The neurodegenerative disease Friedreich's ataxia arises from a deficiency of frataxin, a protein that promotes iron–sulfur cluster (ISC) assembly in mitochondria. Here, primarily using Mössbauer spectroscopy, we investigated the iron content of a yeast strain in which expression of yeast frataxin homolog 1 (Yfh1), oxygenation conditions, iron concentrations, and metabolic modes were varied. We found that aerobic fermenting Yfh1-depleted cells grew slowly and accumulated FeIII nanoparticles, unlike WT cells. Under hypoxic conditions, the same mutant cells grew at rates similar to WT cells, had similar iron content, and were dominated by FeII rather than FeIII nanoparticles. Furthermore, mitochondria from mutant hypoxic cells contained approximately the same levels of ISCs as WT cells, confirming that Yfh1 is not required for ISC assembly. These cells also did not accumulate excessive iron, indicating that iron accumulation into yfh1-deficient mitochondria is stimulated by O2. In addition, in aerobic WT cells, we found that vacuoles stored FeIII, whereas under hypoxic fermenting conditions, vacuolar iron was reduced to FeII. Under respiring conditions, vacuoles of Yfh1-deficient cells contained FeIII, and nanoparticles accumulated only under aerobic conditions. Taken together, these results informed a mathematical model of iron trafficking and regulation in cells that could semiquantitatively simulate the Yfh1-deficiency phenotype. Simulations suggested partially independent regulation in which cellular iron import is regulated by ISC activity in mitochondria, mitochondrial iron import is regulated by a mitochondrial FeII pool, and vacuolar iron import is regulated by cytosolic FeII and mitochondrial ISC activity.

Humans suffering from Friedreich's ataxia are deficient in frataxin (1). This protein binds a multiprotein complex in the mitochondrial matrix that catalyzes assembly of iron–sulfur clusters (ISCs) (2). Yeast frataxin homolog 1 (Yfh1) in *Saccharomyces cerevisiae* acts similarly, which allows budding yeast to serve as a workhorse for probing the pathogenic mechanism of this disease. Frataxin binds cysteine desulfurase, a member of this complex, which transfers the rate by which a sulfur atom is extracted from free cysteine and transferred to the scaffold. Once oxidized and reduced by ferredoxin (Yah1), this sulfur becomes the bridging sulfide of a [Fe3S2]2⁺ cluster bound to the Isu1/2 scaffold proteins (3). Δyfh1 yeast cells grown under aerobic conditions are deficient in mitochondrial ISCs and heme centers, as well as other phenotypic properties that are less easily interpreted (4–7). For example, Yfh1 protects cells against oxidative stress (8).

Fermenting aerobic Δyfh1 yeast cells import excessive amounts of iron which flows into mitochondria and reacts to form FeIII nanoparticles (5, 6). The iron concentration of mitochondria in which nanoparticles have accumulated is 10–15-fold higher than in the WT organelle (4, 9). This accumulation indicates iron dysregulation. Fermenting Δyfh1 cells grow slowly relative to WT cells and are more easily damaged by O2 and other reactive oxygen species (ROS) (10). How these properties are causally related at the molecular level remains incompletely understood. Understanding these relationships, the long-term goal of this study, may reveal better approaches for treating the disease.

The model of Figure 1 describes our current understanding of these relationships, as supported by Mössbauer (MB)-based data presented here. First some background.

In our study, IRON refers to ⁵⁷FeIII citrate which we added to growth media (note: nutrients are capitalized). IRON is typically imported into the cell through the Fet3 transport system on the plasma membrane (11, 12). However, under hypoxic conditions, the Fet4 importer is used. Imported iron ultimately generates the cytosolic labile FeIII pool (13) called FC in the model (note: model components are bold when introduced).

Vacuoles are acidic organelles that reversibly store iron. Under iron-replete conditions, they import FC via the membrane-bound transporter Ccc1 (14). Imported FeII in vacuoles (F2 in the model) is rapidly oxidized to form FeIII.
Cellular iron redistribution in Yfh1-deficient yeast

Figure 1. Model describing iron trafficking and regulation in *S. cerevisiae*, with an emphasis on the cell’s response to a decline in Yfh1 levels under aerobic versus hypoxic growth and high versus low iron in the media. Blue region is cytosol, yellow is mitochondria, and green is vacuole. Points of regulation are indicated by red circles. The dashed red line emanates from the sensed species and terminates at the site of regulation. Regulation was "local"; F5 (the iron regilon) regulates Rcyt, Rvac, and RFS FM (pool of FeII in mitochondria) regulates Rmit, FC (labile iron pool in the cytosol) also regulates Rorg; O2 regulates Rmit, F2, vacuolar FeII; F3, vacuolar FeII; FC, cytosolic iron; FM, mitochondrial FeII pool; FS, iron-sulfur clusters and FeII hemes; MP, mitochondrial nanoparticles; ROS, reactive oxygen species; VP, vacuolar nanoparticles; Yfh1, Yeast frataxin homolog 1.

Polyphosphate complexes, F3 (15–17). F3 can be converted to vacuolar nanoparticles (VPs) (18).

Some FC is converted into CIA, a model component representing all iron species in the cytosol, nucleus, and endoplasmic reticulum except for FC. CIA includes [Fe4S4]-containing proteins that are synthesized by the Cytosolic Iron–sulfur cluster Assembly system.

Much FC is trafficked into mitochondria through two paralogous high-affinity FeII importers (Mrs3 and Mrs4) on the inner membrane (15, 16). Iron entering mitochondria forms an FeII pool (17, 19), referred to as FM, which serves as substrate for the biosynthesis of mitochondrial ISC and hemes, collectively symbolized as FS. Yfh1 is presumed to be a catalyst for the FM → FS reaction; the same reaction is inhibited by mitochondrial component O2.

