Mitochondria regulate intestinal stem cell proliferation and epithelial homeostasis through FOXO

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ABSTRACT A metabolic transition from glycolysis to oxidative phosphorylation is often associated with differentiation of many types of stem cells. However, the link between mitochondrial respiration and stem cells’ behavior is not fully understood. We genetically disrupted electron transport chain (ETC) complexes in the intestinal stem cells (ISCs) of Drosophila. We found that ISCs carrying impaired ETC proliferated much more slowly than normal and produced very few enteroblasts, which failed to further differentiate into enterocytes. One of the main impediments to ISC proliferation and lineage specification appeared to be abnormally elevated forkhead box O (FOXO) signaling in the ETC-deficient ISCs, as genetically suppressing the signaling pathway partially restored the number of enterocytes. Contrary to common belief, reactive oxygen species (ROS) accumulation did not appear to mediate the ETC mutant phenotype. Our results demonstrate that mitochondrial respiration is essential for Drosophila ISC proliferation and lineage specification in vivo and acts at least partially by repressing endogenous FOXO signaling.

INTRODUCTION
Eukaryotes utilize either glycolysis or oxidative phosphorylation (OXPHOS) to extract energy from food and store it as the high-energy ATP molecule. Unlike glycolysis, which partially metabolizes glucose to pyruvate, OXPHOS completely oxidizes glucose to CO₂ and is much more efficient in harnessing energy (Wallace, 2012). OXPHOS is carried out by the electron transport chain (ETC) complexes and mitochondrial ATP synthase located on the mitochondrial inner membrane. Mitochondria are thought to evolve from endosymbiotic bacteria. Over the course of evolution, most mitochondrial genes have migrated to the nucleus, whereas core ETC subunits still remain on the mitochondrial genome (mtDNA) (Wallace, 2012). To accommodate different metabolic profiles and energy demands, cells in different tissues and developmental stages modulate the amount of mtDNA they contain and its expression level (Moraes, 2001). The dynamic regulation of mitochondrial activity is particularly evident during stem cell differentiation. Many types of stem cells favor glycolysis even in the presence of oxygen, a phenomenon termed as “aerobic glycolysis” (Vander Heiden et al., 2009; Rafalski et al., 2012; Lisowski et al., 2018), but undergo a metabolic transition during differentiation (Folmes et al., 2012; Zheng, Boyer, et al., 2016), increasing mitochondrial mass (Cho et al., 2006; Prigione et al., 2010; Zhang, Marsboom, et al., 2013), mtDNA copy number (Cho et al., 2006; Facucho-Oliveira et al., 2007; Wanet et al., 2014), and respiratory activity (von Heimburg et al., 2005; Zhang, Marsboom, et al., 2013; Wanet et al., 2014). Pharmacological or genetic disruptions of ETC complexes often block stem cell differentiation (Chung et al., 2007; Mandal et al., 2011) or interfere with lineage specification (Inoue et al., 2010; Cherry, Gagne. et al., 2013; Diaz-Castro, Pardal, et al., 2015), indicating an involvement of mitochondrial respiration in regulating stem cells’ activity. However, it is largely unknown how ETC complexes interact with the cellular signaling pathways to influence stem cell behavior.

The Drosophila adult midgut is a well-established model to study stem cell behavior. Intestinal stem cells (ISCs) have been found...
interspersed among the basal epithelium of the entire adult fly midgut (Buchon, Osman et al., 2013; Marianes and Spradling, 2013). ISCs proliferate and give rise to new epithelium in response to tissue injury or stresses (Biteau et al., 2011; Jiang and Edgar, 2011; Meng and Biteau, 2015). The ISC usually undergoes an asymmetric division that generates a new ISC and an intermediate progenitor, enteroblast (EB), which can further differentiate into either an absorptive enterocyte (EC) characterized by a polyplid nucleus, or a secretory enterodocrine (EE) cell at a lower frequency (Ohlstein and Spradling, 2007; Biteau et al., 2008; Jiang and Edgar, 2011; Biteau and Jasper, 2014). The simple organization, defined ISC lineage, and ample set of genetic tools available in Drosophila make the Drosophila midgut an ideal model to explore the impact of ETC complexes on stem cells’ activity.

