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# FAM210B is an erythropoietin target and regulates erythroid heme synthesis by controlling mitochondrial iron import and ferrochelatase activity

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Erythropoietin (EPO) signaling is critical to many processes essential to terminal erythropoiesis. Despite the centrality of iron metabolism to erythropoiesis, the mechanisms by which EPO regulates iron status are not well-understood. To this end, here we profiled gene expression in EPO-treated 32D pro-B cells and developing fetal liver erythroid cells to identify additional iron regulatory genes. We determined that FAM210B, a mitochondrial inner-membrane protein, is essential for hemoglobinization, proliferation, and enucleation during terminal erythroid maturation. *Fam210b* deficiency led to defects in mitochondrial iron uptake, heme synthesis, and iron-sulfur

cluster formation. These defects were corrected with a lipid-soluble, small-molecule iron transporter, hinokitiol, in *Fam210b*-deficient murine erythroid cells and zebrafish morphants. Genetic complementation experiments revealed that FAM210B is not a mitochondrial iron transporter but is required for adequate mitochondrial iron import to sustain heme synthesis and iron-sulfur cluster formation during erythroid differentiation. FAM210B was also required for maximal ferrochelatase activity in differentiating erythroid cells. We propose that FAM210B functions as an adaptor protein that facilitates the formation of an oligomeric mitochondrial iron transport complex, required for the increase in iron acquisition for heme synthesis during terminal erythropoiesis. Collectively, our results reveal a critical mechanism by which EPO signaling regulates terminal erythropoiesis and iron metabolism.

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The microarray data for this study were deposited to the NCBI Gene Expression Omnibus (GEO) database under GEO accession number GSE113657.

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This article contains Figs. S1–S3.

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## Fam210b regulates mitochondrial iron import and FECH activity

cellular processes, such as dopamine synthesis, oxygen transport, mitochondrial respiration, xenobiotic metabolism, and maintenance of the circadian rhythm (4–9). However, the majority of iron in the body is used by erythroid cells to synthesize hemoglobin (10, 11). Free iron is highly toxic because of its ability to generate free radicals through the Fenton reaction. Although iron is obtained from outside the cell and is required for reactions that take place in membrane-bound organelles, cellular membranes are not permeable to iron. Organisms have therefore evolved sophisticated mechanisms to efficiently transport iron across membranes (11, 12). In the majority of vertebrate cells, iron is mainly obtained through the transferrin-mediated pathway (13, 14), which is SNX3- and SEC15L1-dependent for efficient transferrin receptor recycling in erythroid cells (15, 16). In the plasma, each transferrin molecule binds two  $\text{Fe}^{3+}$  ions. Iron-bound transferrin interacts with transferrin receptor on the cell surface. This complex is internalized via clathrin-mediated endocytosis. Within the acidified endosome,  $\text{Fe}^{3+}$  is reduced to  $\text{Fe}^{2+}$  by STEAP proteins (17).  $\text{Fe}^{2+}$  is exported from the endosome by endosomal DMT1 (18–20) and may enter the mitochondrial intermembrane space via DMT1, which is localized on the mitochondrial outer membrane (21). Iron that is used for heme synthesis or iron-sulfur [Fe-S] clusters is imported into the mitochondrial matrix by the mitochondrial iron importer mitoferrins 1 and/or 2 (MFRN1/SLC25A37 or MFRN2/SLC25A28) (22–24). In developing red cells, the stabilization of MFRN1 by ABCB10 (25) is a mechanism by which the cell ensures efficient iron import for its utilization in hemoglobin synthesis (26). However, the  $\Delta mrs3\Delta mrs4$  yeast strain that is deficient in the mitoferrins only exhibited a partial defect in mitochondrial iron import. This suggests the existence of other mitochondrial iron importers (27, 28) or of auxiliary mechanisms that augment iron import to compensate for the genetic insufficiency of an iron transporter.

We identified potential EPO-regulated iron transport genes by microarray analysis of the EPO-treated, EpoR-expressing pro-B cell line, 32D, and we compared EPO-responsive genes to genes that were up-regulated during terminal erythropoiesis. One candidate gene, *Fam210b* (*C20orf120*), encoded a protein containing an N-terminal mitochondrial-targeting sequence. *Fam210b* was previously described to be a target of the erythroid transcription factor, GATA1, which encoded a mitochondrial-localized protein (29) that regulates mitochondrial metabolism in nonerythroid cells (30). In this study, we decided to interrogate the role of *Fam210b* in erythroid physiology. *Fam210b* expression is highly up-regulated during terminal erythropoiesis, which requires large quantities of iron. Loss-of-function studies in zebrafish embryos, primary murine fetal liver cells, and Friend murine erythroleukemia (MEL) cell lines show that FAM210B is required specifically to maintain the massive quantities of mitochondrial iron necessary for heme synthesis during terminal erythropoiesis. Although FAM210B is not an iron transporter *per se*, it interacts with terminal heme synthesis enzymes to form an oligomeric complex. Collectively, our data reveal that FAM210B plays a critical role as an adaptor protein in a mitochondrial iron import oligomeric complex. FAM210B is essential for integration of the rapid iron import

with a high rate of porphyrin synthesis that supports hemoglobinization during terminal erythropoiesis. This study describes a novel mechanism by EPO governs enucleation and cellular proliferation, processes underlying terminal erythropoiesis, by its control of mitochondrial iron uptake and heme biosynthesis.

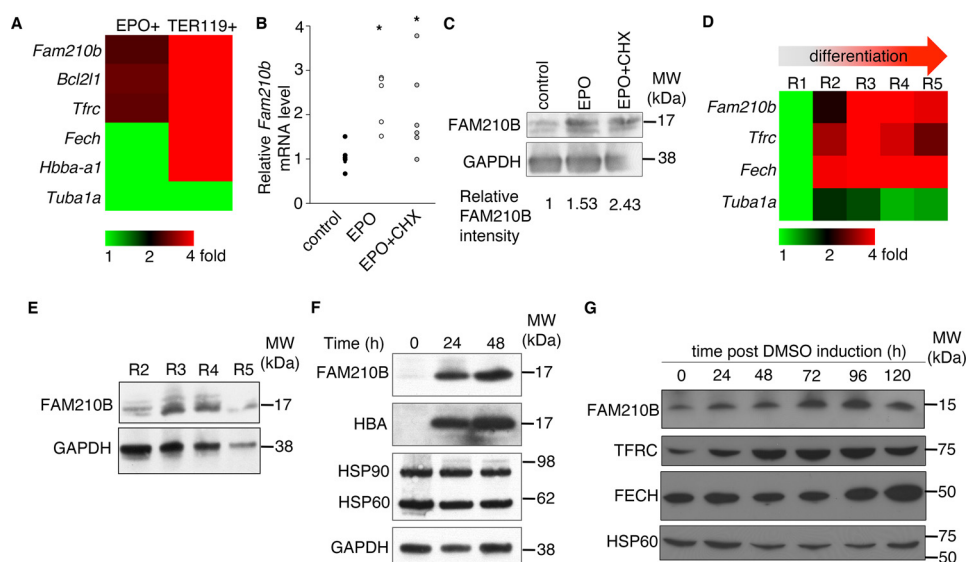
### Results

#### *Fam210b* is an immediate early target of EPO during terminal erythroid differentiation

Maturing erythroid cells acquire large quantities of exogenous iron to keep pace with the enormous demands of hemoglobin production (31, 32). To determine how EPO regulates iron transport, we identified genes up-regulated by EPO in the EpoR-expressing pro-B cell line, 32D (33). Cells were cultured in media without erythropoietin for 6 h prior to treatment with Epo (0.5 units/ml) for 2 h before harvesting. We analyzed global gene expression in control and EPO-treated murine 32D cells by microarray analysis. To determine which EPO targets could also be required for erythroid terminal differentiation, we compared the EPO-responsive genes in 32D cells with genes that were enriched in TER119<sup>+</sup> cells (34), which comprise the terminally differentiating erythroid cell population. We found that *Fam210b* expression was enriched in both datasets (Fig. 1A). To determine whether *Fam210b* was an EPO early response gene, a parallel experiment was set up in which 32D cells were treated with cycloheximide (10  $\mu\text{g}/\text{ml}$ ) 30 min prior to EPO stimulation to inhibit protein translation (35, 36). We observed that *Fam210b* expression was induced by EPO treatment, suggesting that *Fam210b* is an EPO target. The up-regulation of *Fam210b* expression persisted during cycloheximide treatment, indicating that *Fam210b* is an EPO early-response gene (Fig. 1B). The increase in *Fam210b* mRNA levels in response to EPO stimulation and during terminal differentiation translated to increases in the FAM210B protein. Interestingly, FAM210B protein was also up-regulated in the presence of cycloheximide, indicating that EPO increases the post-translational stability of FAM210B (Fig. 1C).

