Orb prevents autophagy in the Drosophila germline through translational repression of Atg12 mRNA

Isabelle Busseau1*, Stéphanie Pierson1, Dany Séverac2, Christelle Dantec2 and Martine Simonelig1*

1Institut de Génétique Humaine, CNRS, 141 rue de la Cardonille, 34396 Montpellier Cedex 5, France.
2Montpellier GenomiX, 141 rue de la Cardonille, 34396 Montpellier Cedex 5, France.

*Corresponding authors:
Isabelle Busseau, E-mail: Isabelle.Busseau@igh.cnrs.fr
Martine Simonelig, E-mail: Martine.Simonelig@igh.cnrs.fr
Tel: 33 4 34 35 99 36

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Abstract

*Drosophila* Orb, the homologue of vertebrate CPEB is a key translational regulator involved in oocyte polarity and maturation through poly(A) tail elongation of specific mRNAs. *orb* has also an essential function during early oogenesis which has not been addressed at the molecular level. Here, we show that *orb* prevents cell death during early stages of oogenesis, thus allowing oogenesis to progress. It does so through the repression of autophagy, by directly repressing, together with the CCR4 deadenylase, the translation of *Autophagy-specific gene 12* (*Atg12*) mRNA. The uncontrolled autophagy observed in *orb* mutant ovaries is reduced when *Atg12* mRNA levels are decreased. These results reveal a role of Orb in translational repression and identify autophagy as an essential pathway regulated by Orb during early oogenesis. Importantly, they also establish translational regulation as a major mode of control of autophagy, a key process in cell homeostasis in response to environmental cues.
Introduction

The regulation of developmental processes and cellular activities largely relies on translational control of mRNAs, and an important mechanism of this regulation involves changes in mRNA poly(A) tail lengths (Weill et al., 2012). Cytoplasmic polyadenylation element binding (CPEB) proteins act sequentially in poly(A) tail shortening and lengthening through the recruitment of deadenylases and poly(A) polymerases (Richter, 2007). They bind to UA-rich short sequences, referred to as cytoplasmic polyadenylation elements (CPEs), located within 3'UTRs of their target mRNAs (Pique et al., 2008; Richter, 2007). CPEB1 has been mostly studied for its implication in vertebrate oocyte maturation, where it is involved in translational repression through the recruitment of PARN deadenylase in immature oocytes, and in translational activation through an interaction with Gld2 poly(A) polymerase during maturation (Igea and Mendez, 2010; Kim and Richter, 2006). The role of CPEBs in translational regulation in somatic tissues has also been established in various contexts, including the control of cell proliferation, senescence, tumor development, synaptic plasticity and glucide metabolism (Alexandrov et al., 2012; Bava et al., 2013; Fernandez-Miranda and Mendez, 2012; Ortiz-Zapater et al., 2012; Udagawa et al., 2012).

Orb is the Drosophila homologue of vertebrate CPEB1 and, consistent with this, it is involved in cytoplasmic polyadenylation and translational activation during oogenesis, for oocyte polarity and maturation (Benoit et al., 2008; Castagnetti and Ephrussi, 2003; Chang et al., 1999; Juge et al., 2002). However, while Orb plays an established role in the initial steps of egg chamber formation, this function is poorly understood (Christerson and McKearin, 1994; Huynh and St Johnston, 2000; Lantz et al., 1994).

Here we show that cell death is the major defect in orb mutant early ovaries. Developmental programmed cell death occurs at three specific stages in the female germline: in the newly formed cysts (region 2 of the germarium, Figure 1A), during mid-oogenesis.
(stages 7-8), and during late oogenesis (stages 12-13) (McCall, 2004; Pritchett et al., 2009). Germ cell death during early and mid-oogenesis is strongly enhanced by starvation (Drummond-Barbosa and Spradling, 2001), and strikingly does not depend on the usual apoptotic activators such as hid, grim, reaper (Foley and Cooley, 1998; Peterson et al., 2007; Pritchett et al., 2009), but involves autophagy (Barth et al., 2011; Nezis et al., 2009). In *Drosophila*, autophagy contributes to developmental cell death in several developmental processes through caspase activation and DNA fragmentation (Denton et al., 2012). A molecular link has been established between autophagy and cell death during late oogenesis as nurse cell death by DNA fragmentation depends on autophagic degradation of the inhibitor of apoptosis (IAP/dBruce) (Nezis et al., 2010).