Applying a selection scheme based on a mitochondrially targeted Xhol endonuclease (MitoXhol) (Xu et al., 2008), we isolated a temperature-sensitive lethal mtDNA mutation (ts), mt:CoI/T300T, that abolishes the sole Xhol site in the cytochrome c oxidase (COX) subunit I locus on the Drosophila mitochondrial genome (Hill, Chen, and Xu, 2014; Chen et al., 2015). The resulting threonine-to-isoleucine substitution destabilizes the association between the mutated Col protein with the cytochrome A heme, which leads to severely reduced COX activity at the restrictive temperature of 29°C (Chen et al., 2015). In tissues or cells harboring a homoplasmic ts allele, mitochondrial inner membrane potential is reduced, but there is no accumulation of reactive oxygen species (ROS) (Chen et al., 2015). Homoplasmic ts flies develop normally at 18°C, but fail to eclose at 29°C. In heteroplasmic flies carrying both the ts mtDNA and wild-type (wt) mtDNA, tissue-specific expression of MitoXhol can effectively remove the wild-type mtDNA and results in ts homoplasmy in those tissues (Chen et al., 2015). In this study, we expressed MitoXhol in the midgut using the esgFlp-Out (esgF/O) method (Jiang et al., 2009). This genetic scheme generated ts homoplasmy in the ISCs and their derivatives in heteroplasmic flies and allowed us to investigate how ETC deficiency impacts ISC’s activity in an overall healthy background. We report that deficiencies in cytochrome chain complexes in midgut stem cells impaired their proliferation and blocked the EC specification, which is partially due to an increase of the activity of the FOXO transcription factor.

RESULTS

COX deficiency in Drosophila ISCs decreases the number of their differentiated progeny

To study how defective respiration might affect the behavior of Drosophila ISCs, we generated clones of ISCs harboring the temperature-sensitive lethal mtDNA mutation (ts): mt:CoIT300T (Supplemental Figure S1). In heteroplasmic ts flies, we drove the expression of a mitochondrially targeted Xhol restriction enzyme (UAS-MitoXhol) specifically in ISCs, using the esgF/O system (esg-GAL4, tub-GAL80ts, UAS-nlsGFP/CyO; UAS-flp, act>CD2-GAL4/TM6) (Jiang et al., 2009), to render the ISCs homoplasmic for the ts mutation (Chen et al., 2015). In this system, GAL4 is specifically expressed in ISCs under the esg promoter, but its activity is hampered by ubiquitous GAL80ts at 18°C. Newly emerged adult flies are subjected to a pulse of heat shock at 37°C and maintained at 29°C afterward. This treatment inactivates the GAL80ts protein and allows esg-GAL4 to activate Flp expression in ISCs (Jiang et al., 2009). Flp in turn induces act>Gal4, which activates UAS transgenes in all progeny derived from ISCs (Jiang et al., 2009). This pulse of heat shock also stresses the midgut to induce ISC proliferation (Strand and Michelli, 2011). Expression of MitoXhol removes the wild-type mtDNA in heteroplasmic cells, rendering them homoplasmic for ts and deficient in COX activity (Chen et al., 2015). Meanwhile, defective cells are labeled with GFP and therefore readily distinguished from healthy cells. As a control, we introduced the same combination of nuclear transgenes in a homoplasmic background harboring an synonymous mtDNA mutation that abolishes the Xhol site on mt:Col locus, but does not change the COX amino acid sequence (mt:Col/w+) (Chen et al., 2015). From here on out, we refer to the ISCs and their progeny carrying the ts allele as mutant and the ISCs and progeny carrying mt:Col/w+ as control.

In control flies, one third of the midgut was occupied by GFP+ cells 3 d after heat shock. The GFP+ area increased in older flies, reaching 40% and 60% at day 10 and day 20, respectively (Figure 1, A and B). In the mutants, the midguts were fragile during dissection and appeared thinner in controls. Less than 20% of the tissue was GFP+ at day 3 (Figure 1B), and the proportion of GFP+ area remained the same in older mutant flies (Figure 1B). We stained midguts for Delta, Su(H)G126-LacZ, Pdm1, and Prospero to mark ISCs, EB, EC, and EE cells, respectively (Supplemental Figure S1, A and C). We also considered the location, shape, and nuclear size of cells to determine cell types. The abundance of ISCs was higher in the mutant than in the control (Supplemental Figure S2A). However, the abundance of GFP+ differentiated cells (EBs, ECs, and EEs) was much lower in mutant midguts compared with control (Figure 1, C and D; Supplemental Figure S2A).

COX deficiency impairs ISC proliferation and EB-to-EC specification

Multipotent ISCs proliferate and generate EB cells, which subsequently differentiate into ECs or EE cells. Thus, the paucity of ISC progeny (EB, EC, and EE cells) in mutant midguts could be caused by either impaired ISC proliferation and differentiation or increased cell death. To test these possibilities, we first assessed apoptosis by staining midguts with an antibody against cleaved-caspase 3, to mark apoptotic cells (King et al., 2015). Overall, very few cells stained positive for cleaved-caspase 3 in either control or mutant midguts (Figure 2, A and B). They mostly located to the anterior part of midguts and had large nuclei, indicating that they were presumably ECs. Of primary significance, the abundance of apoptotic cells was comparable in control and mutant guts (Figure 2, A and B), demonstrating that apoptosis is not increased in mutant midguts, nor should it be considered the cause of the lack of ISC progeny. Additionally, the numbers of GFP+ ISCs were comparable between control and mutant anterior midguts either 8 or 3 d after induction (Supplemental Figure S2B), further substantiating that the lack of ISC progeny in the mutant midgut is not caused by an increase of ISC death.