To determine how *Fam210b* is developmentally regulated, we performed RNA sequencing (RNAseq) on transcripts from murine fetal liver erythroid cells (34) that were sorted into fractions corresponding to their stage of maturation (R1–R5) (15, 37–39). *Fam210b* expression was up-regulated during maturation from R2 to R3, which corresponds to the developmental shift from TER119<sup>−</sup> to TER119<sup>+</sup> expression. This expression pattern was similar to that of genes involved in heme synthesis and iron import, such as *Fech* and *Tfrc* (Fig. 1D), and it is consistent with the enrichment of *Fam210b* in the TER119<sup>+</sup> population (Fig. 1A). FAM210B protein expression in the R2–R3 transition was recapitulated at the protein level (Fig. 1E). Consistent with this, FAM210B protein levels increase during *in vitro* differentiation of primary fetal liver erythroid cells (Fig. 1F) and in DMSO-induced terminal differentiation of MEL cells (Fig. 1G). The increases in protein expression are coordinated with  $\alpha$ -globin (HBA) (Fig. 1F), TFRC, and FECH (Fig. 1G), which are critical to erythroid differentiation, heme, and hemoglobin synthesis, suggesting that FAM210B plays an essential role in erythroid development. Although the

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**Figure 1.** *Fam210b* is an EPO early response gene and is induced in terminally differentiating erythroid cells. **A**, microarray analysis of EPO-treated 32D pro-B cells shows that *Fam210b* is an EPO-responsive gene that is also highly enriched in the terminally differentiating TER119<sup>+</sup> population of fetal liver erythroid cells (34). **B**, qRT-PCR demonstrates that Epo treatment of the EpoR-expressing pro-B cell line, 32D, up-regulated expression of *Fam210b* mRNA. This up-regulation persists in the presence of cycloheximide (CHX), an inhibitor of protein translation, demonstrating that *Fam210b* is an EPO early response gene  $n = 6$ ,  $*$ ,  $p < 0.05$ , Student's  $t$  test. **C**, FAM210B protein levels are up-regulated in response to EPO treatment of the EpoR-expressing 32D pro-B cell line. The increase in protein levels persist with cycloheximide treatment, indicating increased stability. Changes in FAM210B protein expression normalized to GAPDH are quantitated relative to control levels. **D**, RNAseq analysis of primary murine fetal liver cells sorted according to TER119 and CD71 (R1–R5) expression demonstrates up-regulation of *Fam210b* during the R2–R3 transition. **E**, this up-regulation is recapitulated by Western blot analysis of FAM210B protein expression. **F**, FAM210B protein expression is up-regulated during *in vitro* differentiation of primary fetal liver cells. **G**, FAM210B protein is induced upon terminal differentiation of MEL cells in parallel with genes required for heme synthesis, *TFRC* and *FECH*.  $*$ ,  $p < 0.05$ , Student  $t$  test.

expression of iron homeostasis genes is sometimes regulated by intracellular iron status via iron-regulatory proteins, IRP1 and IRP2 (13), we found that this was not the case for FAM210B (Fig. S1).

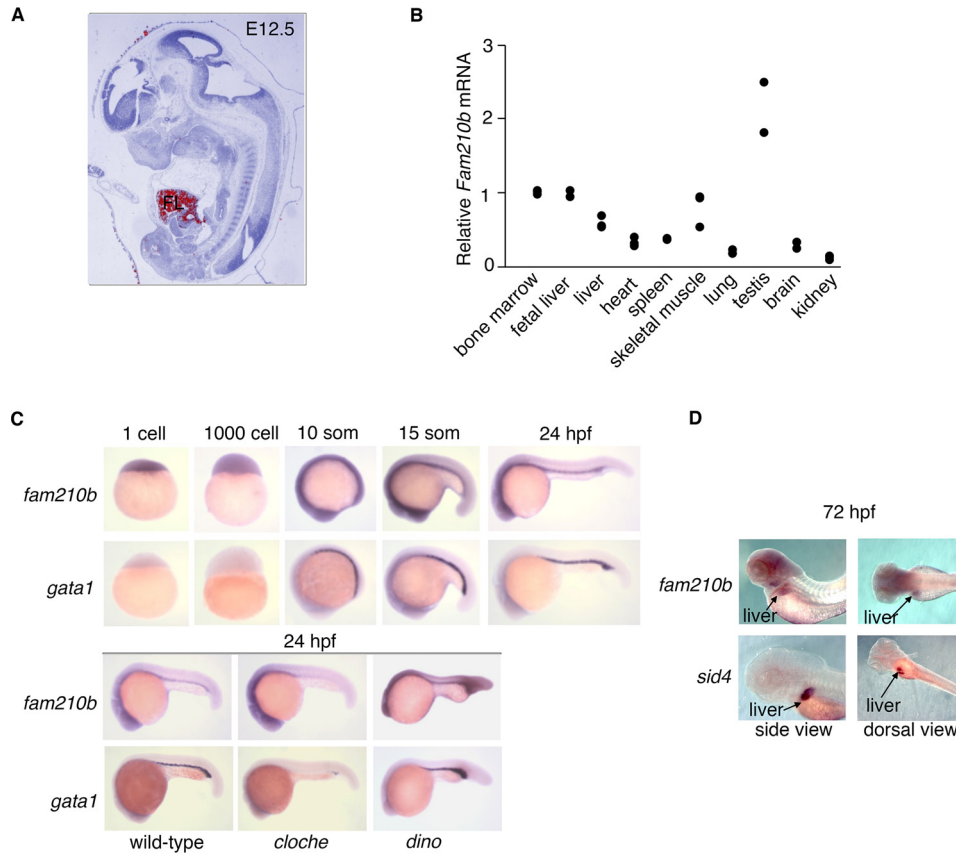
*In situ* hybridization of E12.5 murine embryos (performed as described (40)) indicated that *Fam210b* mRNA is enriched in the murine fetal liver at E12.5, the site of definitive erythropoiesis (Fig. 2A). In the adult mouse, *Fam210b* is enriched in the bone marrow, liver, and skeletal muscle, which are tissues that require large amounts of heme for synthesis of hemoproteins, and the testis, where iron plays an essential role in spermatogenesis (Fig. 2B) (41). In contrast to *gata1*, *fam210b* mRNA is maternally expressed in the developing zebrafish embryo. At 24 hpf, expression of *fam210b* mRNA is restricted to neuronal tissue and the intermediate cell mass (ICM), which is the site of primitive erythropoiesis in teleosts. *fam210b* is not expressed in *cloche* embryos (42, 43), which have defective blood and vascular progenitors. Conversely, *fam210b* expression is expanded in *dino* embryos. In *dino* embryos, the tail is expanded at the expense of the head (44). The tail is also the site of the ICM, which is the site of primitive hematopoiesis in the zebrafish. These zebrafish therefore have expanded blood due to their ventralized phenotype. The enrichment of *fam210b* in early hematopoietic tissues in zebrafish embryos, its absence in mutants that do not make hematopoietic progenitors, and the expansion of its expression in the *dino* mutants, which have an expanded ICM, indicate that *fam210b* expression depends on hematopoietic specification (Fig. 2C). At 72 hpf, *fam210b* expression persists in neural tissue and the liver (Fig. 2D), which are tissues that require large amounts of iron for their function (8, 13).

### *Fam210b* is required for erythropoiesis *in vivo*

To determine the biological function of *fam210b*, we injected antisense morpholinos to knock down its expression in developing zebrafish embryos. As *fam210b* is highly expressed in terminally differentiating erythroid cells (Fig. 1, A and B), we assayed heme synthesis in morphant embryos by *o*-dianisidine staining as a measure of terminal erythropoiesis. Compared with the controls, *fam210b* morphants have significantly decreased heme staining, indicative of their anemia (Fig. 3A). We quantitated the erythropoietic defect by flow cytometry analysis of GFP<sup>+</sup> cells in morphant *Tg(globin-LCR:eGFP)* transgenic zebrafish, which express GFP under the control of a globin locus control region (LCR) enhancer, restricting GFP expression to the erythroid lineage (45). Morphant embryos exhibited decreased erythropoiesis at 24- and 48-hpf, highlighting the importance of *fam210b* in erythropoiesis (Fig. 3B).

