Ferritin is required at multiple stages during the embryonic development of *Drosophila melanogaster*.

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In *Drosophila*, iron is stored in the endomembrane system of cells inside a protein cavity formed by 24 ferritin subunits of two types (Fer1HCH and Fer2LCH) in a 1:1 stoichiometry. Ferritin accumulates in the midgut, nervous system, hemolymph and nephrocytes of *Drosophila* larvae. Here, we show that mutation of either ferritin gene product or deletion of both genes resulted in a similar set of phenotypes of embryonic lethality, ranging from non-deposition of cuticle to developmental defects associated with germ band retraction, dorsal closure and head involution. Maternal contribution of ferritin, which varied reflecting the mother’s iron stores, is used in early development, but zygotic ferritin mutants died with ectopic apoptotic events and disrupted intestinal morphology. The embryonic nervous system of ferritin mutants presented ventral nerve cord disruptions, misguided axonal projections and brain malformations. Finally, ferritin accumulation was also observed in embryonic hemocytes. One ferritin mutant showed no hemocyte ferritin accumulation and this expression was also lost by genetic inhibition of the secretory pathway. Our work suggests that insect ferritin functions in iron storage, intercellular iron transport and protection from oxidative stress at multiple times during the embryonic development of *Drosophila melanogaster*. 
INTRODUCTION

Iron is the most abundant transition metal on earth, commonly found at the active sites of proteins in the form of heme or iron-sulfur clusters or as mono-nuclear and di-nuclear iron (Sheftel et al. 2012). Because of the high reactivity between iron and oxygen, iron has become a key player in aerobic metabolism but also catalyzes oxidative stress when present in excess. Therefore, iron concentration within subcellular compartments and in extracellular fluids is tightly regulated (Cabantchik 2014). The Divalent Metal Transporter 1 (DMT1) is responsible for cellular iron uptake (Gunshin et al. 1997). Ferritin participates in iron homeostasis as the main iron storage complex in both prokaryotes and eukaryotes (Harrison & Arosio 1996). The major form of ferritin in vertebrate animals is cytosolic and consists of 24 subunits of H and L protein chains that assemble into a cage-like structure that sequesters up to 4,500 atoms of iron in its interior core. The H chain contains a ferroxidase center necessary for iron internalization while the L chain contains acidic groups exposed in the interior surface of holoferritin facilitating iron mineralization (Santambrogio et al. 1993). Ferritin genes are regulated during translation by the binding of an iron regulatory protein (IRP) to an iron responsive element (IRE) located in the 5' untranslated region of the mRNA (Pantopoulos et al. 2012). The discovery in mice of ferritin receptors Scara5 (Li et al. 2009) and Tim2 (Todorich et al. 2008) has lead to the idea that ferritin might be involved in iron transport (Meyron-Holtz et al. 2011), however this idea remains controversial (Kell & Pretorius 2014).

In insects, ferritin shells have an H₁₂L₁₂ organization due to inter- and intra- subunit disulfide-bonds ensuring protein folding and assembly (Hamburger et al. 2005). Ferritin intracellular localization in most insects is directed to the endoplasmic reticulum and the Golgi (Missirlis et al. 2006; Missirlis et al. 2007). The Drosophila melanogaster genome encodes for three ferritin genes: Ferritin 1 heavy chain homologue (Fer1HCH) and Ferritin 2 light chain homologue (Fer2LCH) together produce the major ferritin complex (Georgieva et al. 2002; Georgieva et al. 1999; Mandilaras et al. 2013), whereas Ferritin 3 heavy chain homologue (Fer3HCH) encodes the mitochondrial ferritin, which is predominantly expressed in testis (Kumar et al. 2011; Missirlis et al. 2006). The Drosophila Iron Regulatory Protein-1A (IRP-1A), in its iron-sulfur cluster depleted modality, binds IREs thereby regulating the translation of a subset of Fer1HCH mRNA transcripts (Lind et al. 2006). The sole Drosophila DMT1 homolog (Folwell et al. 2006) was originally isolated as a gustatory mutant named...
Malvolio (Mvl) (D’Souza et al. 1999). Previous work has shown that ferritin is required for embryonic and larval development (Missirlis et al. 2007; Tang & Zhou 2013) and that the ferroxidase center of the H chain is essential (Missirlis et al. 2007), but the specific phenotypes of the ferritin mutants have not been studied to date. Drosophila ferritin has been proposed to function as an external source of iron (Li 2010; Nichol et al. 2002; Tang & Zhou 2013; Zhou et al. 2007) but functional analysis in support of this proposition exists only for the secreted ferritin of ticks (Galay et al. 2013; Galay et al. 2014; Hajdusek et al. 2009).