To address whether COX deficiency impairs cell proliferation, we examined DNA synthesis via EdU incorporation. Many EdU+ cells were observed in control midguts (Figure 2C and Supplemental Figure S2C), and the majority of these cells had large nuclei (Figure 2C), indicating that they are EB or newborn EC undergoing endoreplication (Xiang et al., 2017). A small fraction of EdU+ cells had small nuclei, likely the proliferating ISCs. In mutant midguts, the overall numbers of EdU+ cells were much lower than that in controls (Figure 2C and Supplemental Figure S2C). Very few EdU+ cells with small nuclei were detected in mutant midguts (Figure 2, C and D), suggesting that COX deficiency impairs ISCs’ division. We barely observed EdU+ cells with large nuclei (Figure 2C and Supplemental Figure S2C), indicating a lack of endoreplication of EB/EC in the mutant midgut.

The lack of endoreplication could be due to either a lack of EB or impaired EB-to-EC specification. To further test whether ETC disruption impairs EC lineage specification, we replaced the esg-GAL4 used in the esgF/O system with an EB-specific driver.
Su(H)GBE-GAL4, to initiate the expression of Flp, GFP, and MitoX-hol in EBs (Figure 3A). We also used the mtDNA synonymous mutation, mt:Co1^syn, as control. Similar to results in the esg^ts F/O system, the numbers of GFP+ EC were significantly lower in mutant compared with control midguts (Figure 3, B and C), demonstrating that COX deficiency indeed impairs EB-to-EC specification.

ETC deficiencies block ISC proliferation

We initially thought that the ts mtDNA would undergo compensatory replication to restore the mtDNA content to normal after the elimination of the wild-type mtDNA in heteroplasmic ISCs (Supplemental Figure S1B). However, mtDNA content in mutant ISCs remained lower than that in control cells (Supplemental Figure S3A), suggesting an incomplete compensatory replication of mtDNA. Thus, aside from defective Complex IV caused by the ts mutation, mutant ISCs might also possess inadequate complexes I, III, or V because of reduced mtDNA content. To determine whether the lack of ISC proliferation is specifically pertained to defective Complex IV, we disrupted every single complex independently by knocking down the nuclear encoded components (Figure 4A). We used RNA interference (RNAi) lines that were proven effective in a previous study (Teixeira, Sanchez, Hurd, et al., 2015) and activated them in ISCs and progeny specifically using the esg^ts F/O system. While disruption of each individual ETC complex impaired ISC proliferation to some extent (Figure 4B; Supplemental Figure S3, B–D), RNAi against genes in complexes III and IV caused much stronger defects than other complexes (Figure 4B; Supplemental Figure S3, B–D).

Active ETC complexes establish mitochondrial inner membrane potential (ΔΨ_m) that drives ATP synthesis. To understand the biochemical deficiency in mutant ISCs, we assessed the ΔΨ_m, using ratio-metric imaging between TMRM and MitoTracker-deep red, two fluorescent dyes reporting ΔΨ_m and mitochondrial mass, respectively. Consistent with previous studies (Chen et al., 2015; Zhang et al., 2019), ΔΨ_m, indicated as the ratio of TMRM to MitoTracker-deep red, was markedly reduced in mutant ISCs compared with the control (Supplemental Figure S4, A and B). Depolarized mitochondria are less
These observations spurred us to test whether the ROS-FOXO pathway was involved in inhibiting ISC proliferation in mutant midguts.