Here, we show that *orb* prevents autophagy and cell death during early stages of oogenesis, and identify *Autophagy-specific gene 12* (*Atg12*) mRNA as a direct target of translational repression by Orb.

**Results and Discussion**

**Early defects in orb mutant ovaries**

We sequenced *orb*\(^{F343}\) (Lantz et al., 1994) and *orb*\(^{36-53}\) (Morris et al., 2003), two strong or null *orb* alleles and found that both have premature stop codons (Figure S1A, B). For all subsequent analyses, we used *orb*\(^{F343}\) which had the most upstream stop codon. We used meiosis and cell cycle markers to further address the early germ cell defects reported previously (Huynh and St Johnston, 2000; Lantz et al., 1994). Oogenesis stops as pseudo-cysts just after the germarium in *orb*\(^{F343}\). In these pseudo-cysts, expression of C(3)G, a component of the synaptonemal complex, was generally maintained in several cells, reflecting a defect in meiosis restriction and, hence, oocyte determination (Figure 1C, D) (Huynh and St Johnston, 2000). Incorporation of bromodeoxyuridine (BrdU) was used to monitor germ cell
DNA replication. BrdU incorporation that is detected in region 1 of wild-type germarium was not affected in orb mutants (Figure 1E-F'). In contrast, the BrdU incorporation found in wild-type endoreplicating nurse cells was never detected in orb mutant pseudo-cysts (Figure 1E-F'), suggesting that the germ cells in these pseudo-cysts had not entered the nurse cell fate.

In the wild type, follicle cells encapsulate individual cysts to produce egg chambers. They express the transcription factor Traffic Jam (Tj) and Fasciclin 3 (Fas3) and, as they mature, Fas3 is down-regulated except in polar cells (Figure 1G). In orbF343 mutant ovaries, the follicle cells did not express Tj and failed to down-regulate Fas3, suggesting a defect in follicle cell maturation (Figure 1H). We used FLP-mediated FRT recombination to generate orb mutant cell clones and investigate a potential intrinsic function of orb in follicle cells. Ovaries with orb mutant follicle cells produced normal egg chambers (Figure 1I, I'), demonstrating that orb function was not required in the follicle cell lineage. Analysis of orb mutant germline clones showed that orb was dispensable in the germline stem cells for their self-renewal, division rate and differentiation (Figure S1C-F), consistent with orb function being downstream of these events. orb mutant germ cells were co-encapsulated with wild-type cysts in compound egg chambers (70%, n=237) (Figure 1J).

We conclude that orb is required in the germ cells for meiosis restriction and endoreplication of nurse cell nuclei. Defects in these processes prevent germ cell differentiation into oocyte and nurse cells and affect follicle cell maturation non-autonomously.

**Germ cell death is a major defect in orb mutant ovaries**

DAPI staining of orb mutant germline clones revealed pycnotic nuclei that can indicate cell death. We therefore used anti-cleaved caspase 3 and TUNEL assays to record potential cell death in orb mutant ovaries. The staining with both markers revealed cell death in orb mutant
germ cells (89% (n=140) and 69% (n=81) of pseudo-cysts marked with anti-cleaved caspase 3 and TUNEL, respectively) (Figure 2A-D). We used the UAS/Gal4 system to overexpress the known caspase inhibitor DIAP1 in the germ line (Mazzalupo and Cooley, 2006; Peterson et al., 2003). DIAP1 expression reduced the levels of cleaved caspase 3 and TUNEL staining in orb mutant germ cells (Figure 2E-F), and strikingly, rescued the formation of egg chambers in 23 to 27% of ovarioles, with follicle cells expressing Tj and Fas3 normally (Figure 2F', G). This showed that germ cell death contributed to the early oogenesis arrest in orb mutants. Oogenesis did not progress further in these rescued egg chambers, consistent with Orb regulating other processes in the ovary.