To determine whether *Fam210b* was required for mammalian erythropoiesis, we knocked down *Fam210b* in primary fetal liver erythroid cells with shRNA (Fig. 3C). *Fam210b* knockdown caused a reduction in hemoglobinization (Fig. 3D), which was partially ameliorated by co-expression of human FAM210B (Fig. 3E), indicating the functional orthology between the human and mouse FAM210B genes. Knockdown of FAM210B caused defects both in cell proliferation (Fig. 3F) and in enucleation (Fig. 3G), indicating its additional function in erythroid maturation. The independent validations of the erythroid defect in *Fam210b* knockdown zebrafish and murine models indicate that the role of FAM210B in terminal erythroid maturation is highly conserved in vertebrates. In higher vertebrates, continuous FAM210B expression in the erythron is required

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**Figure 2. Expression of *Fam210b* is enriched in vertebrate tissues with high heme content.** *A*, *Fam210b* expression (red pseudocolor) is enriched in the murine fetal liver (FL) at E12.5. *B*, in murine tissues, *Fam210b* expression is enriched in the bone marrow and fetal liver (erythroid), adult liver, skeletal muscle, and testis, all of which require high levels of iron for their function. *C*, zebrafish *fam210b* is maternally expressed during early embryonic development. At 24 hpf, *fam210b* expression is enriched in neural tissue and the intermediate cell mass, the site of primitive hematopoiesis (top). The expression of *fam210b* in the ICM is abolished in *cloche* embryos, which do not form hematopoietic or vascular tissue. In contrast, *fam210b* expression in the intermediate cell mass is expanded in *dino* embryos, which exhibit a ventralized phenotype (bottom), paralleling the increased expression of erythroid *gata1*. *D*, at 72 hpf, *fam210b* expression persists in the neural tissue and is enriched in the liver, delineated by *sid4* expression. *n* = 3.

for optimal erythroid cell proliferation, maturation, hemoglobinization, and enucleation.

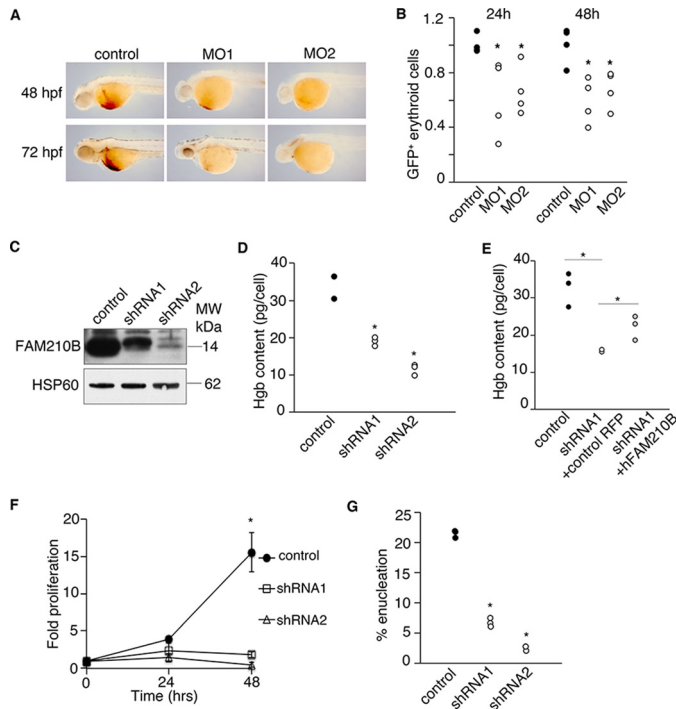
### *Fam210b* is localized to the mitochondrial inner membrane

To delineate the cellular mechanism by which FAM210B regulates erythropoiesis, we transfected *fam210b*-GFP into HEK293T cells and examined its subcellular localization. Fam210b contains three hydrophobic stretches of ~21 amino acids, suggesting that FAM210B is an integral membrane protein. Confocal immunofluorescence microscopy revealed that FAM210B-GFP co-localized with COX4, a mitochondrial resident protein (Fig. 4A). Subcellular fractionation and Western blot analysis of primary fetal liver cells confirmed that FAM210B specifically co-localized with HSP60, a mitochondrial resident protein (Fig. 4B).

We carried out further localization studies in MEL cells stably expressing *Fam210b* to determine the sub-mitochondrial localization of FAM210B. To determine the localization of FAM210B in the mitochondrial outer membrane, purified mitochondria were treated with proteinase K, followed by centrifugation to separate the intact mitochondria (P) from soluble proteins (S) (Fig. 4C). Immunoblotting showed that TOMM20, an outer membrane protein with a large cytosolic domain, was degraded upon protease treatment, whereas FAM210B and

control proteins YME1 (intermembrane space) and PreP (matrix) were intact. FAM210B is therefore not an outer mitochondrial membrane protein.

The mitochondria were then subjected to osmotic shock to determine whether FAM210B was in the intermembrane space or matrix (Fig. 4C). Mitoplasts, which contain the inner membrane and matrix components and intermembrane space/outer membrane components that stick to the mitoplasts, were recovered by centrifugation (P), whereas soluble proteins (S) remained in the supernatant. The mitoplasts were subsequently treated with a low dose of proteinase K to degrade proteins that were in the intermembrane space. FAM210B and PreP were resistant to protease digestion, whereas intermembrane space protein YME1 was degraded as expected. As a control for proteinase K activity, the mitoplasts were treated with Triton X-100, thereby releasing membrane-bound proteins into the supernatant and exposing digestion sites to enzymatic digestion. The addition of proteinase K to the Triton X-100 detergent-treated samples resulted in complete degradation, with the exception of PreP that has a tightly folded domain that is resistant to the low doses of proteinase K. These data suggested that FAM210B was bound to the mitochondrial inner membrane.

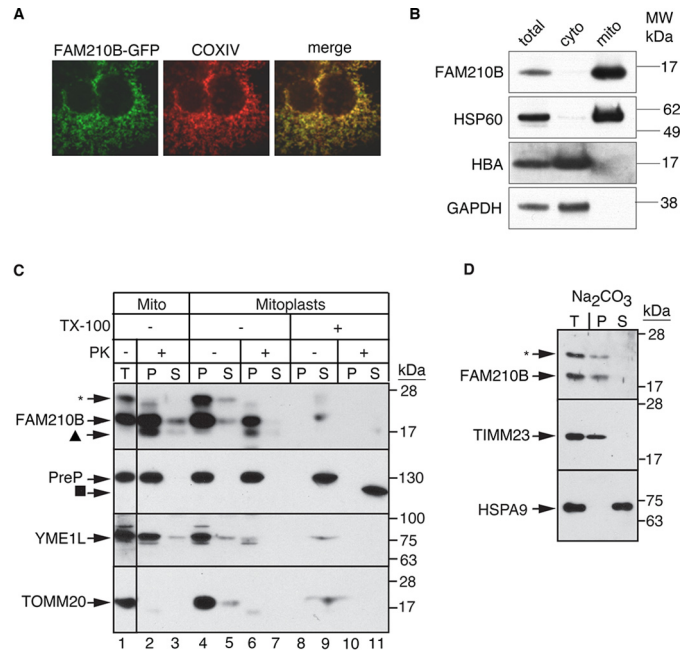


**Figure 3. *Fam210b* is continuously required for terminal erythroid differentiation.** *A*, knockdown of *fam210b* in zebrafish embryos with two independent antisense morpholinos caused a defect in erythroid hemoglobinization. *B*, *Tg(globin LCR:eGFP)*, *fam210b* morphant zebrafish exhibited a decrease in GFP<sup>+</sup> erythroid cells, indicating defective erythropoiesis,  $n = 4$ ; \*,  $p$  value < 0.05 Student's  $t$  test. *C*, shRNA-mediated knockdown of *Fam210b* in primary fetal liver cells decreased FAM210B protein expression. *D*, *Fam210b* knockdown primary fetal liver cells had a decrease in hemoglobin content. *E*, hemoglobinization defect in murine *Fam210b* knockdown cells is complemented by human *FAM210B*. *F*, *Fam210b* knockdown in primary fetal liver cells caused a cell proliferation defect. *G*, *Fam210b* knockdown caused an enucleation defect, indicative of erythroid maturation arrest.  $n = 3$ , \*,  $p$  value < 0.05 Student's  $t$  test.