We first assessed ROS levels by staining midguts with CellROX (Sekihara et al., 2016; Manent et al., 2017), a fluorescent indicator for ROS. CellROX intensity markedly increased in guts pretreated with an organic peroxide, tert-butyl hydroperoxide, but was reduced in guts pretreated with a reducing agent, N-acetylcysteine (Supplemental Figure S6, A and B), suggesting that CellROX indeed properly indicates the level of ROS. We found that intensities of CellROX were comparable between control and mutant midguts (Supplemental Figure S6, A and B). Additionally, co-overexpression of the ROS-scavenging enzymes SOD2 and catalase (Sun et al., 2002) did not rescue the ISC proliferation in the ts mutant background (Supplemental Figure S6, C and D). These results indicate that ROS is not elevated in mutant cells, nor involved in the inhibition of ISC proliferation. By contrast, FOXO protein was markedly increased in ISCs and EBs in the mutant midgut compared with control (Figure 5A) and when Complex III was disrupted (Figure 4A). To assess FOXO activity, we used a lacZ reporter under the control of the promoter of the 4E-binding protein (4EBP) gene, a target of FOXO (Puig et al., 2003; Demontis and Perrimon, 2010). We found that the amount of lacZ protein was markedly higher in mutant cells than in controls (Supplemental Figure S7). Taken together, these results demonstrate that FOXO protein and its transcriptional activity are elevated in mutant cells, in the absence of any detectable increase in cellular ROS.

To understand the consequence of FOXO accumulation, we overexpressed FOXO using the esg^F/O-mitoXhoI flies system and found that it significantly reduced the GFP^+ area (Figure 5B and Supplemental Figure S8), indicating that increased FOXO activity can indeed block ISC lineage specification. Reciprocally, knocking down foxo in ISCs significantly increased the GFP^+ area in mutant midguts (Figure 5, B and C; Supplemental Figure S8). Further classification of cell types demonstrated that foxo knockdown in ISCs significantly restored production of GFP^+ EB and GFP^+ EC but not GFP^+ EE cells (Figure 5, C and D). We also carried out RNAi against foxo in EBs using the Sul(H)GBE^F/O system and found that it restored the number of ECs in the mutant midgut (Supplemental Figure S9). Altogether, these results indicate that FOXO protein is elevated in mutant midguts, blocking the ISC commitment to EB and EB-to-EC specification.

**Thor mediates the impairment in EC specification in mutant midguts**

To understand how FOXO interferes with ISC’s activity, we performed RNA sequencing (RNAseq) on control and mutant ISCs. About 10% of the mRNAs we detected were either up-regulated or down-regulated in the mutant ISCs relative to control (Supplemental

**ETC deficiency blocks the ISC-to-EC specification through FOXO**

Electron flow from Complex III to Complex IV is a major step generating ROS (Starkov, 2008; Bleier and Drose, 2013). A previous study demonstrated that ROS can block the cell cycle progression via a FOXO-dependent inhibition on cyclin E/CDK2 (Owusu-Ansah et al., 2008). FOXO is also believed to play important roles in regulating stem cell quiescence and survival (Cheung and Rando, 2013). Observations spurred us to test whether the ROS-FOXO pathway was involved in inhibiting ISC proliferation in mutant midguts.
The down-regulated pool included many genes involved in cell differentiation, proliferation, and growth regulation (Supplemental Table S2). This result suggests a possible metabolic adaptation in response to theOXPHOS deficiency in mutant midguts. Importantly, knockdown of foxo in mutant ISCs restored the expression level of more than 50% of the up- or down-regulated genes (Figure 6A and Supplemental Table S2), indicating that the transcriptional response to COX deficiency is partially mediated by FOXO.

Thor, the sole homologue of 4EBP in the Drosophila genome, was among the genes whose expression was up-regulated in mutant midguts and restored by foxo knockdown (Supplemental Table S1). This finding is consistent with previous studies showing that Thor is a downstream target of FOXO in the regulation of cell growth and metabolism (Miron et al., 2001; Teleman et al., 2005). 4EBP proteins are highly conserved in metazoans, where they block the initiation of cap-dependent mRNA translation by binding to elongation factor eIF4E (Hernandez et al., 2010; Peter et al., 2015). The transition from EB to EC demands large amount of protein synthesis to drive cell growth, which could explain how the up-regulation of Thor blocks EC specification in mutant midguts.

To test this idea, we knocked down Thor using esg F/O-driven RNAi. We found that the number of GFP+ ECs was partially restored (Figure 6, B and C), confirming that the blockage of EC specification is partially mediated by Thor. Additionally, activating the mTOR pathway that inhibits 4EBP, via Tsc2 knockdown (Kapuria et al., 2012), also led to significantly enlarged GFP positive cells in mutant midguts (Figure 6, D and E), indicating a partial rescue of the EC specification.

So far, our data indicate that ETC deficiencies elevate the FOXO protein level, which blocks EC specification through the 4EBP homologue Thor. However, foxo knockdown only restored the expression of a subset of genes altered in the mutant ISCs (Figure 6A and Supplemental Table S2). Additionally, neither foxo nor thor knockdown restored the number of GFP+ EE cells in mutant midguts (Figures 5D and 6C). These observations suggest that pathways other than foxo are likely to be involved in regulating EE specification.