Understanding the cell’s response to Yfh1 deficiency requires understanding how genes involved in iron homeostasis are regulated (indicated by the red circles and dashed lines in Fig. 1). The iron regilon includes a few dozen genes whose expression is controlled by Aft1 and Aft2 (20–22). These paralogous transcription factors regulate (a) iron import into the cell; (b) iron import into mitochondria (23); and (c) iron export from vacuoles into the cytosol. Thus, iron export from vacuoles to cytosol is commonly thought to be regulated similar to the import of nutrient iron into cytosol (24).

Other systems help regulate iron homeostasis in the cell. Besides Aft1/2, Fet4 expression is regulated by O2 (24). Under iron-replete conditions, vacuolar iron import through Ccc1 is positively regulated by Yap5, another iron-sensing transcription factor (25), as well as by cellular stress factors. Under iron-starved conditions, Cth1 and Cth2 mediate degradation of Ccc1 mRNA (26). Other regulatory factors add further complexity to understanding regulation (27–29). The generation of mitochondrial nanoparticles (MP in model), loss of ISC activity, and excessive oxidative damage are major cellular responses to Yfh1 deficiency (7, 29–31).

One strategy for disentangling the confounding effects of O2 is to minimize its exposure to Δyfh1 cells (15, 30, 32, 33). Hypoxic conditions are more useful than rigorously anaerobic ones because low concentrations of O2 are required for heme biosynthesis. Δyfh1 cells grow slowly under aerobic conditions but faster under hypoxic ones. When grown hypoxically on YPAD, Δyfh1 cells exhibit no obvious phenotype (32) except for low aconitase activities (30). Aconitase is used as a reporter of ISC assembly activity because it requires an [Fe4S4] cluster for activity. Aerobic fermenting Δyfh1 cells lack aconitase activity, so the presence of ISC in the same cells grown hypoxically suggests that Yfh1 is not required for ISC biosynthesis.

Frataxin also minimizes oxidative stress. When anaerobic Δyfh1 and WT cells are exposed to air, oxidative damage to Δyfh1 cells increases. However, ISC-containing ferredoxin is quite stable inside Δyfh1 mitochondria (32, 34) suggesting that O2 inhibits ISC biosynthesis.

The respiratory shield hypothesis (35–37) forms the core of the Figure 1 model. It assumes that the mitochondrial matrix of healthy WT cells grown under aerobic conditions is largely devoid of O2 due to respiratory activity on the inner membrane. In WT cells, the shield protects FM from reacting with O2. In the absence of Yfh1 or other proteins involved in ISC assembly or trafficking, the reaction FM → FS is slowed, which slows the rate of O2 reduction by the respiratory complexes. This allows more O2 to diffuse into the matrix region where it reacts with FM to form MP, thereby limiting the amount of FM available as a substrate for FS. This weakens the respiratory shield and initiates a vicious cycle in which the further decline of FM further slows the synthesis of FS and the rate of respiration. The lack of FS activates the iron regilon which increases iron import into the cell and mitochondria. The net effect is that mitochondria spiral down into a diseased state in which (a) the organelle is filled with nanoparticles; (b) both ISC and heme activities are low; and (c) O2 and ROS flood the cell.

Similar accumulations of nanoparticles in mitochondria are observed for other genetic strains of yeast harboring defects in ISC biosynthesis or trafficking including cells deficient in Yah1 (31), Ggc1 (6), Atm1 (33), and Ssq1 (6). Even more surprising is that a similar phenotype is observed in strains with mutations that are not primarily involved in either ISC assembly or trafficking. This includes cells deficient in Mtm1 (38) and cells...
in which the Aft1 transcription factor is constitutively upregulated (39). Thus, iron dysregulation appears to be a secondary response to defects in various mitochondrial processes including but not limited to ISC assembly, trafficking, and/or regulation. The respiratory shield hypothesis can explain these diverse causes, since the strength of the shield reflects processes such as respiration, bioenergetics, and mitochondrial membrane potential in which many proteins contribute.

Mitochondria from Δyfh1 yeast cells (reportedly) do not accumulate excessive iron when they respire (40). Moreover, aconitase activities in respiring Δyfh1 cells are ~70% of WT levels, whereas they are near zero in aerobic fermenting Δyfh1 cells. Understanding these metabolic differences might reveal new insights into the mechanism of the disease.

For this study, we primarily used MB spectroscopy to probe the relationships giving rise to the Yfh1-deficiency phenotype. We investigated a strain of yeast in which Yfh1 expression is dictated by the estradiol concentration in the growth medium (6, 41). This strain allowed us to probe the effects of gradually reducing the expression of Yfh1. We also examined the effects of O2, nutrient iron, and metabolic mode on cellular iron content. We evaluated the iron content of mitochondria from anaerobic fermenting and respiring Yfh1-deficient cells grown under hypoxic conditions. We then used those and other results to develop the model of Figure 1 and then used the model to simulate the Yfh1-deficient phenotype. Our results offer new insights into the pathogenesis of Friedreich’s ataxia.

**Results**

We employed the mutant strain ERyfh1 in this study because the expression level of yfh1 could be varied according to the concentration of estradiol [EST] added to the growth medium (6, 41). As expected, Yfh1 was undetectable in a Western blot of soluble lysates from cells grown in the absence of EST ([EST] = 0), and expression increased as [EST] increased (Fig. 2).

The [EST] required for full expression was higher than that reported by Seguin et al. (6, 41). They reported that [EST] > 10 nM was sufficient to generate WT expression levels of Yfh1, whereas our samples required roughly 1000 nM [EST] to exhibit WT properties.