To rigorously confirm that this was the case, mitochondria were subjected to sodium carbonate extraction (46), followed by centrifugation, to separate membrane proteins (pellet, P) from soluble (S) proteins (Fig. 4D). FAM210B and membrane-bound TIMM23 were recovered in the pellet, whereas soluble matrix-localized HSPA9 (mortalin) was released in the supernatant. Thus, FAM210B is an integral membrane protein residing in the inner membrane. The collective data indicate that FAM210B is integrated in the inner membrane, and any soluble domains are likely facing the matrix, because protease addition during osmotic shock did not markedly degrade the full-length protein. A smaller fragment, which was sub-stoichiometric, was detected in the pellet (Fig. 4C, marked with a triangle), which may indicate that the N or C terminus is facing the intermembrane space and is digested by protease that may have leaked into the intermembrane space. The smaller fragment sediments with the pellet, suggesting that the segment of FAM210B that is exposed to the protease is tightly associated with the membrane.

**FAM210B facilitates mitochondrial iron import during terminal erythropoiesis**

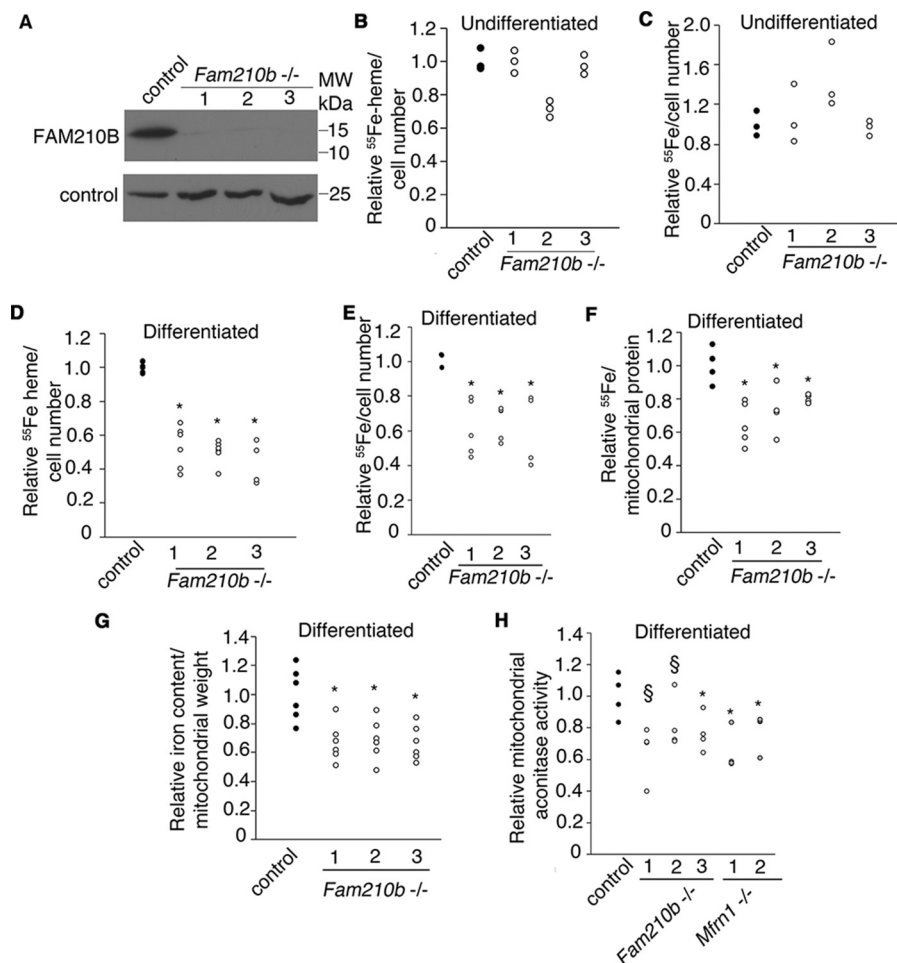
To determine the biochemical function of FAM210B in heme synthesis, we generated *Fam210b*-deficient MEL cell



**Figure 4. FAM210B localizes to the mitochondrial inner membrane.** *A*, confocal fluorescence microscopy of exogenously expressed FAM210B-GFP (green) in HEK293T cells indicated that FAM210B co-localized with COXIV (red), a mitochondrial resident protein. The overlap is indicated in the merge panel (yellow). *B*, subcellular fractionation of primary fetal liver cells showed that FAM210B co-sedimented with HSP60 (mitochondrial) but not with HBA and GAPDH (cytoplasmic), indicating its mitochondrial localization. *C*, isolated mitochondria (Mito, T; lanes 1–3) purified from a MEL cell line stably expressing mouse *Fam210b*, was treated with proteinase K, which degraded outer membrane proteins (TOMM20) but not intermembrane space proteins (YME1L) or matrix (PreP) proteins (lane 2). Inner mitochondrial proteins, which were impervious to proteinase K treatment, (P) were separated from S proteins by centrifugation. FAM210B co-sedimented with the pellet fraction (lane 2). Mitoplasts (lanes 4–11) were generated by subjecting intact mitochondria to osmotic shock, exposing inner membrane and intermembrane space proteins to proteinase K digestion. Mitoplasts were centrifuged to separate the soluble proteins from the mitoplast pellet. Proteinase K treatment degraded a fraction of FAM210B (lane 6) and completely degraded YME1L but not PreP, suggesting that at least a fraction of FAM210B was situated in the mitochondrial inner membrane or intermembrane space. Triton X-100 (TX-100) treatment liberated FAM210B, PreP, and YME1L from the pellet into the soluble fraction (lane 9). These solubilized proteins were digested by proteinase K, demonstrating specificity of assay (lane 10). The asterisk marks a non-specific band that was detected with the mouse FAM210B antibody; the triangle marks a likely FAM210B degradation product; and the square marks a core of PreP that is tightly folded and resistant to protease treatment. *D*, to determine whether FAM210B was an integral inner-membrane protein, isolated mitochondria (T) were analyzed by carbonate extraction. P and S fractions were analyzed by immunoblotting for FAM210B. The blot was probed for TIMM23 as an integral membrane protein control and mortalin as a soluble protein control. An asterisk marks nonspecific bands detected by the FAM210B antibody.

lines by CRISPR/Cas9 genome editing (Fig. 5A). Metabolic <sup>55</sup>Fe-labeling revealed that undifferentiated *Fam210b*<sup>-/-</sup> MEL cells had normal rates of heme synthesis and iron uptake compared with WT controls (Fig. 5, B and C). However, when MEL cells were chemically induced to differentiate, heme synthesis and iron uptake in *Fam210b*<sup>-/-</sup> MEL cells was significantly impaired (Fig. 5, D and E). The heme synthesis defect in *Fam210b*<sup>-/-</sup> MEL cells was due to a deficiency in mitochondrial iron acquisition (Fig. 5, F and G). As FAM210B is required for optimal mitochondrial iron uptake, we assayed the activity of mitochondrial aconitase (EC 4.2.1.3) in FAM210B<sup>-/-</sup> cells. Mitochondrial aconitase activity is sensitive to mitochondrial iron deficiency (47) caused by defects in mitochondrial iron

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**Figure 5. Fam210b is required for heme synthesis by facilitating mitochondrial iron import.** *A*, *Fam210b*<sup>-/-</sup> MEL cell lines generated by CRISPR/Cas9 expressed no detectable FAM210B protein. *B*, <sup>55</sup>Fe metabolic labeling showed that undifferentiated *Fam210b*<sup>-/-</sup> MEL cell clones did not cause changes in heme synthesis ( $n = 3$ ) or iron uptake ( $n = 3$ ) (*C*). *D*, differentiated *Fam210b* knockout MEL cells exhibited defective heme synthesis ( $n = 6$ ). *E*, iron uptake ( $n = 5$ ). *F*, mitochondria from differentiated *Fam210b* knockout cells had decreased <sup>55</sup>Fe-transferrin uptake ( $n = 5$ ). *G*, ICP-MS demonstrated that mitochondria from *Fam210b*<sup>-/-</sup> cells had a decrease in total iron ( $n = 6$ ). *H*, mitochondrial aconitase activity, which depends on [Fe-S] cluster synthesis, is decreased in *Fam210b*<sup>-/-</sup> cells indicating a defect in mitochondrial iron acquisition. *Mfrn1*<sup>-/-</sup> MEL clones served as controls for mitochondrial iron deficiency ( $n = 4$ ). \*,  $p < 0.05$ ; \$,  $p < 0.1$  Student's *t* test. All data are normalized to WT controls.

transport, and this is reflected by the aconitase defect in *Mfrn1*<sup>-/-</sup> MEL cells (Fig. 5H) (23, 28). *Fam210b*<sup>-/-</sup> cells had a decrease in mitochondrial aconitase activity that is comparable with *Mfrn1*<sup>-/-</sup> cells, indicating that it plays a critical role in mitochondrial iron metabolism.