Here, we analyze Fer1HCH and Fer2LCH mutant embryonic phenotypes. We propose that key functions of the ferritin subunits are likely mediated through the ferritin complex, as their respective mutant phenotypes are indistinguishable during embryogenesis. We show that ferritin mutant phenotypes can be enhanced when embryos are deprived of or have a reduced ferritin maternal contribution, by induction of germ line clones or by limiting iron uptake in parental diets, respectively. A severe central nervous system (CNS) defect, ectopic apoptosis and intestinal damage are seen in embryos dying at later stages of development. Blocking the intracellular secretory pathway during embryogenesis results in the mislocalisation of ferritin. We hypothesize that failure of ferritin transport in one of the mutants tested contributes to the resulting phenotype and lethality.

RESULTS AND DISCUSSION

Pleiotropic phenotypes of ferritin mutants revealed from embryonic cuticle preparations

To characterize the embryonic lethal phenotype of ferritin mutants we analyzed the cuticles of previously described transposon-induced loss-of-function alleles Fer1HCH451 and Fer2LCH35 (Missirlis et al. 2007; Tang & Zhou 2013), a null mutation for both ferritin genes Df(3R)Fer (Gutierrez et al. 2013), a GFP-trap line Fer1HCHG188 (Missirlis et al. 2007) and also a new allele described here for the first time Fer2LCHΔ17, which fails to complement Fer1HCH451, Fer1HCHG188 and Fer2LCH35 (see Materials and Methods). Cuticle preparations of ferritin mutants revealed examples of developmental arrest during the key embryonic processes of germ band extension and retraction, dorsal closure and head involution (Figure 1). Nevertheless, quantification of phenotypes suggests that most
mutant embryos presented either a normal cuticle (~75%) or no cuticle at all (~15%). These phenotypes suggested that ferritin is required multiple times during development, consistent with differential requirements for iron to support the metabolic shifts that occur during development (Tennessen et al. 2014). Notably, a number of iron sulfur cluster proteins are induced during the final stages of embryogenesis to support aerobic glycolysis.

A maternal contribution of ferritin is utilized during early embryonic development

Ferritin is maternally contributed (Missirlis et al. 2007). We therefore wondered if our analysis of zygotic mutants could miss early requirements for ferritin fulfilled by this maternal contribution. To analyze how maternal ferritin functioned during embryogenesis we followed two strategies: curtailing iron availability in mothers and generating female germ line clones without ferritin. One way of reducing ferritin expression in adults is to add an iron-specific chelator in their diet (Gutierrez et al. 2013; Missirlis et al. 2006). We hypothesized that reduced overall ferritin levels would result in decreased ferritin maternal contribution and a more severe embryonic phenotype. Homozygous mutant embryos derived from heterozygous adults grown with food containing 200µM Bathophenantrholine Sulfate (BPS) showed a doubling of incidence for the no cuticle phenotype (from ~15% to ~30%; Figure 1). Lack of cuticle deposition was likely due to early embryonic death before epidermal differentiation, although we have not excluded a particular requirement for iron in the differentiation of the epidermis, which could provide an alternative explanation for the same phenotypic outcome. Late embryonic phenotypes also became more frequent: U-shaped embryos, indicative of a failure to retract the germ band, augmented from ~2% to ~4%, and embryos failing to complete dorsal closure increased from ~1% to ~4% (Figure 1F; BPS columns). At the same time, the percentage of “wild type” cuticles decreased. This evidence shows that the extent of ferritin maternal contribution is at least partially regulated in flies via limited iron availability. We note that the changes in phenotypic classes abundances were consistently similar in Fer1HCH451, Fer2LCH35, and Df(3R)Fer. Feeding extra iron to adults did not result in a rescue of the embryonic phenotype of the mutant offspring (Figure 1F, FAC), even though total levels of ferritin in mothers were increased, showing that maternal ferritin contribution was not sufficient to fully rescue embryogenesis.

