changes of the wing discs.

To test whether CHES-1-like directly binds to the \( dpp\) promoter to activate Dpp expression, we used a chromatin immunoprecipitation (ChIP) assay to detect the interaction between CHES-1-like and the \( dpp\) promoter. We expressed HA-tagged CHES-1-like in \( Drosophila\) S2 cells, and immunoprecipitated the protein-DNA complex after cross-linking. Indeed, CHES-1-like was able to interact with the \( dpp\) promoter region (Figure 5H-5J).

**DISCUSSION**

In this study, we used \( Drosophila\) testis as a model system to test the functional relevance of a NOA-associated human gene, \( FOXN3\). Although loss of \( CHES-1\)-like, the fly ortholog of \( FOXN3\), did not result in spermatogenesis defects, the ectopic expression of \( CHES-1\)-like in germ cells blocked spermatocyte differentiation and induced tumor-like structure formation. Our study has revealed CHES-1-like as a novel BMP signaling regulator to promote expression of the BMP ligand Dpp.

In flies, CHES-1-like, together with another fork head transcription factor Jumu, governs cardiac progenitor cell division and specification by regulating Polo kinase activity, as well as the expression of fibroblast growth factor and Wnt signaling pathway receptors [31, 32]. Inactivation of \( FOXN3\) in \( Xenopus\) and mice led to craniofacial defects and was sometimes lethal [33, 34].

It has been well established that BMP signaling is required for the differentiation and proliferation of osteoblasts of the mammalian skull [35]. Interestingly, the expression levels of the BMP pathway ligands BMP2,
Figure 4: Removing one copy of bam enhanced CHES-1-like overexpression phenotypes. A. Male fertility rates are further reduced in the Nos>CHES-1-like; Δ86/+ and Bam>CHES-1-like; Δ86/+ animals. Chi-square test was used. ***: P<0.001. B. The quantification of the clustered elongated spermatids at the proximal ends of the testes with indicated genotypes. Removing one copy of bam further reduced the numbers of elongated spermatids in CHES-1-like overexpression testes. Student’s t test was used, ***: P<0.001. C-E. Whole mount testes with indicated genotypes were stained with Hochest 33342. (C2-E2 and C3-E3) are enlarged views of (C1-E1) to show the proximal end of the testes and clustered spermatids. The boxed regions of (D1 and E1) show cysts with over-proliferated germ cells. F. The quantification of testes containing cysts with over-proliferated germ cells (hyperplasia cysts). Chi-square test was used. ***: P<0.001. G-I. The phase-contrast views and the immunostaining views of the apical tips of testes with indicated genotypes. Loss of one copy of Bam alone does not cause dramatic defects in testes. Removing one copy of bam in Nos>CHES-1-like and Bam>CHES-1-like tests greatly enhanced CHES-1-like overexpressing phenotypes. DNA, germ cells, hub cells, cyst cells, mature cyst cells, fusomes and dividing cells were labeled with Hochest 33342, Vasa, FasIII, DE-Cadherin, Eya, 1B1, and PH3, respectively.
Figure 5: CHES-1-like prevents spermatogonial differentiation through promoting Dpp expression to ectopically activate BMP signaling in germ cells. A1-C6. Over-expressing CHES-1-like in germ cells (Nos>CHES-1-like) but not cyst cells (Tj>CHES-1-like) activates BMP down-stream gene dad reporter dad-lacZ expression in germ cells. DNA, germ cells, cyst cells, dad-lacZ, and HA tagged CHES-1-like were labeled with Hочест 33342, Vasa, DE-Cadherin, LacZ, and HA antibodies. D1-E3. CHES-1-like overexpression in wing imaginal discs ectopically induces dpp expression in wing porch. Dpp expression was monitored by reporter dpp-lacZ. F1-G2. Images of adult wings of animals with indicated genotypes. H-J. ChIP assay indicates that CHES-1-like could bind to dpp promoter region. (H) Chromatin from control (CTRL) and HA-CHES-1-like transfected S2 cells was sonicated to obtain DNA fragments of appropriate size among 100-1000 bp. (I) Scheme shows the genomic region of dpp gene and the primers (P1-F, P1-R) used for CHIP assay. HA antibody could pull down dpp promoter region. (J) Quantification of (I).
BMP4, and BMP7 were greatly reduced in the FOXN3 knockout animals [34]. In our study, we found that CHES-1-like binds to the dpp promoter and activated Dpp expression, suggesting that CHES-1-like might directly regulate BMP ligand expression. Dpp is critical for fly development [36, 37]. However, loss of CHES-1-like did not result in any obvious developmental defect. It is possible that there is redundant molecules or pathways could compensate the effects caused by the loss of CHES-1-like. Jumu is a good candidate since it is playing redundant roles during fly heart development. However, ectopic expressing Jumu in germ cells (Nos>UAS-Jumu) does not have any defects (Supplementary Figure 1), indicating Jumu might not play similar roles as CHES-1-like in testes.

The function of FOXN3 has never been linked to testis tumors or male sterility. The data from human protein atlas (http://www.proteinatlas.org/) revealed that the protein levels of FOXN3 were upregulated in most cancer tissues. A recent study in a cancer cell line indicated that CHES1/FOXN3 decreases cell proliferation by repressing PIM2 and protein biosynthesis, suggesting that FOXN3 is a potential tumor suppressor [38]. However, we found that overexpression of CHES-1-like in fly testis could prevent early germ cell differentiation and lead to tumor-like structure formation. This paradox might arise because of the complex and context-dependent physiological functions of BMP signaling [39]. Therefore, whether FOXN3 promotes or inhibits tumor formation might be determined by the tissue context.