**Cellular iron redistribution in Yfh1-deficient yeast**

We used a shorthand nomenclature to describe the growth conditions for various experiments. Aerobic fermenting wild-type (AFW) cells cultured in glucose-containing minimal medium (MM) grew in accordance with an exponential growth rate $\alpha = 0.2$ to 0.3 h$^{-1}$ where $\alpha$ is the linear slope of ln(A600) versus time plots (Table S1). Aerobic fermenting mutant cells (AFM) grew at similar rates ($\alpha = 0.2$ h$^{-1}$) at [EST] > ~25 nM but slower at [EST] = 0. Supplementing media with Fe$^{III}$ citrate at concentrations between 1 → 100 μM had no effect on growth rate. WT cells cultured on 1 μM Fe$^{III}$ citrate grow at rates comparable to those of iron-replete WT cells (19).

AFM cells grown in media containing [EST] = 0 and [IRON] = 40 will be called AFM0-40 cells, where the first number refers to the nM concentration of EST and the second to the μM concentration of Fe$^{III}$ citrate. Both are final concentrations in the growth media. WT cells averaged ~600 μM iron, similar to reported values (17). The iron concentration of AFM0-40 cells was significantly higher than that of AFW-40 cells (Table S1). The relative intensities of the MB spectra (see below) confirmed that AFM0-40 cells were iron-overloaded.

AFM0 cells contained 500 to 1800 μM Fe depending on growth media. As [EST] increased, the cellular iron concentration declined, eventually reaching concentrations typical of WT cells. This supports the observed decline in MB spectral intensities as [EST] increased (see below). The similar iron concentrations obtained under these conditions suggested that AFM cells expressing WT levels of Yfh1 were not iron-dysregulated, as expected.

When grown under hypoxic conditions, both fermenting WT and mutant cells (called HFW and HFM cells, respectively) grew in accordance with an exponential growth rate $\alpha = 0.2$ h$^{-1}$ regardless of [EST] or [IRON]. The concentration of iron in mutant cells under hypoxic conditions was comparable to that of WT cells (Table S1). Under hypoxic conditions, mutant cells were not noticeably dysregulated.

**MB spectra of AFM cells grown with different [EST]**

We used MB spectroscopy to investigate how the iron content of the mutant cells changed as [EST] decreased. As expected, AFM1000-40 cells exhibited MB spectra typical of WT (W303) cells (compare Fig. 3A to Fig. 1D of Ref (17)). Spectral features included (a) a magnetic feature arising from vacuolar non-heme high spin (NHHS) Fe$^{III}$ polyphosphate (brown line); (b) a central doublet (CD) arising from [Fe$_4$S$_4^{2+}$]clusters and LS Fe$^{III}$ hemes combined (green line); and (c) two minor doublets arising from NHHS Fe$^{III}$ (dark blue line) and HS Fe$^{III}$ hemes (light blue). Some CD originated from mitochondrial iron, while the remainder arose mainly from [Fe$_4$S$_4^{2+}$] clusters in the cytosol and nucleus. A low-intensity doublet likely due to nanoparticles (magenta) was also present. Percentages used in spectral decompositions are in Table S2.

When [EST] was lowered to 100 nM, spectral intensity due to vacuolar NHHS Fe$^{III}$ declined while that of the NHHS Fe$^{III}$ doublet increased (Fig. 3B). AFM25-40 and AFM0-40 spectra
were essentially devoid of the HS Fe\textsuperscript{III} feature. A relatively intense NHHS Fe\textsuperscript{II} doublet was evident in all spectra between AFM100-40 → AFM0-40. A minor shoulder on the inside edge of the low-energy line of this doublet (see arrow) probably arose from the CD (represented as the sum of FS + CIA in the model), but resolution was limited. According to the model of Figure 1, the early loss of vacuolar NHHS Fe\textsuperscript{III} (F3 in the model) indicates that the earliest titration event that occurs as Yfh1 levels decline is the reduction of F3 → F2. The observed NHHS Fe\textsuperscript{II} doublet arose from some combination of modeling components F2, FC, and FM; the three cannot be distinguished by MB. Although this doublet represented just 9% of the intensity of the AFM0-40 spectrum, the absolute cellular concentration of the Fe\textsuperscript{II} species giving rise to it was significant. A similar doublet has been observed along with the dominating nanoparticle doublet in Yah1- and Atm1-decient mitochondria (33). The Fe\textsuperscript{II} doublet remained throughout the titration. Most of the NHHS Fe\textsuperscript{II} intensity probably arose from vacuolar Fe\textsuperscript{II} (F2 in the model), but this iron might have also been exported from vacuoles forming cytosolic Fe II (FC in the model).

When [EST] was reduced to 10 nM in an AFM sample, the resulting spectrum (Fig. 3E) remained dominated by the NHHS Fe\textsuperscript{II} doublet. However, the low-energy line of this doublet was broadened suggesting the initial development of nanoparticles (MP in model). This feature increased in the 5 nM and 2.4 nM samples (Fig. 3, F and G). When [EST] = 0, in cells grown with 40 μM \textsuperscript{57}Fe\textsuperscript{III} citrate, the overall spectral intensity (Fig. 3H) increased dramatically (more so when MM rather than complete synthetic medium was used as the growth medium). Virtually, all the extra spectral intensity arose from a quadrupole doublet attributed to nanoparticles.

Seguin et al. 2010 (6) reported that mutant aerobic cells accumulated large amounts of iron when [EST] = 0 but not when [EST] = 2.5 nM (at that concentration there was a modest rise in cellular iron concentration relative to in WT cells). They also reported that aconitase activity was undetectable at [EST] = 0 but was at WT levels when [EST] = 2.5 nM. Their results indicate a causal correlation between (a) loss of ISC activity; (b) increased rate of iron import into the mitochondria; and (c) increased level of mitochondrial nanoparticles.