To further explore whether maternal ferritin was partially rescuing zygotic ferritin mutants, especially...
during early development, we generated homozygous mutant germ line clones for the ferritin genes. Most germ line clones had no cuticle or bore cuticles with defects. The ‘no cuticle’ phenotype percentage changed spectacularly from ~15% to ~80% (Figure 1F). Heterozygous embryos without maternal contribution developed normally into adults (data not shown). We hypothesize that this rescue was due to zygotic ferritin genes being overexpressed as a response to lack of maternal ferritin and that iron in these embryos was delivered to the developing oocyte by other means.

Central Nervous System (CNS) phenotypes of ferritin mutants

The majority of zygotic ferritin mutant embryos derived from heterozygous mothers died at the final stages of embryogenesis with an apparently normal cuticle. Therefore, we sought to study internal tissues that might be affected by lack of ferritin. Since ferritin protein in first instar larvae is concentrated in the intestine and the CNS (Mehta et al. 2009), we asked whether the CNS developed normally in ferritin mutants. We performed immunofluorescence with antibodies against neuronal markers. The application of an antibody against Elav, which marks all neuronal nuclei, revealed that ferritin mutants harbor holes in the abdominal segments of the CNS (Figure 2). More severe phenotypes were also present, albeit in fewer embryos, including aberrant condensation of the CNS, twisted CNS, and complete lack of parts of the brain and peripheral nervous system (PNS). Importantly, and consistent with our analysis of the cuticle phenotypes discussed above, these phenotypes were observed with both ferritin alleles and with the 2.2 kb genomic deletion that specifically deletes both Fer1HCH and Fer2LCH (Figure 2). We interpret this to mean that, in wild type embryos individual ferritin subunits function in concert to assemble the ferritin complex and do not carry out subunit-specific functions.

It was previously reported that in adult flies, RNA interference in subsets of neurons against Fer2LCH but not against Fer1HCH disrupted circadian rhythms (Mandilaras & Missirlis 2012). Furthermore, some cell types, including commonly used cell culture lines (Metzendorf et al. 2009; Missirlis et al. 2003a), only express Fer1HCH and not Fer2LCH. Finally, overexpression of either Fer1HCH or Fer2LCH or both subunits simultaneously in Drosophila glia (Kosmidis et al. 2011) or neurons (Wu et al. 2012b) resulted in qualitatively different responses. Thus, the question of how each cell type regulates the two ferritin genes and subunits and whether they always act in concert, as the analysis of
mutant embryonic phenotypes suggests, requires further investigation.

Further markers were used in embryos homozygous for \textit{Df(3R)Fer}. We tested whether neuronal axons were projecting normally using α-BP102, which stains axons. BP102-dependent fluorescence revealed that CNS axons are frequently misguided (Figure 3A, D). The developing CNS consists of at least four types of cells: neuroblasts, ganglion mother cells, neurons and glia. Neuroblasts give rise to ganglion mother cells, and these, in turn, give rise to neurons and glia \cite{Biffar&Stollewerk2014}. In order to test whether neuroblasts were affected in ferritin mutant embryos, we performed antibody staining with Deadpan and Evenskipped (Eve) antibodies \cite{Boone&Doe2008,Kohwi2013}. Our preparations with antibodies against Deadpan, which marks all neuroblasts, indicate that CNS defects are already present within neuroblast cell lineages, including misplaced neuroblasts (Figure 3B, E). As expected for early CNS defects, Eve positive neuroblasts were also heavily misplaced (Figure 3C, F). In conclusion, the developing CNS of embryos lacking the ferritin genes is affected from the time neuroblast cell lineages are specified giving rise to contorted and aberrant CNS in late embryos.