In this study, we found that the ortholog of a NOA-associated gene could modulate testis tumor formation. It is striking since infertility is a high risk factor of testis tumors. The misregulation of FOXN3 could contribute to both spermatogenesis defects and tumor genesis. Further investigation is necessary to dissect the role of FOXN3 in TGCTs.

**MATERIALS AND METHODS**

**Fly strains**

All flies were cultured on standard corn meal food at 25 °C. Information for alleles and transgenes used in this study can be found either in FlyBase or as noted: Nos-Gal4 (BDSC, #4937), Tj-Gal4 (DGRC, #104055), Upd-Gal4 (BDSC, #26796), MS1096-Gal4 (THFC, #TB00038), CHES-1-like RNAi (THFC, #THU2388), Dpp-lacZ, Dad-lacZ, BMPGFP/+, and Bam-Gal4;Δ86/+ are gifts from DH Chen.

**Generation of CHES-1-like transgenic fly**

CHES-1-like CDS (BDGP, clone LD24749) was subcloned into the pUAS-attB-HA vector. Primers used for PCR are: forward: 5- AAAAGCGGCCGCGCAATG GTTCTAGATTACCGCATCCAGGCGGACT-3 and reverse: 5-ATAAGAATGCGGCCGCGATGTTCGAACTAGAG TTCCACAGATAATCCCACACAG-3 and reverse: 5-CCGCTCGAGCTAGAGATTATTCGA-3. The transgenic flies were generated using standard procedures.

**CRISPR/Cas9-mediated genome editing**

CHES-1-like mutant was generated by an optimized CRISPR / Cas9-mediated genome editing method as described before [40]. Two sgRNAs targeting CHES-1-like exon regions were designed to generate an about 1.2kb DNA fragment deletion. Genomic DNA PCR and sequencing were used to confirm the deletions.

**Light and phase-contrast microscopy**

Fly testes were dissected in 1x phosphate-buffered saline (PBS) and washed several times. Testes were observed on slides by a phase-contrast microscope after gently squashing them with a cover slip. For an overall view of wing morphology, adult wings were observed directly under light microscope.

**Immunofluorescence and antibodies**

Fly testes were dissected in 1x PBS and fixed for 30 min in 4% paraformaldehyde. After washing three times in 1x PBS with 0.1% Triton X–100 (PBST) and blocking for 1hr in 5% bovine serum albumin (BSA), the samples were incubated with primary antibodies overnight at 4°C. After washing three times for 10 min in 0.1% PBST, the samples were incubated for 1 hr with secondary antibodies at room temperature followed by three times washing in 0.1% PBST. Testes were then stained with Hoechst 33342 (1.0 mg/ml, Invitrogen) for 5 min before mounting. The antibodies used were as follows: mouse anti-Eya (DSHB, 1:20); mouse anti-FasIII (DSHB, 1:50); rat anti-DE-cadherin (DSHB, 1:20); mouse anti-1B1 (DSHB, 1:75); rabbit anti-Bam C (a gift from DH Chen, 1:2000) [41]; rabbit anti-Vasa (1:1000, Santa Cruz); rabbit anti-PH3 (1.0 mg/ml, Invitrogen) for 5 min before mounting. The antibodies used were as follows: mouse anti-Vasa (1:1000, Santa Cruz); rabbit anti-PH3 (CST, 1:400); mouse anti-lacZ (Promega, 1:1000); rabbit anti-HA-tag (CST, 1:1000), rat anti-Zfi1 (C Tong lab, 1:2000), mouse anti-Ptc (DSHB, 1:100), chicken anti-GFP (Abcam, 1:1000). Secondary antibodies conjugated to A488, Cy3, A594, or A647 (Molecular Probes and Jackson Immunologicals) were diluted at 1:1000.

**Chromatin immunoprecipitation (ChIP) assay**

Formaldehyde cross-linking and chromatin immunoprecipitation (ChIP) assays of S2 cells were performed using a protocol as described before [42]. S2 cells transfected with HA-CHES-1-like were subjected for CHIP assay. Chromatin was sonicated on ice to obtain
DNA fragments of appropriate size among 100-1000 bp. Twenty percent of total supernatant was used as a total input control. Following removal of bound proteins, immunoprecipitated DNA was subjected to PCR.

The primer pair used for the CHES-1-like CHIP assays are: 5′-CACACACGCTCAGAGACACA-3′ and 5′-CAAGCGGGACTATAGGG-3′.

**Immunoprecipitation (IP)**

pUAS-Myc-Mad and pUAS-Flag-Medea were made by inserting PCR products of CDS of Mad and Medea into pUAS-Myc and pUAS-Flag constructs. Following primers were used: Mad-Myc F: 5′-AATTGCGGCCGCATGGACACCGACGATGTG-3′, Mad-Myc R: 5′-ACGTCTAGATTAGGCGGCGGCACGCGG-3′, Medea-Flag F: 5′-AATCTCGAGATGGGCGGCGGCTCGGG-3′, Medea-Flag R: 5′-ATTCTAGATTAGGCGGCGGCGCACCGCGG-3′. S2 cells transfected with Myc-Mad and HA-CHES-1-like or Flag-Medea were subjected to IP assays. IP were performed as described [43].

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**CONFLICTS OF INTEREST**

The authors declare no conflicts of interest.

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