We also examined the electron paramagnetic resonance (EPR) spectra of mutant cells with different Yfh1 expression levels. AFM0-10 cells exhibited an intense broad EPR signal near g = 2.0 due to nanoparticles (Fig. 4, Panel A, dashed line). Similar signals have been observed in spectra of Yah1- and Atm1-depleted cells (31, 33). In contrast, AFM2000-10 cells

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**Figure 3. Mössbauer spectra of AFM cells with decreasing [EST] and [IRON] = 40.** [EST] in nM was: A, 1000; B, 100; C, 50; D, 25; E, 10; F, 5; G, 2.5; H, 0. Red lines are composite simulations based on parameters in Table S2. Solid lines above A are simulations for nanoparticles (magenta), non-heme HS Fe\textsuperscript{II} (dark blue), heme HS Fe\textsuperscript{II} (light blue), the central doublet (green) and vacuolar Fe\textsuperscript{III} (brown). All MB spectra included in the paper were collected at ~5 K and 0.05 T and with gamma radiation parallel to the field. Note scale differences between spectra. Pie charts show the iron distribution for the corresponding spectra (as given in Table S2), using the same color coding as for the solid spectral line simulations. AFM, aerobic fermenting mutant cells; EST, estradiol; MB, Mössbauer.
confirmed that and revealed that most iron in such cells was 
NHHS FeIII (F2 + FC + FM combined). An O2-dependence for 
oxidizing F2 → F3 had not been previously recognized. Such 
an O2-dependence is included in the model of Figure 1.

Manganese species in Yfh1-deficient cells

Although not the focus of this study, EPR spectra of mutant 
cells exhibited a noticeable hyperfine-split signal in the g = 2 
region due to mononuclear MnII ions. Its intensity was dimin-
ished in HFM2000-10 cells relative to AFM2000-10 cells (Fig. 4, 
Panel B red versus blue spectra). Under aerobic conditions, 
the parent strain contained 20 ± 5 μM Mn (n = 2) which is similar 
to the average of previous reports for aerobic WT fermenting cells 
(17, 19, 42), namely 26 ± 7 μM. AFM0-40 cells contained sub-
stantially higher Mn concentrations (70 ± 11 μM; n = 4) which 
may be related to an increased expression of Mn superoxide 
dismutase in response to oxidative stress (43). Consistent with 
this, the cellular Mn concentration in our titration series 
increased 2- to 5-fold as the [EST] concentration decreased 
(data not shown). HFM0 cells contained 27 ± 5 μM Mn (n = 2) 
which may reflect a return of Mn superoxide dismutase to WT 
levels under hypoxic conditions (44).

Nanoparticle loading declined in aerobic Yfh1-deficient cells 
grown at high [IRON]

WT cells grown on media containing 40 μM 57FeIII citrate 
store high-spin FeIII polyphosphate complexes in vacuoles 
(45); these ions exhibit magnetic MB features that span 
from −10 to +10 mm/s velocity (Fig. 5B). WT cells grown with 
1 μM 57FeIII citrate did not contain much vacuolar FeIII 
(Fig. 5A); this is the standard state for WT yeast cells (17). 
Rather, these WT cells exhibited an intense NHHS FeII 
doublet for cells grown on [IRON] = 1 μM. Based on our 
model, this doublet arose from the sum of [FC] + [F2] + [FM] 
(caused by activation of the Iron-Regulon). We are not sure 
which of these species dominates, but it is probably [FC] or 
[F2]. Holmes-Hampton et al. observed similar behavior under 
iron-deficient conditions and attributed the strong NHHS FeII 
doublet mainly to F2 (17).

Under the same low-iron conditions, mutant Yfh1−deficient 
cells exhibited a fundamentally different MB spectrum, one 
dominated by nanoparticles. Essentially identical spectra were 
observed for AFM0-1 (Fig. 5C) or AFM0-10 (Fig. 5D) cells. 
Surprisingly, spectral intensity was independent of nutrient 
iron concentration within this range. This differs from 
Auchere et al. (46) who reported that the extent of iron 
accumulation increases with increasing nutrient iron levels. 
However, comparisons are difficult because of differences in 
experimental details.

We were also surprised that spectra of Yfh1-deficient 
mutant cells grown aerobically on 100 μM 57FeIII citrate 
were 5-fold less intense than those of cells grown in media 
containing lower iron concentrations (in Fig. 5, compare ab-
sorption [%] scale in E versus in C or D). We observed a similar 
reduction in nanoparticle spectral intensity in three indepen-
dent preparations. Curiously, this result implies that Yfh1−
deficient cells grown on media containing high nutrient iron are less dysregulated than when grown on low nutrient iron. Consistent with this, Seguin et al. found that excess nutrient iron (up to 2.5 mM) improved the growth of Δyfh1 cells (6).

AFM0 cells also lacked the vacuolar Fe\textsuperscript{III} species (Fig. 5, C–E) that is present in WT cells even at high iron (Fig. 5B). The AFM0 spectra also exhibited significant HS Fe\textsuperscript{II} features. Percentages of spectral features are listed in Table S2. Vacuolar iron export should accelerate in AFM cells for the same reason that these cells accumulate iron—i.e., activation of the iron regulon increases expression of both Fet3 and Fet5. However, the regulatory details may be more complicated.

**MB of hypoxic (HFW and HFM) cells**

We performed similar experiments on fermenting cells grown under hypoxic conditions. MB spectra of HFW cells were dominated by a NHHS Fe\textsuperscript{II} doublet (Fig. 6, A and B) and lacked features due to vacuolar Fe\textsuperscript{III}. Δmtm1 and adenine-deficient WT cells exhibit similar spectra (38, 47). The minor shoulder at ca. +1 mm/s in Figure 6, A and B was likely the high-energy line of the CD.