The heme synthesis (Fig. 6A) and iron transport (Fig. 6B) defects in *Fam210b*<sup>-/-</sup> MEL cells were functionally complemented to the level of WT cells by hinokitiol (CAS 499-44-5; ChEBI: 10447), a natural product that binds iron and autonomously transports both Fe(II) and Fe(III) across lipid membranes to bypass defects in iron transport (48). Iron-hinokitiol functionally complemented the erythropoietic defect in *fam210b* morphant zebrafish embryos (Fig. 6C), demonstrating that the primary role of FAM210B is to facilitate mitochondrial delivery of iron for the synthesis of heme and [Fe-S] clusters in the developing erythroid cell. C2-deoxyhinokitiol, which does not bind or transport iron, did not chemically complement the anemia in *Fam210b*<sup>-/-</sup> cells (Fig. 6, A and B), indicating that the ability of the iron-hinokitiol to rescue heme synthesis was directly dependent on the ability of hinokitiol to chelate and transport iron across membranes (48). The specificity of the

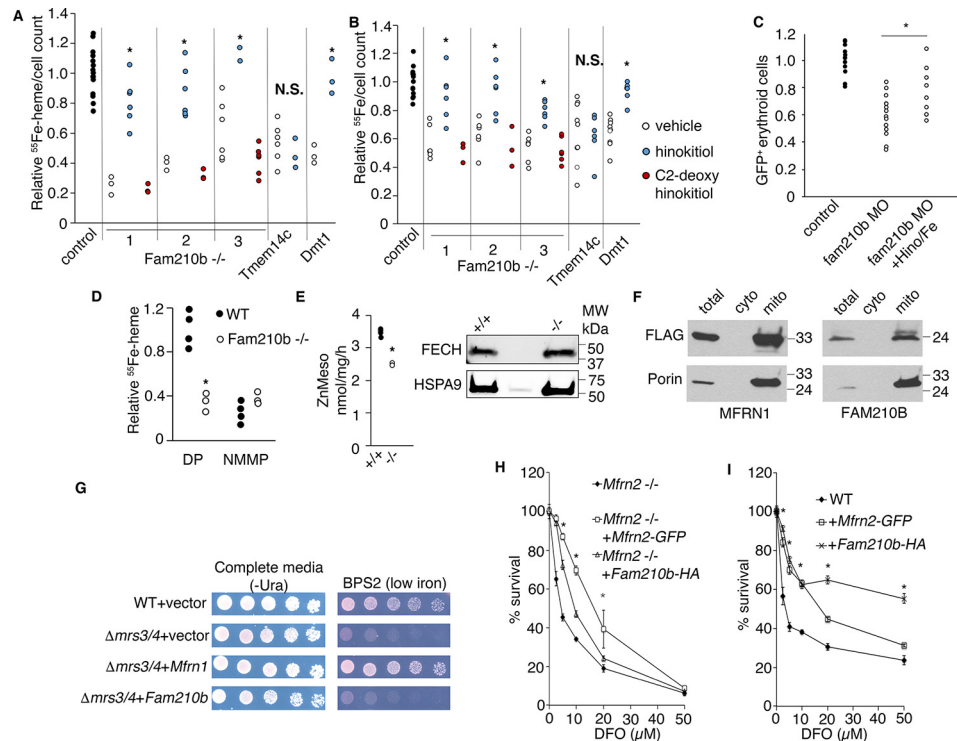
functional complementation is further demonstrated by the restoration of both <sup>55</sup>Fe and <sup>55</sup>Fe-heme only in cells defective for iron transport, such as the *Dmt1* knockdown cell line (19, 20), but not in cells with a porphyrin transport defect, such as the *Tmem14c* CRISPR cell line (38).

HPLC analysis of protoporphyrinogen IX (PPIX) content in differentiating *Fam210b*<sup>-/-</sup> cells indicated that FAM210B was not required for synthesis of the tetrapyrrole ring in the mitochondria (Fig. S2A). This was further confirmed by the inability of deuteroporphyrin IX (DP), a PPIX analog that can chemically complement porphyrin synthesis defects (38), to rescue heme synthesis in *Fam210b*<sup>-/-</sup> cells (Fig. S2B).

To exclude the possibility that the heme defect in FAM210B deficiency was caused by a lesion in mitochondrial biogenesis or physiology, we stained control and *Fam210b*<sup>-/-</sup> cells with Mitotracker dye and analyzed mitochondrial staining by flow cytometry. The percentage of Mitotracker-positive cells and average Mitotracker dye staining did not differ between control and *Fam210b*<sup>-/-</sup> cells, indicating that *Fam210b* deficiency did not cause defects in cell survival (Fig. S2C), mitochondrial membrane potential, and mitochondrial mass (Fig. S2D).



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**Figure 6. Fam210B increases the kinetics of mitochondrial iron import in differentiating MEL cells by functioning as an auxiliary factor.** *A*,  $^{55}\text{Fe}$  metabolic labeling confirmed a decrease in heme synthesis in differentiating *Fam210b* (CRISPR1–3), *Tmem14c*, and *Dmt1* deficient MEL cells (black bars). The addition of hinokitiol, a lipid-soluble iron carrier (white bars), restored heme synthesis in *Fam210b*- and *Dmt1*-deficient cells, but not *Tmem14c*-deficient cells. C2-deoxyhinokitiol, which does not chelate iron, did not complement heme synthesis in the *Fam210b* knockout cells (gray bars) ( $n = 6$ ). *B*,  $^{55}\text{Fe}$  metabolic labeling confirmed an iron uptake defect in differentiating *Fam210b*-, *Tmem14c*-, and *Dmt1*-deficient cells (black bars). The iron uptake defect in *Fam210b*- and *Dmt1*-deficient cells was chemically complemented by hinokitiol (white bars). Hinokitiol did not rescue iron uptake in *Tmem14c*-deficient cells, which have a porphyrin synthesis defect. C2-deoxyhinokitiol did not rescue the iron deficiency in *Fam210b* $^{-/-}$  cells (gray bars) ( $n = 6$ ). *C*, treatment of *fam210b* morphant zebrafish embryos (MO) with iron citrate and hinokitiol (MO + Hino/Fe) rescued their anemia ( $n = 8$ ). *D*, FECH activity was measured in intact mitochondria isolated from WT and *Fam210b* $^{-/-}$  MEL cells, using  $^{55}\text{Fe}$  and DP as substrates. FECH activity in *Fam210b* $^{-/-}$  mitochondria was  $\sim 1/3$  that of WT controls. Both WT and *Fam210b* $^{-/-}$  mitochondria had very little measurable FECH activity when NMMP was used as a substrate ( $n = 4$ ). *E*, FECH activity was measured in isolated mitochondria from WT and *Fam210b* $^{-/-}$  MEL cells that were treated with detergent and disrupted by sonication, allowing access of reaction substrates to FECH in the absence of intact membranes. FECH activity was decreased in *Fam210b* $^{-/-}$  mitochondria, but not to the extent of intact mitochondria ( $n = 3$ ). *F*, FLAG-tagged MFRN1 and FAM210B co-localized with porin in  $\Delta mrs3/4$  yeast, indicating correct mitochondrial localization. *G*, *Mfrn1* expression complemented the growth defect of  $\Delta mrs3/4$  yeast in low-iron media, whereas *Fam210b* expression did not. *H*, survival of *Mfrn2* $^{-/-}$  fibroblasts in DFO is significantly increased by expression of *Mfrn2-GFP*, but less so by expression of *Fam210b-HA* ( $n = 3$ ). *I*, expression of *Fam210b-HA* and *Mfrn2-GFP* in WT fibroblasts increase cell survival in the presence of DFO, with *Fam210b* overexpression possessing a protective effect over a larger dose range than *Mfrn2-GFP*. Mean  $\pm$  S.E.,  $n = 3$ , \*,  $p < 0.05$ ; §,  $p < 0.1$  Student's *t* test. N.S., not significant.