A ferritin enhancer trap shows expression in the CNS

\textit{Fer1HCH^{451}} carries a \textit{LacZ} element that serves as reporter of \textit{Fer1HCH} expression. To test whether this reporter recapitulates known changes in \textit{Fer1HCH} expression, we monitored \textit{LacZ} activity in the anterior midgut upon iron feeding of larvae, as occurs for endogenous \textit{Fer1HCH} (Figure S1). Iron-dependent induction of expression was confirmed with this reporter and we therefore used \textit{LacZ} detection to monitor \textit{Fer1HCH} expression in the embryo. \textit{LacZ} driven from \textit{Fer1HCH^{451}} is strongly expressed in the neuroectoderm (Figure 4A). The co-localization between the neuronal marker Elav and the β-galactosidase reporter was also observed in homozygous mutant embryos with a disrupted CNS (Figure 4B). We note that the use of the \textit{Elav-Gal4} driver to silence either \textit{Fer1HCH} or \textit{Fer2LCH} resulted in viable adults with disturbed circadian behavior \cite{Mandilaras&Missirlis2012} and apparent neurodegeneration \cite{Tang&Zhou2013}. RNA interference is known to cause reduced expression but not complete silencing of its targets, which may explain why the RNAi flies survived to adulthood. In addition, overexpression of ferritin subunits with \textit{Elav-Gal4} failed to rescue their respective mutants \cite{Tang&Zhou2013}. Nevertheless, our findings suggest that ferritin is expressed in the embryonic CNS and is required for its development.
Ectopic apoptotic activation in ferritin mutants

Several links have been drawn in the last decade between iron metabolism and apoptosis (Gambis et al. 2011; Kosmidis et al. 2011; Pham et al. 2004). We hypothesized that the disrupted CNS would result following cell death by an apoptotic mechanism. To test the hypothesis we used an antibody that recognizes solely the cleaved, activated caspases, and has been used in flies to mark apoptotic cells (Denton et al. 2008). In contrast to control embryos at stage 12 where there was no apoptotic signal detected (Figure 5A), ectopic apoptotic activation appeared in the neuroectoderm region in mutant embryos (Figure 5B). By stage 15 of embryogenesis control embryos have a weak and restricted apoptotic signal (Figure 5C) whereas in the mutant embryos this signal was massive and covered most of the embryo (Figure 5D). Similar widespread immunoreactivity against activated caspase was also observed with Fer1HCH^{451} and Fer2LCH^{35} homozygous mutant embryos (data not shown). Thus, the early apoptotic activation seen in ferritin mutants, mainly restricted to the neuroectoderm region, suggests that the CNS may be more susceptible to the lack of the ferritin complex, although, ultimately, many other tissues become affected.

The Drosophila DMT1 homolog, Malvolio, is upregulated in ferritin mutants

Mutations in Mvl result in reduced iron within intestinal iron storage cells and in the whole body (Bettedi et al. 2011). Mvl mutants can suppress both intestinal iron accumulation resulting from ferritin (Tang & Zhou 2013) and Multicopper Oxidase-1 (MCO1) (Lang et al. 2012) misregulation. MCO1 catalyzes the oxidation of ferrous iron (Fe^{2+}) to ferric iron (Fe^{3+}) and is thought to participate in cellular iron export in insects (Lang et al. 2012). In view of the above findings, we used Mvl^{97f}, a P-element insertion mutant carrying LacZ reporter gene activity (Rodrigues et al. 1995) to assess β-Galactosidase activity staining. In control embryos, LacZ showed a very restricted pattern of Mvl expression consistent with previous studies (Kumar et al. 2011; Rodrigues et al. 1995), however in the absence of ferritin, Mvl-driven LacZ is upregulated (Figure 6A). We hypothesize than in the absence of functional ferritin, iron depleted cells upregulate Mvl. Strikingly, the addition of one copy of Mvl^{97f} hypomorphic allele into a ferritin-depleted embryo resulted in the appearance of melanotic spots, which were never observed in Mvl^{97f} or in ferritin homozygous mutants alone (Figure 6B, C). The interaction between
DMT1 and ferritin is reminiscent of the recent finding that the mammalian ferritin receptor SCARA5 is likewise upregulated in the absence of transferrin receptor (Li et al. 2009).