Spectra of HFM0 cells were also dominated by a NHHS Fe\textsuperscript{II} doublet (Fig. 6, C and D). There was a small shoulder resolved from the low-energy line of the main doublet which probably arose from the CD (Fig. 6C, arrow). The major difference, relative to WT cells, was that the spectral intensity for the NHHS Fe\textsuperscript{II} doublet was quite variable, with the intensity of some spectra comparable to WT spectra, while the intensity of other mutant spectra was greater than WT spectra (Fig. S1). We suspect that this variability was due to small differences in the percentage of O\textsubscript{2} used in growing hypoxic mutant cells.

Spectra of HFW-1, HFW-40, HFM0-1, and HFM0-40 cells did not exhibit the magnetic feature due to vacuolar Fe\textsuperscript{III}. With low [O\textsubscript{2}] bubbling through the growth medium, the rate of F2 \textrightarrow F3 oxidation appears to have been slowed. A sensitive balance of factors, including [O\textsubscript{2}] levels, appears to control the oxidation state of vacuolar iron and the rate of iron import into the cell.
MB spectra of respiring cells

Although respiration requires O₂, the hypoxic conditions used here (~5% O₂) allowed both WT and mutant cells to respire slowly on glycerol/ethanol while limiting ROS damage. Thus, we investigated the iron content of hypoxic respiring cells. MB spectra of aerobic respiring ARW-40 cells (Fig. 7A) were comparable to those of fermenting AFW-40 cells (Figs. 3A or 5B) except that the CD contributed a higher spectral percentage, consistent with a greater use of mitochondria during respiration.

The spectrum of hypoxic HRW-40 cells (Fig. 7B) was dominated by the CD and a NHHS FeII doublet. The loss of vacuolar FeIII (probably due to reduction to FeII) as respiring WT cells transitioned to hypoxic conditions indicated a sensitivity to metabolic mode. This suggested that the redox state of vacuolar iron in WT cells is determined by the O₂ status during cell growth (with lower O₂ favoring FeII vacuolar iron).

The spectrum of aerobic ARM0-40 mutant cells was dominated by nanoparticles, with significant vacuolar FeIII and little NHHS FeII (Fig. 7C). There is little doubt that the nanoparticles in both respiring and fermenting aerobic cells were in mitochondria; this accumulation indicated that the iron regulon under both metabolic modes was activated. Thus, aerobic mutant cells were strongly iron dysregulated, regardless of metabolic mode. However, respiring mutant cells must have been dysregulated differently than fermenting mutant cells because vacuolar FeIII was present in respiring mutant cells but lacking in fermenting ones. Since both Fet3 and Fet5 are part of the iron regulon, both genes should have been strongly expressed in both the respiring and fermenting mutant cells. Our results suggest that iron regulon activation is not the only factor that determines the redox status of vacuolar iron.

Spectra of hypoxic respiring HRM0-40 cells exhibited a dominant vacuolar FeIII signal, not a nanoparticle doublet (Fig. 7D). The NHHS FeII doublet was also intense. In contrast, the spectrum of hypoxic fermenting HFM0 cells were dominated by NHHS FeII (e.g., Fig. 6, C and D) with no NHHS FeIII and little if any CD. The spectra of respiring hypoxic mutant cells overall were quite intense (compared to Fig. 3A), suggesting that the cells are dysregulated. In any event, these results suggest that the redox state of vacuolar iron and perhaps the rates of iron import/export into vacuoles are independent of the iron regulation/dysregulation status of the mitochondria or whole cell.

Isolated Yfh1-deficient mitochondria from hypoxic cells

Mitochondria isolated from aerobic Yfh1-deficient fermenting cells accumulate large quantities of FeIII oxyhydroxide nanoparticles (5). In contrast, mitochondria isolated from corresponding hypoxic cells in the current study were not iron-overloaded, regardless of whether mutant cells were grown with [IRON] = 1 or 40 (Fig. 8, C and D). Rather the spectra were dominated by the CD, as in spectra of mitochondria from WT cells (17, 48–50). Mitochondria from analogous HFW cells exhibited similar spectra (Fig. 8, A and B). In all four spectra, 50% to 60% of the intensity was due to the CD. Much of the remaining intensity was due to a NHHS FeII doublet. Previously reported spectra of mitochondria isolated from Yah1-deficient and Atm1-deficient cells grown under anaerobic/hypoxic conditions were similar (31, 33).

We also examined mitochondria isolated from HRM0-40 and HRW-40 cells. In these cases, the dominating feature in both spectra was the CD (Fig. 9, A and B), similar to spectra of mitochondria from aerobic respiring WT cells (48–50). The mitochondria from these hypoxic cells were not iron-overloaded and did not display spectral features due to nanoparticles.

Aconitase activities were determined in four other batches of mitochondria isolated from cells grown under hypoxic conditions, one each for HRW-1, HRW-40, HRMO-1, and HRMO-40. Activities were 1.14 ± 0.01, 1.07 ± 0.22, 2.5 ± 0.37, and 1.31 ± 0.06 units/mg protein, respectively (uncertainties are standard deviations from four measurements). We conclude that mitochondria from both WT and mutant cells,

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**Figure 7. Mössbauer spectra of respiring WT and Yfh1-deficient cells grown under aerobic and hypoxic iron-sufficient conditions.** A, ARW-40; B, HRW-40; C, ARM0-40; D, HRM0-40.
grown under hypoxic conditions, had significant and comparable ISC activities.

Taken collectively, these results provided unambiguous spectroscopic and enzymatic evidence that [Fe₄S₄] clusters can be assembled and are stable in cells lacking Yfh1—but only if hypoxic conditions are used.