Discussion
Pharmacological disruptions of ETC complexes in cultured embryonic stem cells impair their proliferation and differentiation (Chung et al., 2007; Mandal et al., 2011). However, how mitochondria exert these regulations is still enigmatic. Besides the fact that these studies were carried out in vitro and were complicated by potential side effects of drugs, little evidence has been provided to clarify the role of mitochondrial respiration in regulating stem cell activity in vivo, especially for somatic stem cells that are necessary to maintain tissue homeostasis during aging. To address this question, we genetically disrupted ETC in Drosophila intestine stem cells by manipulating their mitochondrial genomes while keeping the environmental conditions controlled.

Table S1). The down-regulated pool included many genes involved in cell differentiation, proliferation, and growth regulation (Supplemental Table S2). The up-regulated pool included genes participating in various metabolic processes (Supplemental Table S2). This result suggests a possible metabolic adaptation in response to theOXPHOS deficiency in mutant midguts.
epithelial niche intact. We found that ETC deficiencies impaired ISC's proliferation and lineage specification. The defects could be a common consequence of ETC disruption, as similar phenotypes were observed among genetic disruption of different complexes, although the penetrance varied (Figure 4B; Supplemental Figure S3, B–D). Disruption of complexes I and II had a mild effect on ISC proliferation and lineage specification, whereas disruption of complexes III or IV resulted in much stronger defects (Figure 4B; Supplemental Figure S3, B–D), similar to that in ISC carrying the "ts" allele. Mitochondrial respiration begins with the oxidation of either NADH or succinate by NADH dehydrogenase (Complex I) and succinate dehydrogenase (Complex II), respectively (Figure 4A). The partial redundancy between complexes I and II may explain mild phenotypes in RNAi against these two complexes.

The most obvious consequence of ETC disruption is ATP deficiency. Although AMPK, the classical energy sensor, is not involved in the manifestation of the ISC phenotype (Supplemental Figure S5, B and C), we cannot rule out the possibility that ATP deficiency might inhibit ISC proliferation through other unnoted processes independent of AMPK. Alternatively, the ISC phenotype may specifically pertain to inhibition of the cytochrome chain in complexes III and IV.

The leakage of free electrons from the cytochrome chain to oxygen is a main source of cellular ROS (Starkov, 2008; Bleier and Drose, 2013). However, in our system, disruption of ETC did not lead to ROS accumulation (Supplemental Figure S6, A and B). Additionally, overexpression of ROS scavengers did not rescue the mutant phenotype (Supplemental Figure S6, C and D), further supporting that ROS accumulation is not the cause of the ISC proliferation defect. Nonetheless, in mutant ISCs, we found elevated expression of FOXO, a transcription factor known to be up-regulated in response to ROS stress and to mediate stem cell behavior (Cheung and Rando, 2013; Klotz et al., 2015). Importantly, we found that the elevated FOXO activity is partially responsible for the mutant defect. Hyperactivation of FOXO and elevated FOXO levels have been documented in human disease accompanied by mitochondrial dysfunction and increased oxidative stress (Kannike et al., 2014). The increased FOXO protein levels in these contexts result from FOXO autoregulation via a positive feedback loop. Such a positive autoregulation is also reported in the Drosophila adult gut (Alic et al., 2014) and may remain active in the mutant ISCs. How ETC disruption leads to elevated FOXO activity in our system is currently unclear. Aside from ROS, the mitochondrial unfolded protein response (UPR<sub>mt</sub>) causes lasting FOXO activation during Drosophila development (Borch Jensen et al., 2017). Could the reduced availability of mtDNA-encoded subunits in mutant ISCs result in UPR<sub>mt</sub> and eventually activate FOXO? In this case, ETC deficiencies may impact ISC activity indirectly. Recent studies have suggested that some ETC complexes may have other functions besides respiration (Teixeira, Sanchez, Hurd, et al., 2015; Meng, Yamashita, et al., 2017). The possibility of a specific impact of complexes III and IV on FOXO can therefore not be eliminated, though the exact mechanisms remain obscure at this point.
**Figure A**

Images showing GFP-FOXO and FOXO expression in cellular contexts.

**Figure B**

Bar graph showing GFP+ area in whole midgut (%).

**Figure C**

Images showing GFP-β-gal expression under different conditions.