**Identification of mRNA targets of Orb**

To address whether orb could regulate cell death directly we performed Orb RNP immunoprecipitation-microarray (RIP-Chip) analysis (Keene, 2007) to identify mRNAs associated with Orb. We used either mature ovaries or early ovarian stages dissected from newly eclosed females (germarium to stage 8) (Figure S2). Significance analysis of microarrays (SAM) with a false discovery rate (FDR) of 0.01 then identified 421 and 603 mRNAs that were enriched at least 1.5 fold in Orb RIP from early and mature ovaries, respectively, compared to mock RIP (Figure 3A, Table S1). Gene ontology (GO) term enrichment analysis using DAVID with a p-value <0.05 (Benjamini corrected) identified the terms "translation", "cell cycle" and "mitochondria" as enriched among the mRNAs present in Orb RIP (Figure 3B). While CPEs have been defined in *Xenopus* (Pique et al., 2008), they remain uncharacterized in *Drosophila*. We used the software designed to identify CPEs in *Xenopus* (Pique et al., 2008) to identify those in mRNAs from Orb RIP. CPEs were not found enriched in Orb RIP mRNAs compared to mRNAs expressed in ovaries (16%, versus 22% in 6614 mRNAs expressed in ovaries from FlyBase). This could indicate that the software did
not reveal all Orb binding motifs. It appears unlikely because Orb possesses most aminoacids shown in CPEB1 to be involved in the interaction with the CPE (Afroz et al., 2014). It is important to note that Orb is part of large ribo-nucleoprotein complexes, where it is associated with many other RNA binding proteins (Weill et al., 2012). It is therefore very likely that a proportion of mRNAs in Orb RIP could coprecipitate through interactions with these other RNA-binding proteins.

Among the Orb mRNA early targets that might be involved in cell death, we identified Autophagy-specific gene 12 (Atg12) functionally annotated for "autophagic cell death". Since autophagy is thought to contribute to cell death in Drosophila oogenesis and Atg12 encodes an effector of autophagosome formation (Gorski et al., 2003; Scott et al., 2004), we focused on this mRNA and studied its potential direct regulation by Orb.

**Atg12 mRNA is a direct target of Orb**

Independent Orb RIP and quantification of Atg12 mRNA levels by RT-qPCR confirmed the presence of Atg12 mRNA in complex with Orb (Figure 3C). The presence of two potential CPEs in the Atg12 3'UTR (Figure S3) led us to use RNA pull-down assays to address a potential direct binding of Orb to Atg12 3'UTR. oskar (osk) 3'UTR known to interact with Orb (Chang et al., 1999) was used as a positive control, and the 3'UTR of thread (th) which encodes DIAP1 protein was used as a negative control. Both Atg12 and osk 3'UTRs were able to pull down Orb protein from an ovarian extract, while th 3'UTR was not to the same extent (Figure 3D). Competition assays were used to test the binding specificity of Orb to Atg12 3'UTR. osk 3'UTR and the TRI-Xef RNA were used as CPE-containing competitors, and TRI-XefΔCPE as a competitor which did not contain CPE. Unlabeled competitor RNAs were added in excess to the binding reactions (4X or 20X). The presence of 20X CPE-containing competitor RNAs substantially decreased the binding of Orb to Atg12 3'UTR, whereas the
non-CPE competitor did not (Figure 3D). These results are consistent with the direct binding of Orb to Atg12 3’UTR.