### FAM210b is not a direct mitochondrial iron carrier in vertebrate erythroblasts

To determine whether FAM210B functions as a mitochondrial iron importer or whether it played a role in regulation of FECH, we carried out an *in vitro* heme synthesis assay to compare the amount of iron transported across the mitochondrial membrane to synthesize deuteroheme (49).  $^{55}\text{Fe}$  and DP were added to isolated mitochondria from WT and *Fam210b* $^{-/-}$  MEL cells. DP was added in excess. As DP is lipid-soluble, any differences in the synthesis of  $^{55}\text{Fe}$ -DP are caused by alterations in FECH activity or iron transport across the mitochondrial membrane. In the absence of FAM210B, *in vitro* heme synthesis decreased to background levels comparable with cells treated with NMMP, a PPIX analog that inhibits FECH and cannot be metallated (Fig. 6D) (50). These data suggested that FAM210B is required for either iron import or heme synthesis. To determine which was the case, we performed a FECH activity assay as described (81). Using this method, cells are disrupted by sonication and treatment with Tween 20 detergent, allowing the substrates, mesoporphyrin IX, and zinc access to FECH with-

out the need to cross membranes. FECH activity is measured by quantification of the product zinc-mesoporphyrin IX by HPLC. In the absence of a mitochondrial membrane, we observed that *Fam210b* $^{-/-}$  cells exhibited a decrease in FECH activity (Fig. 6E, left). This was not attributable to a decrease in FECH protein levels (Fig. 6E, right). Based on Fig. 6, D and E, FAM210B is required for optimal FECH activity in erythroid cells. However, we observed that the decrease in FECH activity was far less significant when membranes were disrupted (Fig. 6E) compared with when the mitochondrial membrane was intact (Fig. 6D). We concluded that FAM210B is required both for full activation of FECH but also plays a role in facilitating iron transport across the mitochondrial membrane for heme synthesis.

To determine whether FAM210B is a *bona fide* iron transporter, we expressed epitope-tagged *Fam210* or *Mfrn* in the  $\Delta mrs3\Delta mrs4$  yeast strain (denoted as  $\Delta mrs3/4$ ), deficient in the *Mfrn* orthologs (23), or in *Mfrn2* $^{-/-}$  primary murine fibroblasts,<sup>7</sup> which are deficient in mitochondrial iron transporter

<sup>7</sup> D. M. Ward and J. Kaplan, unpublished data.



and FECH or potential bridging proteins that can mediate indirect interactions, it is likely that the physical interactions between FAM210B, PPOX, and FECH are direct. Notably, PPOX and FECH are in the same complex as MFRN1 (Chen *et al.* (26) and Medlock *et al.* (79)). The specificity of these protein–protein interactions is ensured by the lack of CPOX co-immunoprecipitation with FAM210B (Fig. 7B). To further determine whether FAM210B can interact with FECH at endogenous levels, we generated a MEL cell line that stably expressed FAM210B–FLAG. We immunoprecipitated FAM210B with FLAG antibodies in differentiating cells. Western blot analysis of the immunoprecipitated protein showed that FAM210B–FLAG interacted with endogenous FECH *in vivo* (Fig. 7C). FAM210B does not homooligomerize, as would be expected if FAM210B was a membrane-embedded solute carrier (Fig. 7C). Collectively, these results are consistent with a role for FAM210B as an auxiliary factor in an iron transport oligomeric complex that includes the terminal heme biosynthesis enzymes in the mitochondria. The intact oligomeric complex is essential for maintaining the high rate of mitochondrial iron import that is required to fulfill the demands for heme synthesis in maturing red cells.

## Discussion

Our study demonstrates that FAM210B, a protein of previously unknown function, is required for mitochondrial iron import to support sustained synthesis of heme, iron–sulfur cluster, and hemoglobin during erythroid maturation. In addition, FAM210B regulates FECH enzyme activity. Our results are consistent with and extend the findings of a recent study identifying *Fam210b* as a GATA1 transcriptional target that is required for erythroid differentiation (29). Mitochondrial mass, membrane potential, and porphyrin levels are normal in *Fam210b*<sup>−/−</sup> cells, precluding a role for FAM210B in housekeeping mitochondrial homeostasis or tetrapyrrole synthesis, respectively. Although heme synthesis and iron import was decreased in *Fam210b*-deficient cells, we did not observe an accumulation in porphyrin, similar to the phenotype in *Mfrn1* deficiency (51, 52). This is attributable to impaired [Fe-S] cluster biosynthesis in *Fam210b*-deficient cells (Fig. 5H), which elevates IRP1 RNA binding and suppresses *Alas2* mRNA translation and porphyrin accumulation (52, 53).

FAM210B deficiency can be functionally complemented by the addition of hinokitiol, which acts as a small-molecule mimetic of iron transporters (48). However, *Fam210b* overexpression cannot genetically complement the iron transport defect in  $\Delta$ *mrs3/4* yeast or *Mfrn2*<sup>−/−</sup> fibroblasts. We therefore conclude that the primary function of FAM210B is to facilitate mitochondrial iron import in differentiating erythroid cells by functioning as an auxiliary factor in promoting formation of an oligomeric iron–transport complex with terminal heme synthesis enzymes, thereby increasing the kinetics of mitochondrial iron import to keep pace with its cellular demands for heme. Our studies on FAM210B are the first to describe an EPO-induced mitochondrial chaperone that is tissue- and differentiation stage-specific, revealing a novel mechanism by which cells regulate intracellular iron metabolism. We previously identified a mitochondrial chaperone, CLPX, which facilitates the delivery of the vitamin B<sub>6</sub> cofactor required for the

catalytic activity of ALAS, the enzyme that catalyzes the committed step of protoporphyrin synthesis (54). The oligomerization of MFRN1, ABCB10, and FECH integrates the efficient import of iron with the metallation of PPIX to form heme during erythroid maturation (26). We further note that the use of small molecules that autonomously perform protein-like functions as a distinct type of biological probe, as demonstrated herein with the transmembrane iron transporter hinokitiol, may be more widely applicable and have complementary strengths to other experimental approaches (48, 55).

*Fam210b* is transcriptionally activated by GATA-1 (29, 56), consistent with its expression in erythropoietic tissues. Iron chelation by DFO or iron supplementation by iron citrate did not cause changes in its steady-state protein expression, indicating that FAM210B is not regulated by cellular iron status (Fig. S1).

EPO regulates FAM210B expression both at the levels of mRNA transcription and protein stability, suggesting that FAM210B is an essential component of EPO's function and signaling in erythropoiesis. The up-regulation of *Fam210b* by EPO in EpoR-expressing 32D pro-B cells suggests that *Fam210b* is direct target of EPO. Although FAM210B is localized in the mitochondria, it is required for cellular processes that take place during terminal erythroid differentiation, such as proliferation, and finally enucleation. These data suggest a link between nuclear processes in terminal erythroid differentiation and mitochondrial iron metabolism and also demonstrate a novel mechanism by which EPO regulates the biology of terminal erythropoiesis via its regulation of cellular iron uptake.

Although published GWAS studies did not reveal any extant association between *FAM210B* mutations and hematologic traits (57–61), the profoundly anemic phenotype of the *fam210b* morphant embryos and the heme defects in primary murine fetal liver and in MEL cells predict that mutations in *FAM210B* may lead to hematologic and iron metabolism defects in humans. As *MFRN1* genetic or splicing defects have been shown to be modifiers of porphyria (62) or sideroblastic anemia (63), we predict that detailed genetic studies will reveal the function of *FAM210B* mutations in patients suffering from anemias and porphyrias of unknown etiology. As *Fam210b*<sup>−/−</sup> cells only exhibited heme and iron metabolism defects in differentiating erythroid cells, it is expected that *FAM210B* mutations will exhibit a phenotype specifically in the terminally differentiating erythroid cell population.

To adapt to their role as oxygen carriers in vertebrates, erythroid cells have evolved several mechanisms to efficiently transport and utilize iron within the cell for adequate heme synthesis. This is critical as mitochondrial iron import is tightly coupled with heme synthesis, and cells cannot preload iron in the mitochondria (64). In addition to the housekeeping mitochondrial iron importer MFRN2, erythroid cells express MFRN1 during their terminal maturation (22, 23). The tissue-restricted regulation at the transcriptional level for *MFRN1* at “erythroid super-enhancer” by GATA-factors (65, 66), is reminiscent of the transcriptional regulation of ALAS, the first and rate-limiting enzyme in the heme biosynthetic pathway (67). Like the *MFRN* genes, these *ALAS* genes are encoded by two separate genes, encoding distinct ALAS isoenzymes. The ubiqu-

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uitous *ALAS1* is regulated by heme via the peroxisome proliferator-activated 1 $\alpha$  (PGC-1 $\alpha$ ) (68). The erythroid-enriched *ALAS2* and *MFRN1* genes are not regulated by heme. The post-translational regulation of *ALAS2* is mediated by the intracellular levels of iron and iron-responsive protein 1 (IRP1) binding to its cognate iron-responsive element (IRE) in the 5'-untranslated region (UTR) of the *ALAS2* mRNA (69); however, no such regulation by iron on IRE have been reported and ascribed to the *MFRN* genes.