**Figure D**

Graph depicting the number of GFP+ cells per mm² under control (ctrl) and FOXO knockdown (KD) conditions.
As for the impact of elevated FOXO on the ISC lineage, several scenarios are possible. Drosophila FOXO functions as a key negative regulator of growth and proliferation, probably via 4EBP transcription (Puig et al., 2003). Indeed, we found that overexpression of FOXO in normal ISCs blocked their lineage specification (Figure 5B and Supplemental Figure 5B) and inhibition of FOXO in mutant ISCs partially rescued the defect (Figure 5, C and D), as did down-regulation of 4EBP (Figure 6, B and C). Considering the large size of EC cells that demand prodigious protein synthesis, it is not far-fetched to propose that activation of the FOXO-4EBP cascade could inhibit the formation of EC cells through its inhibition on protein translation and cell growth. Additional support for this notion stems from our observation that activating mTOR, a protein kinase that inactivates FOXO, and further support our finding that the intact ETC and aerobic metabolism are essential for ISCs’ activity and midgut epithelial homeostasis.

MATERIALS AND METHODS

Fly stocks and transgenic lines

Fly strains used in this study including the following: w;esg-GAL4, tub-GAL80Ts, UAS-nlsGFP;CyO;UAS-flp,Act-cD>Gal4/TM6B; Myosin II-Gal4; UAS-SOD2; Catalase; AMPKα RNAi (BDSC#25931); foxo RNAi (BDSC #27656); UAS-FOXO (BDSC #42221); ND30 (30D) RNAi (VDRC #103412); ND812 (B12) RNAi (VDRC #104890); ND23 (Tkyk) RNAi (VDRC #21748); Sdha RNAi (VDRC #110440); Sdch RNAi (VDRC #6031); Sdch RNAi (VDRC #101739); Cytochrome c1 (cyt c1) RNAi (VDRC #109809); UQCR-C1 (cp1) RNAi (VDRC #101350); UQCR-C2 (cp2) RNAi (VDRC #100818); COX5A (Va) RNAi (VDRC #109070); COX7C (VIIc) RNAi (VDRC #104970); COX8B (VIII) RNAi (VDRC #104047); ATPsynB (b) RNAi (VDRC #100758); ATPsynC (c) RNAi (VDRC #106834); ATPsynD (d) RNAi (VDRC #104353); Thor-lacZ (4EBP-lacZ) (BDSC #49558); Thor RNAi (BDSC #36815); Tsc2 RNAi (BDSC #35401); UAS-pAbp (BDSC #9420, for tissue-specific mRNA tagging (Yang et al., 2005)); and Luciferase RNAi (BDSC #31603, control for RNAi lines). Female flies of heteroplasmic mt:Co1T303 and homoplasmic mt:Co1pm background as previously described (Xu et al., 2008; Chen et al., 2015) were used to generate the mutant (M) group and control (C) group flies for assessments, respectively. All flies and crosses were maintained at 18°C.

Tissue preparation, immunofluorescence staining, and confocal imaging

All fly crosses and the offspring were kept at 18°C. Two days after eclosion, the adult flies for assay were heat-shocked at 37°C for 2 h and maintained at 29°C until dissection at different day points. The entire midgut tissues were dissected out in room temperature Schneider’s medium (Thermo Fisher Scientific) supplied with 10% heat inactivated fetal bovine serum (FBS; Thermo Fisher Scientific). The midguts were then rinsed by the same medium and used for either direct imaging or further fixation and staining.

Immunofluorescence staining was done after tissue fixation in phosphate-buffered saline (PBS) containing 4% Paraformaldehyde (PFA) and three times PBS washing. The antibodies and reagents used for staining were mouse anti-Prospero (MR1A; Developmental Studies Hybridoma Bank [DSHB]); mouse anti-delta (C594.9B; DSHB); chicken anti-beta galactosidase (ab9361; Abcam); rabbit anti-Pdm1 (gift from Yu Cai, Temasek Life Sciences Laboratory, National University of Singapore); rabbit anti-cleaved caspase-3 (9661; Cell Signaling); rabbit anti-FOXO antibody (ab19597; Abcam); GFP tag polyclonal antibody, Alexa Fluor 488; Alexa Fluor secondary antibodies (Thermo Fisher Scientific), and Vectashield mounting medium with 4′,6-diamidino-2-phenylindole (DAPI) (Vector Laboratories).

Imaging analysis was performed using the Carl Zeiss Axio Observer equipped with the Perkin Elmer spinning disk confocal system. For live imaging, a Zeiss incubation system was used to maintain proper temperature and humidity. The image processing and quantification were performed by Velocity 6.1.1 software (Perkin Elmer) and ImageJ software (National Institutes of Health [NIH]). Ten midguts were analyzed in each group. The region of the posterior midgut, about 500 μm from the junction of the midgut and hindgut, was used for the quantification of GFP+ cell numbers.