Thus, the lack of ISCs in AFM cells is not directly due to the lack of Yfh1. Such clusters are synthesized in the absence of Yfh1, but at a slower rate (16). As long as O₂ concentrations in the matrix are low, clusters can be assembled. Perhaps the persulfide formed on Nfs1 is more stable under hypoxic conditions. Under aerobic conditions, perhaps more O₂ enters the mitochondrial matrix of mutant cells (due to a weakened Respiratory Shield) where it inhibits ISC assembly. O₂ also likely reacts with the FeII pool in the mitochondria. Both effects would weaken the shield in a vicious cycle.

Although the spectral features exhibited by mitochondria isolated from hypoxic respiring mutant (Fig. 9A) and WT (Fig. 9B) cells were qualitatively similar, the percent-effects differed significantly; the WT spectrum was more intense than the mutant spectrum (22% versus 14%). Additionally, the MB cup for the WT sample was only ¼ filled, whereas that for the mutant cells was ¾ filled. Taking these differences into account indicates that respiring mitochondria isolated from hypoxic WT cells contained nearly 5× as much iron as comparable mutant mitochondria (assuming the same purity).

Mathematical model

We began with the ordinary differential equation (ODE)-based model used to simulate the ΔΔmrs3/4 phenotype (19, 36) and minimally modified it as necessary to simulate the Δyfh1 phenotype. The chemical model shown in Figure 1 included 10 reactions (Table S3) and nine components (defined in the Introduction). At the core of the model was the respiratory shield which can simulate an abrupt disease ↔ healthy transition. The assumed rate-law expressions for all reactions of the model are in Table S3. These expressions, along with the reaction network, were used to generate a set of ODEs describing the change in component concentrations with time (Table S4).

In silico WT cells and hypoxic Δyfh1 cells were defined to grow at exponential rate α = 0.20 h⁻¹, whereas aerobic Δyfh1 cells grew more slowly. Cell growth was simulated by adding the term -α[Cᵢ] to the ODE for each component Cᵢ. This term allowed the system to be solved under an "expanding steady-state" condition (51) in which the volume of a cell is presumed to increase at a constant rate and the rates of all of the modeled reactions are time invariant. This was done by integrating the ODEs, along with a set of initial conditions, using the NDSolve function in Mathematica 10 (Wolfram Software) at long integration times. In silico cells grew on fixed nutrients [IRON] = 1, 2, 11, or 41 μM and [RO2] = 5 (hypoxic) or 100 (aerobic) μM. To account for the effects of hypoxia, O₂ was assumed to inhibit the reaction FM → FS and to control the rate of vacuolar iron oxidation, R₂₃. As before, Rᵐᵖ, Rₑ₃ and
$R_{O2}$ had O$_2$ dependences included in their rate-law expressions.

The major modification relative to our previous model was to include Reg functions which allowed the system to be regulated. Previously, rates of regulated reactions were manually adjusted depending on cellular state. Here, they were augmented with Reg functions (35). Reg functions are surrogates of complex signal-transduction regulatory mechanisms, characterized by (a) the sensed species S (a component of the model); (b) a set-point concentration of S ($[S]_{sp}$); and (c) sensitivity factor $n$.

We initially attempted “global” regulation by assuming Reg±(FS) functions for reactions $R_{cyt}$, $R_{mit}$, $R_{vac}$, and $R_{23}$. Doing so implied control by the Aft1/2-dependent iron regulon, the most well-established mechanism of iron regulation in yeast cells. However, this attempt proved incapable of simulating observed behavior. A Reg-(FS) function successfully regulated $R_{cyt}$, but the same function regulating $R_{mit}$ afforded excessively high [FM] and low [FS]. This mismatch was corrected by regulating $R_{mit}$ with a Reg-(FM) function. $R_{vac}$ and $R_{23}$ were best regulated by assuming the “dual” function (Reg+(FS)-Reg+(FC)) in which both high [FS] and [FC] contributed to increasing the flow of iron into vacuoles. We also added a Reg-(O2) function to regulate $R_{stir}$.

We focused on five different cellular states, including aerobic WT, hypoxic WT, aerobic Δyfh1, hypoxic Δyfh1, and aerobic ΔΔmrs3/4. The model included 35 adjustable parameters (Table S5). For these parameters, we initially assumed previous values (36) and adjusted them conservatively to achieve desired behavior. Values of 14 parameters remained invariant (36). Values for 14 other parameters were changed relative to our previous study but were invariant for all five current cellular states.

The sensitivity of the WT aerobic model to variations in parameters revealed that four of the nine most sensitive parameters involved the ISU reaction in mitochondria and three involved iron import into the cell (Table S8). Of the six least sensitive parameters, four involved vacuoles.

We assessed the viability of simulations qualitatively by their ability to mimic certain phenotypic behaviors while adhering to experimental iron concentrations. This approach was effective in generating the desired qualitative behavior. However, we cannot guarantee that the parameters in Table S5 are unique in eliciting these behaviors or are the globally best at doing so.

We first optimized the behavior of WT and ΔΔmrs3/4 simulations and then focused on WT hypoxic, Δyfh1 aerobic, and Δyfh1 hypoxic states. To simulate these latter states, we adjusted the easily justifiable parameters first and then minimally adjusted remaining parameters as needed. Of the remaining seven adjustable parameters, four were easily justified according to the established properties of the particular cellular state. Specifically, $R_{O2}$ was lowered to create hypoxic conditions, $R_{stir}$ was lowered to create Yfh1-deficient conditions, as expected for Yfh1-deficient conditions, and $R_{mit}$ was lowered to slow mitochondrial iron import in ΔΔmrs3/4 cells, again as expected for Mrs3/4-deficient conditions. The growth rate ($\alpha$) for aerobic Δyfh1 cells was set lower than for WT cells, hypoxic Δyfh1 cells, and ΔΔmrs3/4 cells grown at high nutrient iron, consistent with observations. This left only three parameters that were adjusted without clear justification. Of those, $K_{m}$ (Michaelis–Menten–like parameter describing growth rate) was adjusted for only one of five states, and $K_{N}$ (same for iron import into the cell) and $k_{cia-max}$ (rate-constant for generating CIA) were adjusted for only two of five states. The optimized model simulated the phenotype of these five cellular states with overall semi-quantitative fidelity (Table S6).