**Ferritin expression, localization and trafficking during development**

*Fer1HCH<sup>G188</sup>* is a mutant *Fer1HCH* allele, shown to generate a chimeric GFP-Fer1HCH that faithfully mimics endogenous Fer1HCH pattern in heterozygous condition (Gutierrez et al. 2010; Mehta et al. 2009; Missirlis et al. 2007; Uhrigshardt et al. 2013). During stages 16-17 of embryonic development GFP tagged Fer1HCH is present in hemocytes (Figure 7A). Hemocytes are large cells that are loosely associated with peripheral tissues and can circulate in the hemolymph, where they function as both phagocytic and immune cells (Evans & Wood 2011). In order to confirm that the large, ferritin-accumulating cells were actually hemocytes we used Cg-Gal4 line to drive expression, exclusively in hemocytes, of a nuclearRFP in *Fer1HCH<sup>G188/+</sup>* embryos (Figure 7A). GFP-Fer1HCH is present in the same cells as Cg-nRFP, but not in FB-nRFP (Figure 7B), where mRNA expression is seen (Missirlis et al. 2007). The GFP tag is thought to block the correct function of ferritin in homozygous *Fer1HCH<sup>G188</sup>* embryos, because if all the H-subunits carry a GFP tag, embryonic development fails (Missirlis et al. 2007) and even *Fer1HCH<sup>G188/+</sup>* flies show a mild reduction in iron accumulation within ferritin (Gutierrez et al. 2013). We noticed that homozygous mutant *Fer1HCH<sup>G188</sup>* embryos showed a dramatically reduced expression of GFP-Fer1HCH than similarly staged heterozygous *Fer1HCH<sup>G188</sup>* embryos (Figure 7E, F). Lower and restricted GFP-Fer1HCH expression in homozygous *Fer1HCH<sup>G188</sup>* embryos, mainly seen in the intestinal region (Figure 7F), suggested either degradation of the mutant protein or that mutant, non-functional heteropolymers (composed exclusively of GFP-Fer1HCH and Fer2LCH subunits) would not be trafficked. Furthermore, we noted that the intestinal morphology of homozygous *Fer1HCH<sup>G188</sup>* was affected (Figure 7F). To test further whether ferritin is delivered to hemocytes from other embryonic tissues and whether ferritin trafficking can be observed, we blocked the intracellular secretory pathway by means of a lethal mutation in Sec23, *sec23<sup>j13C8</sup>* , a P-element insertion in the 5’ UTR of *sec23*, expected to eliminate or severely attenuate gene function (Abrams & Andrew 2005). If ferritin is indeed transported during embryogenesis, blocking the secretory pathway will impede its exit from the cell where it was originally transcribed. Ferritin was not detected in hemocytes of *sec23<sup>j13C8/j13C8</sup>* mutants; rather, GFP-Fer1HCH aggregates were detected mainly around the midgut (Figure 7D) in a similar expression pattern as observed in homozygous *Fer1HCH<sup>G188</sup>*
mutants. Thus, in embryos impaired in the secretory pathway hemocytes fail to accumulate ferritin, suggesting that its source in wild type embryos may come from hemolymph ferritin. Taken together, our observations suggest that ferritin traffics between tissues during embryonic development and that the hemocytes could play a key role during this trafficking process, by importing ferritin from one tissue and delivering it to another. Such a communicating role for hemocytes has been previously suggested in the context of tissue communication in the innate immune response (Wu et al. 2012a).

Conclusions

Insect embryos must course through development with limited amounts of iron, provided during oogenesis by the mother in part via ferritin. It seems reasonable to assume that tissues developing at different rates and with different metabolic states present different iron requirements. Therefore, iron storage and transport are of vital importance for normal development. Ferritin appears to be serving both functions in *Drosophila*.

EXPERIMENTAL PROCEDURES

Fly stocks

As a control strain *y,w* flies were used. *Fer1HCH*<sup>451</sup> and *Fer2LCH*<sup>35</sup> are P(ry[<sup>+t7.2</sup>] = PZ) insertion alleles generated during a large-scale mutagenesis screen (Spradling et al. 1999), they have been partially characterized elsewhere (Missirlis et al. 2007) and they were obtained from the Bloomington Drosophila Stock Center (BDSC) stock numbers #11497 and #11483, respectively. *Fer2LCH*<sup>Δ17</sup> was generated from an imprecise excision of *Fer2LCH*<sup>EP1059</sup> (described in Flybase) and interferes with expression of both genes, as confirmed by complementation crosses. Df(3R)*Fer* was a gift from Alexis Gambis, Bertrand Mollereau and Hermann Steller and is a 2.2 kb deletion disrupting specifically *Fer1HCH* and *Fer2LCH* (Gutierrez et al. 2013). To generate germline clones, *Fer1HCH*<sup>451</sup> and *Fer2LCH*<sup>35</sup> were recombined unto FRT82 containing chromosomes (Xu & Rubin 1993). *Fer1HCH*<sup>G188</sup> is a protein trap line and has been extensively described elsewhere (Missirlis et al. 2007). *Mvl*<sup>97f</sup> is an homozygous viable *P(lacW)* insertion was obtained from BDSC #5151 (Rodrigues et al. 1995).
Sec23^{13c8} mutant is a P(lacW) insertion within the 5'UTR of Sec23 (Abrams & Andrew 2005), BDSC stock #10218. Cg-Gal4 (BDSC stock #7011) was used to drive expression in the hemocytes,drm-Gal4 (BDSC stock #7098) in embryonic gut and scattered cells around the epidermis, FB-Gal4 in the fat bodies (Missirlis et al. 2006; Missirlis et al. 2003b). In cases where recombinant or double balanced stocks were needed they were generated following conventional crossing schemes.