Detection of mitochondrial membrane potential

Mitochondrial potential was detected using a protocol adopted from a previous study (Zhang et al., 2019). Briefly, after dissection, intact guts were incubated in Schneider’s medium containing 2% (v/v) heat inactivated fetal bovine serum (FBS; Thermo Fisher Scientific) and MitoTracker Deep Red (200 nM; Thermo Fisher Scientific) at 29°C for 20 min and rinsed with medium three times. Samples from different groups were mounted with medium on the same coverslip set in a custom-made metal frame and then covered with a small piece of Saran Wrap before imaging. Images were captured with fixed
FIGURE 6: The impaired EC specification is partially mediated by 4EBP in mutant midguts. (A) Diagrams summarizing the number of genes up- or down-regulated by ETC disruption and restored by foxo knockdown (foxo\textsuperscript{KD}). The cut-off was 0.5 CPM (counts per million) fold change averaged across two replicas for regulation by ETC disruption, and 0.25 CPM fold change on average for restoration by foxo\textsuperscript{KD}. Full raw data are presented in Supplemental Table S1. (B) Views of posterior midguts expressing a 4EBP knockdown construct (\textit{esg}\textsuperscript{ts}F/O\textendashmitoXhoI, 4EBP\textsuperscript{KD} flies) compared with a control RNAi (\textit{esg}\textsuperscript{ts}F/O\textendashmitoXhoI, luciferase\textsuperscript{KD} flies) showed partial rescue of GFP\textsuperscript{+} EC numbers in mutant (M) background. Green: GFP; Red: Prospero; Blue: DAPI. Scale bars: 20 µm. (C) Quantification of the GFP\textsuperscript{+} ECs and GFP\textsuperscript{+} EEvs in the posterior parts of control (C) and mutant midguts (M) in the presence or absence of 4EBP\textsuperscript{KD}, 10 d after the heat shock. \(n = 10\) midguts each group, error bar: SD; *: \(p < 0.001\), NS: not significant. (D) Activation of mTOR pathway...
confocal settings. These small, isolated, and basally localized GFP+ cells within the intestinal epithelium were identified as ISCs in esg−/+ GFP flies based on a previous study (Hochmuth et al., 2011). A single section of basal epithelium was captured, and TMRM and MitoTracker signal intensities were quantified using ImageJ. Mitochondrial membrane potential was computed as the ratio of the mean intensity of TMRM to MitoTracker fluorescence with background correction.

**ATP measurement**

The tissue ATP level was measured using the ATP Bioluminescence Assay Kit HS II (Roche) following the manufacturer’s protocol. Wild-type and homoplasmic ts female flies were raised at 18°C for 2 d after eclosion and then shifted to 29°C for 3 d, or continuously maintained at 18°C as control. For each genotype, individual midgut was dissected out and immediately homogenized in 100 μl lysis buffer. The lysates were boiled for 5 min and centrifuged at 20,000 × g for 1 min. Cleared lysate (10 μl) was used for ATP measurement using the kit through a 96-well plate luminometer (Berthold). The amount of ATP was then determined and normalized with the protein level measured by BCA protein assay (Thermo Fisher Scientific).

**Detection of ROS levels**

ROS levels were detected using CellROX Deep Red following the manufacturer’s protocol (Thermo Fisher Scientific). Briefly, the guts were incubated in Schneider’s medium containing 5 μM CellROX Deep Red for 45 min at 29°C followed by three washes with PBS. The samples were then fixed with 3.7% formaldehyde for 15 min and mounted for imaging analysis. For comparison and quantification, the dissected guts were preincubated in Schneider’s medium with or without drugs for 1 h at 29°C before CellROX staining. Control gut incubated in the medium containing 200 μM tert-butyl hydroperoxide (TBHP) was used as the positive control, and gut incubated with 1 mM N-acetylcysteine (NAC) was used as the negative control. Images were captured with a spinning disk confocal system. These small, isolated, and basally localized GFP+ cells within the intestinal epithelium were identified as ISCs in esg−/+ GFP flies based on a previous study (Hochmuth et al., 2011). A single section of basal epithelium was captured, and CellROX intensity was quantified using Volocity 6.1.1 software (Perkin Elmer).

**GFP-positive ISCs sorting and mtDNA level**

The GFP+ ISCs were sorted as previously described (Dutta et al., 2013). Briefly, midguts of “w; esg-GAL4, tub-GAL80Δ UAS-nlsGFP/ UAS-mitoXhol” flies with different mtDNA background, which were heat-shocked and maintained at 29°C for 7 or 15 d, were dissected in PBS and dissociated for 2 h by incubating with 0.8 mg/ml elastase at 29°C. ISCs were then sorted based on intensity of the GFP marker with the help of the Flow Cytometry (FACS) Core Facility of the National Heart, Lung, and Blood Institute (NHLBI), NIH. To quantify the mtDNA level, total DNA was prepared from the sorted samples using the DNeasy Blood kit (Qiagen). Quantitative real-time PCRs were performed in triplicate using the Roche LightCycler 480 system and SYBR Green I Master. Primers based on mtCol (F: ATTTGAGTGAATTTACGATTTTCCTCAA; R: AGTT-GATAAAATTTATGTTG- GTAAG) and His4 (F: TCCAG-GGATTCAGAAAGCC; R: AACCTTCAGAACCCAC) sequences were used for the mtDNA and nuclear DNA, respectively.