**Orb and CCR4 repress Atg12 mRNA translation through deadenylation**

We addressed whether Orb was involved in the control of Atg12 mRNA poly(A) tail lengths using PAT assays. In contrast to the shorter poly(A) tails that were reported in orb mutants for several mRNAs (Benoit et al., 2005; Castagnetti and Ephrussi, 2003; Juge et al., 2002), we found that Atg12 mRNA had elongated poly(A) tails in orb mutant early ovaries (Figure 3E), suggesting a role of Orb in Atg12 mRNA poly(A) tail shortening that could lead to its translational repression. The role of Orb as a translational repressor of Atg12 mRNA was confirmed by the increased levels of Atg12 protein in orb mutant pseudo-egg chambers compared to wild-type (Figure 4A, B).

While a function of Orb in translational repression has not been reported, CPEB has been shown to act as a translational repressor prior to its action in cytoplasmic polyadenylation, and the presence of a deadenylase in the CPEB complex appears to contribute to this repressor function (Hosoda et al., 2011; Kim and Richter, 2006). We therefore analyzed the potential role of the CCR4-NOT deadenylase complex in Orb-dependent poly(A) shortening of Atg12 mRNA. PAT assays of Atg12 mRNA in ovaries mutant for twin, the gene encoding CCR4 deadenylase (Temme et al., 2004) revealed longer than wildtype poly(A) tails, indicating a role of CCR4 in shortening Atg12 mRNA poly(A) tails (Figure 3E). Co-immunoprecipitation experiments used to determine if Orb and CCR4 could be in complex in early ovarian stages showed that Orb was able to co-precipitate CCR4 in the presence or the absence of RNAs (Figure 3F), indicating that both proteins are part of the same complex. Importantly, we also uncovered a strong genetic interaction between orb and twin. Whereas single heterozygous orb or twin mutant females displayed normal
oogenesis, double *twin orb* heterozygous mutant females showed about 50% of ovarioles arrested at mid-oogenesis (Figure 3G, H). Arrested egg chambers expressed Atg12 protein, consistent with Orb and CCR4 acting together to repress *Atg12* mRNA translation (Figure 3I). Moreover these egg chambers underwent cell death as indicated by the expression of cleaved caspase 3 and staining by TUNEL assays (Figure 3J, K).

These data strongly suggest that Orb acts with CCR4 to repress *Atg12* mRNA translation by poly(A) tail shortening.

**Orb represses autophagy and cell death during oogenesis**

*Atg12* protein expression in *orb* mutant ovaries correlated with the induction of autophagy indicated by the presence of autophagosomes visualized using the Lysotracker marker (100% (n>100) of *orb* F343 ovarioles) and punctate staining of Atg8 protein (Barth et al., 2011) (Figure 4C-F). To address the functional importance of *Atg12* mRNA regulation by Orb in autophagy and cell death, we reduced *Atg12* expression in *orb* mutant ovaries using RNAi. Germline expression of *Atg12*-RNAi in *orb* F343 mutant ovaries reduced the amounts of *Atg12* protein, thus validating the RNAi transgene (Figure 4G). This led to decreased autophagy visualized with Lysotracker and Atg8-labeled autophagosomes, showing that *Atg12* expression had an important role in autophagy induction (Figure 4H, I). Strikingly, reduced expression of *Atg12* resulted in a strong rescue of egg chamber formation in *orb* F343 mutant ovaries, with up to 62% of ovarioles able to produce egg chambers surrounded by follicle cells expressing Tj and Fas3 normally (Figures 2G, 4J-L). Staining with anti-cleaved caspase 3 and TUNEL showed that cell death was reduced in these egg chambers (62% (n=47) and 72% (n=18) of rescued egg chambers with no or weak anti-cleaved caspase 3 and TUNEL staining, respectively) (Figure 4J, K).
These results demonstrate that Atg12 expression plays an important role in the induction of autophagy in oogenesis and that Orb acts to directly repress Atg12 mRNA translation thereby preventing autophagy and, to some extent, cell death.