Iron diets

Flies were raised for 3 successive generations on standard medium supplemented with 200 μM Bathophenanthrolinedisulfonic acid disodium salt (SIGMA #B1375) referred to as BPS in the text or with 1 mM ammonium iron (III) citrate (SIGMA #F5859) referred to as FAC. Adults were used for embryo collections. Protein extracts from female adults were also analyzed by non-reducing SDS-PAGE, confirming the differential accumulation of ferritin in flies raised on the respective diets (data not shown).

Immunohistochemistry and confocal imaging

Following dechorionation with a commercial bleach solution, embryos from overnight collections were devitellinized and fixed in a 1:1 mixture of heptane and 36% formaldehyde for 5 minutes and then washed in methanol. Embryos were then stored at -20°C or rehydrated, and used for staining. Primary antibodies used were: rat α-Elav 1:100, mouse α-BP102 1:100, mouse α-Eve 1:100 (Developmental Studies Hybridoma Bank); α-activated Caspase 3 1:100 (Cell Signaling); and rat α-Deadpan 1:2, a gift from Cheng-Yu Lee. Secondary antibodies used were: Alexa flour 546 α-rat 1:100 (Santa Cruz Biotechnology), Cy5 α-mouse 1:1000, Cy3 α-mouse 1:1000, FITC α-rabbit 1:1000 (Zymax). Signal from α-deadpan staining was increased with the ABC kit from Vectastain. A 510 Meta confocal microscope (Zeiss) was used for fluorescent imaging, and images were processed with Zeiss software and ImageJ.

Cuticle preparations and X-Gal staining

Embryos were collected in agar containing plates for 12 hours and incubated for another 36 hours at
25°C. Viable first instar larvae were removed from cultures. The cuticles of unhatched (dead) embryos were dechorionated and mounted in Hoyer's medium and incubated for 24 hours at 50°C to digest soft tissues. Resulting cuticles were then viewed and photographed with dark field optics in a compound microscope (Nikon). For X-Gal staining embryos were fixed and stained with X-Gal using standard procedures. Both controls and experimental embryos were incubated in parallel for the same amount of time to allow for direct comparisons.

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FIGURE LEGENDS

Figure 1. Ferritin mutants result in a variety of cuticle phenotypes quantified by different colors. Examples are shown of (A) wild type cuticle (green), (B) head involution defects (red), (C) dorsal closure defects (yellow), (D) germ band retraction defects (orange), (E) no cuticle deposition (blue). (F) Percentages of cuticular phenotypes of ferritin mutants. An enhancement of the earlier phenotypes was seen in mutant embryos whose mothers were fed BPS. 80% of embryos derived from ferritin mutant germline clones failed to develop cuticle. C: normal diet, FAC: high iron diet, BPS: low iron diet, GL: germline clones, n: number of cuticles examined.

Figure 2. Ferritin mutants result in CNS phenotypes revealed by α-Elav immunofluorescence. All embryos were oriented with anterior to the left and were visualized from a ventrolateral view. (A) control y, w embryo, (B-D) three examples of Fer1HCH451, (E, F) Fer1HCH35 and (G, H) Df(3R)Fer homozygous mutant embryos. The CNS appears twisted (C, E, H) and holes are seen within the ventral nerve cord (white arrows, B, D, F, H) in mutant embryos from all three genotypes.

Figure 3. In Df(3R)Fer homozygous mutant embryos the CNS appears contorted and with gaps in its organization. (A) Axons of the brain and ventral nerve cord have a stereotyped pattern in normal development. (B, C) Two different neuroblast populations, marked by Dpn and Eve, respectively, show a characteristic spatial organization in control y, w embryos. In homozygous Df(3R)Fer mutants, (D) the axons form but are disorganized and (E, F) the neuroblast populations are misplaced.