In Figure 10, simulations were compared to MB spectra. Contributions from [FC], [FM], and [F2] were summed and normalized to simulated cellular iron concentrations (Table S7), as this allowed comparison to the NHHS Fe$^{III}$ species observed by MB spectroscopy. Likewise, [FS] and [CIA] simulations were summed and normalized in the same way, so that they could be compared to the central quadrupole doublet (CD) observed by MB. Similarly, [MP] and [VP] were summed and normalized, so that they could be compared to the nanoparticle contribution observed in MB spectra. Finally, [F3] was normalized and compared to the NHHS Fe$^{III}$ feature in MB spectra.

**Figure 10. Simulations of MB-related species, given as a percentage of cellular iron, versus nutrient iron concentration.** Maroon, CD; green, NHHS Fe$^{III}$; blue, NHHS Fe$^{II}$; red, nanoparticles. Nutrient IRON is plotted as log base two for better visualization. CD, central quadrupole doublet; MB, Mössbauer; NHHS, non-heme high spin.
For aerobic WT cells, simulations correctly predicted that vacuolar Fe\textsuperscript{III} would dominate MB spectra of these cells at high [IRON]. They also correctly predicted similar relative intensities of the CD, NHHS Fe\textsuperscript{II}, and nanoparticles, in that order (Fig. 10A). As [IRON] declined, simulated vacuolar Fe\textsuperscript{III} levels declined causing the CD to increase percentagewise, as observed in Figure 5, A and B. Simulated Fe\textsuperscript{II} and nanoparticle percentages were low, as observed.

For hypoxic WT cells, simulations at high [IRON] were dominated by Fe\textsuperscript{II}, as observed (compare Figs. 10B to 6, A and B). This was followed by the CD, and little Fe\textsuperscript{III} or nanoparticle intensities. Simulations indicated more CD (relative to Fe\textsuperscript{III}) than was observed.

For aerobic mutant cells, nanoparticles dominated as observed (compare Figs. 10C to 5, C–E), and they declined somewhat at high [IRON], also as observed. For hypoxic mutant cells, simulated Fe\textsuperscript{II} dominated as observed (compare Figs. 10D to 6 C, and D). Relative to WT cells, the CD was lower, also as observed.

The simulations shown in Figure 11, Panel A should be compared to the titration of Figure 3; both show the effects of lowering the expression level of Yfh1. The right-side of the Figure 11, top panel plots represent aerobic WT conditions while the left-side represents aerobic Yfh1-deficient conditions ([EST] = 0). Simulations showed an immediate drop in the CD followed by a decline of Fe\textsuperscript{III}. In actual titrations, Fe\textsuperscript{III} declined before the CD. In the simulated titration, as Yfh1 levels declined, Fe\textsuperscript{II} increased followed by nanoparticles.

The other panels of Figure 11 are plots showing the effect of decreasing oxygen levels on WT and mutant cells. For both strains, simulated [O2] concentrations declined as the oxygen concentration in the growth medium (RO2) declined. For WT cells, this occurs with a shift from Fe\textsuperscript{III} to Fe\textsuperscript{II} (as observed) while the concentration of the CD remained constant at ~30%. In real spectra, the CD contribution is less intense. For simulations of mutant cells, nanoparticles dominated, as observed. As [O2] levels declined, simulations showed that nanoparticles declined as Fe\textsuperscript{II} increased (as observed). Simulated [FS] was low, whereas in reality, the CD in mitochondria isolated from hypoxic cells had an intensity comparable to in WT mitochondria (See Figs. 8 and 9). Encouragingly, simulated [FS] did increase as in silico cells approached hypoxia. ∆Δmrs3/4 simulations still showed the recovery of mitochondrial iron distribution with increasing nutrient iron and a large amount of vacuolar Fe\textsuperscript{III} with respect to WT (Fig. S2).

Given the complexity of the model and the different phenotypes observed for the five states, the simulations overall were qualitatively consistent. They provide significant support for the mechanism of Figure 1 and the major assumptions made in its development.

Discussion

Disentangling the Yfh1 phenotype in yeast is notoriously difficult; it has been the topic of >400 research articles since it was first described in 1997 (52). We used the results of this study, primarily obtained from MB spectra, along with published results from the literature, to develop a chemical/mathematical model (Fig. 1) that can explain this phenotype on the molecular level and in semiquantitative terms. Apart from our related earlier math models (35, 36), no comparable models have been published. Although not perfect, our model represents the best current mechanistic understanding of how healthy yeast cells transition to the diseased state associated with Friedreich’s ataxia.