Of the twelve autophagy-specific genes found in the Drosophila genome, seven (including Atg12) contained CPEs (UUUUAAU, UUUUAU, UUUUACU or UUUUAAGU) in their 3’UTR (Figure 4M, Figure S3). We used Orb RIP and RNA quantification by RT-qPCR to analyze the presence of these seven mRNAs in Orb RNP complexes. Six out of the seven were found enriched in Orb RIP suggesting that Orb might globally co-regulate the autophagic pathway (Figures 3C, 4N).

Conclusion
The role played by autophagy and Atg genes in a wide range of biological processes is now emerging (Boya et al., 2013). Post-translational modulation of Atg proteins is recognized as an important mode of regulation of autophagy and crosstalk with other cellular processes, in response to cellular and environmental cues. Here, we add translational control as another key mechanism of regulation of autophagy. We have demonstrated the direct regulation of Atg12 mRNA by Orb: Orb represses Atg12 mRNA translation through its deadenylation by CCR4, thus preventing autophagy.

Autophagy is highly regulated by the levels of nutrients. During Drosophila oogenesis, autophagy is activated upon starvation or inhibition of the insulin/TOR signaling pathway, through the upregulation of Atg protein levels (Barth et al., 2011). On the other hand, Orb is part of the highly dynamic RNP granules defined as P (processing) bodies that are essential for translational regulations in germ cells (Weil et al., 2012). These P bodies (including Orb) undergo massive reorganization upon reduction of nutrient availability (Snee and Macdonald, 2009). Therefore, an intriguing implication from our data is that Orb,
potentially with other translational regulators within P bodies, would act as a sensor of environmental cues to regulate autophagy. Consistent with this, we found that decreasing orb gene dosage increased cell death induced by amino-acid starvation (Figure S4). In agreement with the implication of CPEB proteins in interpreting environmental conditions, such a function has been proposed for CPEB1 in the regulation of glucose homeostasis in mouse liver (Alexandrov et al., 2012).

Translational regulation of autophagy might have a dramatic impact in many contexts, including in the expanding group of degenerative diseases involving RNA granules. Pathological RNA granules that form in neurodegenerative disorders have been proposed to be targeted by autophagy (Buchan et al., 2013; Ramaswami et al., 2013) which is conversely altered in some neurodegenerative diseases (Rubinsztein et al., 2012). In these disorders, translational deregulation of autophagy through affected RNA granules may induce a positive-feedback loop leading to enhanced production of pathological RNA aggregates.

**Experimental Procedures**

*Drosophila stocks and genetics*

Fly stocks used in this study and clonal analysis are described in Supplemental Experimental Procedures.

**Fluorescent labeling and immunostaining**

Primary antibodies for immunostaining and procedures for fluorescent labeling are described in Supplemental Experimental Procedures.

**Immunoprecipitations and RNA analyses**
Immunoprecipitations were performed as described (Zaessinger et al., 2006) and were followed by either RNA extraction and RT-PCR or by western blots as detailed in Supplemental Experimental Procedures. Poly(A) tail length analysis by PCR (PAT assay) and RT-qPCR using the LightCycler System (Roche Molecular Biochemical) were performed as described previously (Benoit et al., 2005; Benoit et al., 2008; Zaessinger et al., 2006) using primers listed in Supplemental Informations.

RNA pull-down assays

UTP-biotinylated RNAs and unlabeled competitor RNAs were synthesized using T7 RNA polymerase on PCR fragments synthesized from genomic DNA with primers indicated in Supplemental Informations. RNA pull-down experiments were performed as published previously (Besse et al., 2009). For each experimental point, 20 µL of ovarian extract from 20 one day-old females were used.

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References


Figure legends

Figure 1: Cell cycle and meiosis defects in orb mutant germ cells

(A) Schematic drawing of a wild-type germarium. Germ cells are represented in white and the
differentiated oocyte in red (GSC, germline stem cells). Somatic follicle cells are in grey.