During terminal erythropoiesis, *MFRN1* protein stability is also enhanced by oligomerization with *ABCB10* (25, 26). *MFRN1* and *ABCB10* are in a physical complex with *FECH*, which kinetically favors the coupling of iron transport and heme synthesis. The addition of *FAM210B*, which forms an oligomer with *PPOX* and *FECH* to enhance mitochondrial iron import and heme synthesis, fits into this experimental model where *FAM210B* can nucleate or stabilize the formation of a complex between *PPOX* and *FECH*. In this manner, it can facilitate the transfer of *PPIX* to *FECH* and promote iron import coupled with heme synthesis (Fig. 7D). While *PPOX* and *FECH* physically interact, they do not form a tight complex. This supports the idea that bridging molecules, such as *FAM210B*, may mediate the strength of the interaction, hence modulating the rate of iron import and heme synthesis (70).

*FAM210B* plays a second role in heme synthesis by activating *FECH* enzymatic activity (Fig. 6, D and E). It is well-established that red cells increase the expression of ferrochelatase protein during erythropoiesis. This occurs either by increase in *FECH* mRNA transcription (71) or stabilization of *FECH* protein by binding of iron-sulfur clusters (72). In contrast, *FAM210B* deficiency does not alter *FECH* protein levels (Fig. 6E), suggesting that its effects on *FECH* activity may be mediated by the requirement for *FAM210B* for efficient iron delivery to *FECH*, and/or by playing a role in allosteric activation of the *FECH* enzyme. These data add to our understanding of the mechanisms by which Epo signaling regulates heme synthesis by post-translational regulation of *FECH* (49) and suggest mechanisms by which the activity of heme synthetic enzymes, in this case, *FECH*, may be rapidly modulated in response to metabolic requirements.

As *Fam210b* is also expressed in nondifferentiating erythroid cells and neural and hepatic tissues, it may also play a role in these tissues that is independent of heme synthesis and iron metabolism. Because *FAM210B* is a mitochondrial protein, detailed sequencing studies may reveal mutations in *FAM210B* that play a role in modifying the severity of mitochondriopathy in these tissues. Our identification of *Fam210b* as a novel mitochondrial protein that is required for optimal mitochondrial iron uptake during terminal erythropoiesis thus provides a novel genetic tool for further studies on vertebrate erythropoiesis and mitochondrial biology.

## Materials and methods

### Cell lines

MEL DS19 subclone cells were obtained from Arthur Skoultschi (Albert Einstein College of Medicine, New York). Human HEK293T cells were used for transient transfections.

### GenBank™ accession numbers

Zebrafish *fam210b* (*c20orf108*) is GenBank™ accession number NM\_001033751; mouse *Fam210b* is GenBank™ accession number NM\_025912.

### Knockdown of *Dmt1* and *Fam210b* by shRNA hairpins in mouse cells

*Dmt1* (GenBank™ accession number NM\_008732.2) stable knockdown MEL clones were obtained by stable transfection of an shRNA hairpin, TRCN0000079535. *Fam210b* (GenBank™ accession number NM\_025912) stable knockdown MEL clones were obtained by transfection of shRNA hairpins: shFam210b-1 and TRCN0000103732; shFam210b-2 TRCN0000103734. shRNA hairpins were obtained from Sigma. MEL clones were electroporated with shRNA plasmids by electroporation, and stable clones were obtained by limiting dilutions and selection with G418 (*Dmt1*).

### CRISPR design and cloning

CRISPR guide sequences were designed to direct cleavages at genetic loci to generate chromosomal deletions (73, 74). CRISPR guide sequences were designed to have a unique 12-bp seed sequence 5'-NNNNNNNNNNNNNGG-3' in the mouse genome to minimize off-target cleavages. Exons 1 and 3 of the *Fam210b* locus were targeted. The exon 1 tgt1 targeting sequence was as follows: 5'-GCTCGCGGCGGCCATGGCC-3'; exon 1 tgt2 targeting sequence was 5'-TCGGGCGTCCGGAACCGGAT-3', and the exon 3 targeting sequence was 5'-TACCGGGAGCAAACAGCTTATGGA-3'. Exons 2 and 4 of the *Mfrn1* locus were targeted. The exon 2 targeting sequence was as follows: 5'-GATGCTTGTATACCGGGCTT-3'; the exon 4 targeting sequence was 5'-GAAGAACTCATAAACGGACC-3'. CRISPR guides were cloned into pX330 plasmid (Addgene) with *BbsI* ligation as described previously (75).

### CRISPR/Cas9 genome editing in MEL cells

CRISPR/Cas9 constructs were delivered to MEL DS19 cells by electroporation. CRISPR/Cas9 constructs were co-electroporated with pEF1 $\alpha$  at a 9:1 ratio. Two days later, cells were resuspended in puromycin media (5  $\mu$ g/ml). Cells were plated by limiting dilution at 0.3 cells per well in 96-well plates and cultured at 37 °C. Clones were screened by gDNA isolation (QuickExtract, Epicenter), and PCR for the deletion allele (CRISPR 1), or PCR for the WT allele (CRISPR 2 and CRISPR 3). Clones were validated by Western blot analysis for *FAM210B* protein; *Mfrn1* gene expression was assayed by qRT-PCR (48).

The PCR primers to genotype *Fam210b* CRISPR 1 were as follows: mFam210b\_F 5'-CTATAGAGCCCCGCCTATCC-3' and mFam210b\_R 5'-TCGGTCAGGTGCAATGTAAA-3'. The PCR primers used to genotype *Fam210b* CRISPR 2 and 3 were as follows: E3F 5'-GCATAGACATGTCTGCAATCC-3' and E3R 5'-AGCCACCUUGCACACAGCCC-3'.

### Iron radiolabeling and radio iron-heme measurements

<sup>55</sup>FeCl<sub>3</sub> (specific activity: 1.28 Ci/mmol) (PerkinElmer Life Sciences) were loaded onto transferrin as described previously

(80). Metabolic labeling with  $^{55}\text{Fe}$ -transferrin and quantitation of the incorporation of  $^{55}\text{Fe}$  into heme were carried out as described previously (38).

### ICP analysis of mitochondrial iron content

Mitochondrial iron was measured as described previously (76). Isolated mitochondria were treated with nitric acid, and sample iron content was determined by using an ICP optical emission spectrometer (PerkinElmer Life Sciences). The results were normalized to mitochondrial protein content.

### HPLC analysis of heme and porphyrins

HPLC analysis was carried out with MEL cells that were differentiated for 72 h with 1.5% DMSO. A cell pellet spun down from a 30–50-ml culture was mixed with water to about 200  $\mu\text{l}$  in a microcentrifuge tube and sonicated for 12 cycles with 5-s intervals at 50% duty (about 2.5 s on, 2.5 s off) using a microtip. A 50- $\mu\text{l}$  aliquot was mixed vigorously with 200  $\mu\text{l}$  of an extraction mixture of ethyl acetate (4 volumes) and glacial acetic acid (1 volume). The phases were separated by microcentrifugation for 1 min at maximum speed. The upper organic layer was immediately analyzed simultaneously for protoporphyrin IX and heme in the HPLC (38, 51).

### Silencing of Fam210b in mouse primary fetal liver cells

Silencing of *Fam210b* and hemoglobin quantification in mouse primary fetal liver cells was carried out as described (77). The retroviral plasmids expressing shRNAs for *Fam210b* were transfected in a packaging HEK293T cell line. The collected retroviral supernatants were added to erythroid precursor cells purified from E14.5 mouse fetal liver (C57BL/6J). The retrovirally-transduced cells were sorted for GFP expression using a FACSAria machine (BD Biosciences), followed by *ex vivo* erythroid terminal development in Epo-only medium for 48 h. After culture, cells were collected for analysis on enucleation and hemoglobin content. The hemoglobin content was quantified with Drabkin's reagent used human hemoglobin as standards.

### Fetal liver proliferation and enucleation assays

To measure proliferation rates, GFP-positive cells were plated in 24-well plates with a density at 100,000 cells/per ml, cultured in Epo-only erythroid differentiation medium. The number of cells was counted at the 24 and 48 h using a hemocytometer. In each time point, the cell numbers were the average of three biological samples, which were the average of two technical duplicates. To measure the rate of enucleations, after 48 h of culture, one million cells were spun down and resuspended with 100  $\mu\text{l}$  of FACS buffer (PBS + 2% fetal bovine serum + 100  $\mu\text{M}$  EDTA). Anti-Ter119 antibodies conjugated with APC (eBioscience) and Hoechst 33342 (Sigma) were added to stain the cells for 10 min at room temperature following manufacturer protocols. After staining, cells were washed with cold PBS twice and resuspended with the FACS buffer containing propidium iodide (Sigma) to distinguish between live and dead cells and were further analyzed on a FACS

Fortessa machine. The details of protocols were previously described (78).