**EdU Incorporation**

EdU incorporation was determined following a protocol previously described (Zhang et al., 2015), using the Click-iT EdU Alexa Fluor 555 Imaging kit (Thermo Fisher Scientific). Briefly, intact guts were dissected from adult female flies and were incubated in 10 μM EdU in Schneider’s medium containing 10% FBS for 2 h at 25°C. Tissues were then fixed with 4% PFA, washed in PBS with 0.1% Triton X-100 (PBST), processed with the kit, and mounted for imaging analysis.

**ISC specific mRNA profiling**

A modified mRNA tagging technique (Yang et al., 2005) was used to isolate mRNA from ISCs of fly midguts. Briefly, the FLAG-tagged poly(A)-binding protein (pAbp) was expressed in C and M group flies under esg-GAL4 driver. About 200 midguts from C and M group “esg-GAL4,tub-GAL80Δ,UAS-nlsGFP/UAS-mitoXhol;UAS-pAbp-FLAG/+” or “esg-GAL4,tub-GAL80Δ,UAS-nlsGFP/UAS-mitoXhol;UAS-pAbp-FLAG/UAS-FOXO-IR” flies, which were heat-shocked and maintained in 29°C for 5 d, were fixed in 1 ml of PBS containing 1% formaldehyde (Sigma-Aldrich) and 0.5% Igepal CA-630 (Sigma-Aldrich) for 1 h on ice and then incubated for 5 min after the addition of 140 μl of 2 M glycine. Samples were rinsed three times with ice cold PBS and homogenized by sonication in 2 ml of homogenization buffer (150 mM NaCl, 50 mM HEPES buffer at pH 7.6, 1 mM ethylene glycol tetraacetic acid (EGTA), 15 mM EDTA and 10% glycerol) supplied with 8 mM vanadyl ribonucleoside complex (Sigma-Aldrich), 50 U/ml SUPERase-In (Thermo Fisher Scientific), and 1 tablet/50 ml protease inhibitor cocktail tablet (Roche). After centrifugation at 13,000 × g for 30 min, the supernatant was incubated with anti-FLAG-M2 affinity agarose beads (Thermo Fisher Scientific) at 4°C overnight (O/N). The beads were then washed four times with homogenization buffer. The RNAs bound to the pAbp were recovered in 100 μl of elution buffer (50 mM Tris- HCl, pH 7.0, 10 mM EDTA, 1.3% SDS, and 50 U/ml SUPERase-In) by proteinase K (40 μg/ml) treatment for 2 h at 60°C and O/N at 65°C after addition of NaCl (200 mM). Eluant was then mixed with 4 vol of Trizol (Thermo Fisher Scientific) and 1 vol of chloroform (Sigma-Aldrich) with vigorous shaking, incubated at room temperature for 10 min, and then centrifuged at 12,000 × g for 15 min at 4°C. The supernatant was transferred and mixed with 2.5 vol of isopropanol, chilled at −80°C O/N, and centrifuged at 12,000 × g for 30 min at 4°C. The RNA pellet was rinsed twice with 70% ethanol and then dissolved in RNase free water. Poly(A) capture libraries were generated at the DNA Sequencing and Genomics Core, NHLBI, NIH. RNA sequencing was performed with a HiSeq3000 (Illumina), and 75–base pair pair-end reads were generated at the DNA Sequencing and Genomics Core, NHLBI, NIH. RNA sequencing data were analyzed by the Bioinformatics and Computational Biology Core, NHLBI, NIH. After quality control (QC) assessment of FASTQ files using FastQC toolkit (v0.11.5) with default parameters. paired-end reads were aligned against the Drosophila melanogaster reference genome by Tsc2 knockdown in ISCs showed enlarged GFP+ cells in mutant midguts (esg−/F/O-mitoXhol, Tsc2ΔO flies), indicating a partial rescue. Assessed 10 d after the heat shock. Scale bars: 20 μm. (E) Quantification shows that Tsc2 knockdown (KD) in ISCs significantly increased the GFP+ midgut area due to enlarged GFP+ cells in mutant midguts, 10 d after the heat shock. n = 10 midguts each group, error bar: SD; *: p < 0.001.
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acknowledgments

statistical analysis

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