(B) Schematic drawing of the mitotic divisions producing a cystoblast (CB) from one GSC,
and a 16 germline cell-cyst from one CB. Red dots indicate meiosis which initiates in four of
the 16 germ cells within a cyst, and becomes restricted to the oocyte.

(C-H) Defects in meiosis restriction and cell cycle in orbF343 germ cells. Wild type (left
panels) and orbF343 (right panels) ovarioles stained with anti-Vasa (germ cell marker, red) and
anti-C(3)G (green) (C, D); anti-Vasa (red) (E, F) or DAPI (E’, F’) and BrdU incorporation
(green) (E-F’); anti-Tj (red) and anti-Fas3 (white) (G, H). White arrows indicate the lack of
C(3)G restriction to one cell in (D) and the lack of BrdU incorporation in pseudo-cyst nuclei
in (F, F’).

(I, I’) orbF343 follicle clone labeled with DAPI (blue) and GFP (green) (I), and with anti-Tj
(red) and anti-Fas3 (white) (I’).

(J) orbF343 germline clones labeled with DAPI (blue) and GFP (green). White arrowheads
indicate orbF343 germ cells. Scale bars: 20 µm.

Figure 2: Cell death in orb mutant ovaries

(A, B) Wild-type ovarioles labeled with DAPI (blue), TUNEL (green), anti-cleaved caspase 3
(Casp, red) and anti-Fas3 (white). The white arrowhead indicates TUNEL staining in follicle
cells.

(C-F’) orbF343 (C-D’) and UASp-DIAP1/+; nos-Gal4 orbF343/ orbF343 (nos>DIAP1 orbF343) (E-
F’) ovarioles labeled with DAPI (blue) and TUNEL (green) (C, E); DAPI (blue), anti-cleaved
caspase 3 (Casp, red) (D, F); and anti-Tj (green), anti-cleaved caspase 3 (Casp, red) and anti-Fas3 (white) (D’, F’). Scale bars: 20 µm.

(G) Quantification of ovarioles containing egg chambers in orbF343 either expressing DIAP1 (UASp-DIAP1/+; nos-Gal4 orbF343/orbF343 (nos>DIAP1)) or Atg12-RNAi (UASp-TRIP-Atg12 orbF343/nos-Gal4 orbF343 (nos>Atg12-RNAi)).

Figure 3: Atg12 mRNA is a direct target of Orb

(A) Venn diagram of mRNAs significantly enriched in Orb versus mock RIP, from early (blue outline) and mature (green outline) ovaries.

(B) Gene ontology term enrichment using DAVID.

(C) RT-qPCR experiments showing enrichment of Atg12 mRNA in Orb RIP versus Mock RIP. Mean ratios of three quantifications of Atg12 mRNA levels in Orb/Mock, normalized with RpL32 mRNA. Error bars: SEM. Two independent experiments (Exp 1, Exp 2) are shown.

(D) Biotin RNA pull-down experiments showing that Atg12 and osk 3’UTRs pull down Orb more efficiently than th 3’UTR (top panels). Biotin RNA pull-down competition assays showing efficient competition with 20X of unlabeled osk 3’UTR or pTRI-Xef which contain CPEs, and the lack of efficient competition with 20X of unlabeled RNA fragment devoid of CPE (pTRI-XefΔCPE) (bottom panel).

(E) PAT assays showing elongated poly(A) tails of Atg12 mRNA in orbF343 and twinDG23102 early ovaries. RpL32 mRNA was used as a negative control. The profiles of Atg12 PAT assays using ImageJ are shown.

(F) Orb immunoprecipitation in early ovaries showing CCR4 co-precipitation, both in the absence (-RNase) and in the presence (+RNase) of RNase A and micrococcal nuclease.