Figure 4. Fer1HCH451 lacZ enhancer trap is expressed in the embryonic CNS. Using an antibody against β-Galactosidase (green) and an antibody against the neuronal marker Elav (red), colocalization is observed in (A) heterozygous Fer1HCH451/+ and (B) homozygous Fer1HCH451 embryos.

Figure 5. Ferritin mutants cause apoptosis in the CNS and other tissues. Whole embryos were treated with an α-CSP3act marking apoptotic cells (green), and an α-Elav marking neurons (red). (A) Absence of staining for CSP3act in stage 12 y, w embryos, whereas (B) apoptotic markers appear in homozygous Df(3R)Fer embryos from stage 12 onwards mostly restricted to the neurogenic region. C) At stage 15 limited apoptotic events are seen in y, w embryos. D) At stage 15 homozygous Df(3R)Fer
embryos apoptosis can be observed in all embryonic tissues. (E) Higher magnification of a control ventral nerve cord and (F) a hole in the ventral nerve cord. DAPI was used to mark nuclei (blue), α-Elav for neurons (red) and α-CSP3act (green) for apoptosis.

**Figure 6.** Genetic interaction between ferritin and the DMT1 homolog Mvl. (A) The Mvl^{97f}-LacZ line shows a spatially restricted expression pattern for Mvl, mainly in head region, the brain, and a segmental repeated pattern. (B) In a Fer2LCH^{A17} mutant background, Mvl^{97f}-LacZ expression is induced. Black arrows denote the head region, white asterisk the embryonic brain, red asterisk the ventral nerve cord, and red arrows mark the segmented expression pattern. Cuticle preparations of (C) a homozygous Fer2LCH^{A17} embryo and (D) a double mutant Mvl^{97f}, Fer2LCH^{A17}. (E) Quantification of the appearance of melanotic spots in the cuticle following the introduction of one or two Mvl^{97f} alleles into a ferritin mutant background.

**Figure 7.** Ferritin accumulation in embryos with different genetic backgrounds. Ferritin protein was visualized in all embryos from the Fer1HCH^{G188} GFP trap line. In stage 17 heterozygous Fer1HCH^{G188/+} embryos that successfully complete development, ferritin mainly accumulates in the midgut and in hemocytes. (A) The GFP-Fer1HCH signal is found in hemocytes marked by cg>nRFP, but not (B) in the fat bodies marked by FB>nRFP. (C, D) Blocking the secretory pathway using a homozygous mutant Sec23^{13C8} background reveals that ferritin is absent from the hemocytes, where it normally resides, but accumulates in intestine. (E, F) Fer1HCH^{G188} homozygous embryos, which die like other ferritin mutants, also show ferritin accumulation in the intestine. These embryos also lose drm>nRFP staining suggesting intestinal disruption and lack ferritin accumulation in hemocytes suggesting ferritin trafficking defects.

**Figure S1.** Fer1HCH^{451} is a functional ferritin enhancer trap line. β-Galactosidase expression is normally restricted to the iron region in the larval midgut but if the expression is enhanced in the anterior midgut (AMG) when iron fed, as occurs in wild type larve (Mehta et al. 2009).
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cuticle preparation from ferritin mutants

Figure 1
Figure 2
Figure 3
Figure 4

(A) Fer1HCH^{451/+}

- LacZ
- α-Elav

(B) Fer1HCH^{451/451}

- LacZ
- α-Elav
Figure 5

A. S12
B. S15
C. y, w
D. Df(3R)Fer

E. α-Elav
F. DAPI
G. α-CSP3
H. merge

Df(3R)Fer
Figure 6

Panel A: Image of Mvl^{97f} embryos with LacZ expression. Panel B: Image of Mvl^{97f}, Fer2LCH^{Δ17} embryos with LacZ expression. Panel C: Image of Fer2LCH^{Δ17} embryos. Panel D: Image of Mvl^{97f}, Fer2LCH^{Δ17} embryos. Panel E: Bar graph showing the percentage of melanotic spot embryos for different genotypes: Mvl^{97f}, Fer2LCH^{Δ17}, Mvl^{97f/+}, Fer2LCH^{Δ17}, Mvl^{97f/97f}, Fer2LCH^{Δ17}. The graph shows a significant increase in melanotic spot embryos for the Mvl^{97f}, Fer2LCH^{Δ17} genotype compared to the others.
Figure 7