### Functional complementation assays with chemicals in cultured cells

MEL cells were differentiated chemically with 2% DMSO for 72 h. For functional complementation assays with hinokitiol, cells were concurrently treated with 10  $\mu\text{M}$  iron citrate and hinokitiol (1  $\mu\text{M}$ ) or C2dOHino (1  $\mu\text{M}$ ) (48). Chemical complementation with DP was carried out by treating cells with 5  $\mu\text{M}$  DP (Frontier Scientific) as described (38).

### Sub-mitochondrial localization studies

Crude mitochondria were isolated from MEL or primary fetal liver cells using a mitochondrial isolation kit (Pierce).

For mitochondrial sublocalization studies, MEL cells stably expressing mouse *Fam210b* were harvested at confluency and lysed in homogenization buffer (20 mM HEPES, pH 7.4, 220 mM mannitol, 70 mM sucrose) supplemented with 2 mg/ml BSA and 0.5 mM PMSF using a Teflon–Dounce homogenizer to release mitochondria. The lysates were centrifuged at  $100 \times g$  at 4 °C for 5 min. The post-nuclear supernatant was centrifuged at  $10,000 \times g$  for 10 min to obtain mitochondria pellets. The pellets were subsequently washed with homogenization buffer without BSA and PMSF, and mitochondria were collected by centrifugation. The protein concentration was measured using BCA assay according to the vendor's protocols (ThermoFisher Scientific).

For mitochondrial fractionation, mitochondria (equivalent of 50  $\mu\text{g}$  of mitochondrial protein per reaction) were suspended in homogenization buffer, 20 mM HEPES (for osmotic shock) or 20 mM HEPES with 1% Triton X-100 in the presence or absence of 5  $\mu\text{g}/\text{ml}$  proteinase K for 30 min at 4 °C. Proteinase K was inactivated with the addition of 1 mM PMSF. Samples were centrifuged at  $10,000 \times g$  at 4 °C to obtain the pellet that contained mitoplasts (the matrix and inner membrane). The supernatant was collected and precipitated with trichloroacetic acid (TCA, final concentration of 15% TCA) to obtain soluble proteins. The proteins (50  $\mu\text{g}$  per lane) were separated by SDS-PAGE followed by immunoblot analysis.

Mitochondria (equivalent of 50  $\mu\text{g}$  of mitochondrial protein per reaction) were suspended in homogenization buffer or 0.1 M sodium carbonate, pH 11.5, for 30 min at 4 °C. Then, the samples were centrifuged at  $28,000 \times g$  for 30 min at 4 °C to obtain the pellet of precipitated membrane proteins. The supernatant with soluble proteins was TCA-precipitated. The proteins (50  $\mu\text{g}$  per lane) were separated by SDS-PAGE followed by immunoblot analysis.

### Western blot analysis

The following antibodies were used in this study: TIMM23 (BD Biosciences catalog no. 611222); TOMM20 (Santa Cruz Biotechnology clone FL-145); Mortalin (Proteintech catalog no. 1488-14-AP). PreP and Yme1L polyclonal antibodies were raised against recombinant proteins (Pacific Immunology). Immunoblotting of FAM210B was performed using a custom anti-mouse FAM210B polyclonal antibody generated against three peptides (C-LSHPVPDARLLRTARGDC, C-TGTEK-

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KLSTRTQQLKKVC, and C-KLGFKESLVQSKMAC), and immunoaffinity-purified against these antigenic peptides (Genemed Synthesis, Inc., San Antonio, TX). Anti-FECH polyclonal (C-20), HA (Y-11), hemoglobin a (H-80), and HSP60 (K-19) antibodies were obtained from Santa Cruz Biotechnology (Athens, GA). Anti-TFRC (H68.4) was obtained from Invitrogen. Anti-GAPDH was obtained from Pierce (GA1R). Anti-FLAG M2-HRP was obtained from Sigma (catalog no. A8592). HSP90 antibody was obtained from Cell Signaling (C45G5). For Fig. 7C, we used anti-FECH polyclonal antibodies from Kerfast (EGA191).

### qRT-PCR

qRT-PCR probes were obtained from Invitrogen. Murine *Fam210b* probe is Mm00508881\_m1; *Hprt* probe is Mm01545399\_m1. Zebrafish *fam210b* probe is Dr03426307\_m1; *hprt* probe is Dr03138604\_m1. The mouse tissue cDNA array was purchased from Clontech.

### Chemical hinokitiol and functional complementation in zebrafish

24-hpf morphant embryos in the transgenic *Tg(globinLCR:eGFP)* line were dechorionated with Pronase as described (54) and incubated in the presence of 1  $\mu$ M hinokitiol + 10  $\mu$ M iron citrate for another 48 h (48). For specificity, the control, inactive C2 deoxyhinokitiol was 1  $\mu$ M final concentration with iron citrate. Vehicle-treated embryos were exposed to 0.01 mM DMSO. Control or morphant embryos at roughly 72 hpf were mechanically homogenized as described previously for flow cytometry (45, 54).

### Genetic complementation assays

Genetic complementation of the  $\Delta$ *mrs3/4* yeast strain with chimeric vertebrate *Fam210b* and *Mfrn1*, containing the first N-terminal 23 amino acid residues from yeast *Mrs3* subcloned in the pRS426-ADH vector, was performed as described previously (23). Western blot analysis of the total, mitochondrial, and cytosolic fraction was described previously with anti-FLAG and anti-porin antibodies.

Control and *Mfrn2*<sup>-/-</sup> mouse embryonic fibroblasts (MEF) were isolated from the skin of 3-month-old mice and immortalized by forcing through a senescence crisis. *Mfrn2*-GFP or *Fam210b*-HA was transiently expressed in these MEF cells. Transfected MEF cells were treated with DFO (10–50  $\mu$ M). Cellular viability was then assayed using a cell titer blue assay, which measures conversion of resazurin to resorufin on a 96-well format (Promega).

### Immunoprecipitation of FAM210B and its interacting proteins

CPOX-FLAG, PPOX-FLAG, FECH-FLAG, and FAM210B-HA were cloned into the pEF1 vector and expressed under the control of the EF1 promoter. pEF1/FAM210B-HA was co-transfected with pEF1 empty vector, pEF1/CPOX-FLAG, pEF1/PPOX-FLAG, or pEF1/FECH-FLAG into 10-cm<sup>3</sup> dishes of HEK293T cells using a Lipofectamine 2000 reagent-based method (Invitrogen). Cells were harvested 48 h post-transfection and lysed in a buffer containing 1% Nonidet P-40, 300 mM NaCl, 50 mM Tris, pH 8.0, and protease

inhibitor mixture (Pierce). Cell lysates were precleared with mouse IgG-agarose (Sigma) for 60 min at 4 °C. Precleared lysates were then incubated with anti-FLAG or anti-HA-agarose beads (Sigma) for 2 h at 4 °C. Beads were washed three times for 15 min in lysis buffer. Immunoprecipitated proteins were subsequently eluted with Laemmli buffer and resolved by SDS-PAGE.

To determine whether FAM210B interacted with endogenous FECH, we electroporated pEF1/FAM210B-FLAG into MEL cells and selected FAM210B-FLAG expressing single cell clones by limiting dilution. pEF1/FAM210B-HA and pEF1 empty vector expressing cell lines were differentiated with 1.5% DMSO for 72 h. For each immunoprecipitation experiment, we lysed 100 million differentiated cells with 1 ml of lysis buffer (77). 2 mg of protein was used as the starting material for each immunoprecipitation sample. Samples were precleared with mouse IgG beads and immunoprecipitated with anti-FLAG M2-agarose for 4 h at 4 °C. Beads were washed three times for 15 min at 4 °C. Proteins were eluted with Laemmli buffer, resolved by SDS-PAGE, and analyzed by Western blotting.

### Statistics

Statistical analysis was carried out using two-tailed paired or unpaired Student's *t* test. Significance was set at *p* < 0.05. All data points were plotted as scatter plots, representing all replicates of each experiment.

### Study approval

All zebrafish experiments were performed in accordance with the Institutional Animal Care and Use Committee (IACUC) regulations of Brigham and Women's Hospital. WT AB zebrafish were used for all zebrafish experiments. All mouse experiments were performed in accordance with IACUC regulations at the Massachusetts Institute of Technology and University of Utah.

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