(G) DAPI staining of a orbF343 +/+ twin8115 ovariole showing mid-oogenesis arrest.
(H) Quantification of ovarioles with mid-oogenesis arrest in control and orb\textsuperscript{F343} +/- twin\textsuperscript{8115} females.

(I-K) orb\textsuperscript{F343} +/- twin\textsuperscript{8115} ovarioles labeled with DAPI (blue) and anti-Atg12 (green) (I); DAPI (blue) and anti-cleaved caspase 3 (Casp, red) (J); DAPI (blue) and TUNEL (green) (K). Scale bars: 20 \(\mu\)m.

**Figure 4: orb regulates autophagy**

(A-L) Wild-type (A, C, E), orb\textsuperscript{F343} (B, D, F) and UASp-TRIP-Atg12 orb\textsuperscript{F343}/nos-Gal4 orb\textsuperscript{F343} (nos>Atg12-RNAi orb\textsuperscript{F343}) (G-L) ovarioles labeled with DAPI (blue) and anti-Atg12 (A, B, G); DAPI (blue) and Lysotracker (LTR, red) (C, D, H); DAPI (blue) and anti-Atg8 (E, F, I); DAPI (blue) and anti-cleaved caspase 3 (Casp, red) (J); DAPI (blue) and TUNEL (green) (K); anti-Tj (green) and anti-Fas3 (white) (L). White arrows indicate rescued egg chambers. Scale bars: 20 \(\mu\)m.

(M) Presence/absence of CPEs in 3'UTRs of autophagy-specific genes.

(N) RT-qPCR experiments showing enrichment of Atg mRNAs in Orb RIP. Mean ratios of three quantifications of Arg mRNA levels in Orb/Mock, normalized with RpL32 mRNA. Error bars: SEM.
Figure 1

- **A**: Diagram of a wild type germ line cyst showing the organization of germ cells and follicle cells.
- **B**: Diagram illustrating the process of cyst encapsulation and cell division.
- **C** and **D**: Immunofluorescent images showing Vasa expression in wild type and orb$^{343}$ animals.
- **E** and **F**: Images showing BrdU incorporation to mark cell division in wild type and orb$^{343}$.
- **G** and **H**: DAPI staining of germ line clones in wild type and orb$^{343}$.
- **I** and **J**: GFP expression in germ line clones under DAPI staining.
- **L**: Further images showing Tj Fas3 expression in germ line clones.

Legend:
- C(3)G: Germ line clone
- orb$^{343}$: Mutant genotype
- DAPI: DNA dye
- GFP: Green fluorescent protein
- Vasa: Germ cell marker
- BrdU: Cell division marker
- Tj: Cell type marker
wild type

orb\textsuperscript{F343}  
nos>DIAP1 orb\textsuperscript{F343}

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Ovarioles with egg chambers in orb\textsuperscript{F343} background.
Figure 3

A

421 genes (early)

603 genes (mature)

95

326

277

B

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C

Atg12 mRNA in Orb IP / Mock IP

D

Extract th 3'UTR Atg12 3'UTR osk 3'UTR

Orb

Biotin-Atg12 3'UTR + competitor RNA:

pTRI-Xef

osk 3'UTR

ΔCPE

pTRI-Xef

- 4x 20x

4x 20x

Orb

E

F

Atg12

62A

12A

Rpl32

wild type

Orb+F343 + /

Mock IP

Exp 1 Exp 2

wild type

Orb+F343 - RNase + RNase

Input - RNase + RNase

Orb

CCR4-

F343 + / + twin8115

G

DAPI CaspDAPI Atg12 DAPI TUNEL

H

Ovarioles with mid-oogenesis arrest

w1118 5% (n=193)

orbF343 / + 5% (n=182)

twinDG23102 / + 54% (n=502)

I

DAPI Atg12

J

DAPI Casp

K

DAPI TUNEL

orbF343 / twin8115
Table 1: Gene CPEs

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Figure 4