Changes during Yfh1-titration reveal the cell’s response to Yfh1-deficiency

In the titration reported here, the earliest MB-observable event as Yfh1 expression in AFM cells declined from WT levels was the reduction of vacuolar Fe\textsuperscript{III} to Fe\textsuperscript{II}. This reduction occurred prior to the accumulation of iron in the cell or mitochondria and prior to the loss of [Fe3S4]\textsuperscript{2+} clusters and LS Fe\textsuperscript{II} hemes (as reflected in the CD), much of which is in mitochondria. We were unable to determine whether F2 remained in the vacuole or moved into the cytosol. Nor could we determine whether the observed spectral changes arose...
from changes in the genetic expression of the iron transporters on the vacuolar membrane or from epigenetic effects, e.g., changes in the concentrations of a metabolite that affected the rate of reduction or transport (e.g., NADPH). In any event, these results reveal an independence between the iron status of vacuoles and the activity of Yfh1 in mitochondria; amazingly, vacuolar iron is more sensitive to modest declines in Yfh1 expression than is mitochondrial iron (even though Yfh1 is a mitochondrial protein). One speculative possibility is that the redox properties of the vacuoles are connected to the activity of mitochondria through metabolites such as NAD(P)H. For example, reduced consumption of NADH due to impaired respiration might increase NADH which might be converted to NADPH by dedicated kinases thus promoting reduction of vacuolar FeIII.

Later titration events included the accumulation of cellular iron and the loss of the CD. These two processes occurred synchronously, suggesting a tight mechanistic connection between the rate of cellular iron import (Rcyt), mitochondrial iron import (Rmit), ISC assembly (Rm), and nanoparticle formation (Rnp). Rcyt probably increased when the declining concentration of Yfh1 began to inhibit Rmit. The reduction in [FS] caused by this inhibition probably activated the Aft1/2-dependent iron regulon causing Rcyt to increase. From these results alone, one might conclude that the same regulatory mechanism controls Rmit but the rate of mitochondrial iron import may be regulated differently. Our model assumes that Rnp is controlled by O2 levels in mitochondria and is not genetically regulated.

Seguin et al. estimated that the Yfh1 threshold for activating the iron regulon is ~7% of WT levels (6). Below this, aconitase activity drops precipitously. At [EST] = 2.5 nM, aconitase activity was similar to WT activity, whereas at [EST] = 0 nM, activity was near zero. We observed similar disproportionate behavior in terms of iron accumulation and nanoparticle formation, in that both developed abruptly with a small change in [EST]. The respiratory shield aspect of the Figure 1 model allows such abrupt behavior due to its vicious-cycle nature.

In 2010, Moreno-Cermeño et al. examined a strain of yeast in which Yfh1 could be depleted over time by adding doxycycline to the growth medium (53). Iron accumulated soon after Yfh1 started to deplete. This was followed by a decline in respiration and then a decline of ISC enzyme activities. They concluded that the iron regulon activates and cellular iron accumulates before ISC activity declines suggesting that Yfh1 is not primarily involved in either ISC or heme biosynthesis. In contrast, our results suggest that Yfh1 primarily accelerates the rate of ISC/heme assembly and that the other events associated with a Yfh1 deficiency arise secondarily because of a decline in ISC/heme assembly rates.

**Iron homeostasis partially recovers in Yfh1-deficient cells grown on high nutrient iron**

Unexpectedly, AFM0-100 cells accumulated less iron as nanoparticles than similar cells grown with lower FeIII citrate supplementation. The mitochondrial membrane potential in Yfh1-deficient cells was 20-fold lower than in WT cells, but it could be partially restored by growing Yfh1-deficient cells in media containing high levels of iron (32). Accordingly, we suggest a connection between membrane potential, Rmit, and Rnp. The respiratory shield hypothesis implies such a connection since respiration establishes a proton gradient and hence a membrane potential. We hypothesize that Yfh1-deficient mitochondria in aerobically grown cells synthesize ISCs at a rate that is limited by [FM] and that [FM] is influenced by [FC] (via rate Rmit). We further hypothesize that ISCs are not observed by MB spectroscopy or evidenced by aconitase activity because the high levels of O2 in mitochondria under these conditions inhibit the FM → FS reaction. When aerobic cells are grown in high-iron media, the rate of ISC assembly increases because [FM] increases (due to increased [FC]). This fortifies the respiratory shield in ΔΔmrs3/4 cells grown under iron-replete conditions (36). The observed recovery at high nutrient iron would be difficult to explain without the respiratory shield concept.

**Hypoxic growth reveals that Yfh1 is not required for ISC assembly**

The dominance of the CD in MB spectra of mitochondria isolated from HFM0 and HRM0 cells was typical of WT mitochondria, regardless of whether mutant cells were grown with [IRON] = 1 or 40. This indicates that Yfh1 is not required for ISC synthesis. We cannot distinguish whether ISCs in aerobic Yfh1-deficient cells are synthesized and then degraded by O2 or not synthesized at all (due to inhibition by O2). Previous results suggest the latter (32, 54), and so we assumed that O2 inhibits the FM → FS reaction. Doing so also strengthens the vicious-cycle nature of the respiratory shield.

Yfh1-deficient cells can generate ISCs and hemes when grown under anaerobic/hypoxic conditions (16, 30, 32), and our results support this. ΔΔyfh1 cells display no growth defects when cultured aerobically and ISCs can be assembled (as evidenced from by aconitase activities), though probably at lower rate than for WT cells. These mutant cells have increased levels of oxidative damage when grown aerobically.

ISC synthesis in Yfh1-deficient cells may be more O2 sensitive than WT cells. Yfh1 forms an ISC assembly complex with Nfs1, Isd11, Acp1, and Isu1/2. In that complex, Yfh1 may guide the persulfide on the Nfs1 flexible loop to its destination on a recipient cysteine residue of Isu1/2. The Nfs1 persulfide and the cysteine recipient may be particularly O2 sensitive, leading to a greater requirement for Yfh1 in air versus hypoxic conditions.

Ast et al. recently confirmed the effect of hypoxia on Yfh1-deficient yeast and extended such studies to frataxin-deficient human cells, nematode worms, and mice (54). They reported that frataxin was not required to achieve WT levels of ISC assembly activity under hypoxic conditions. They also found that hypoxia boosts ISC biosynthetic activity rather than suppresses ISC degradation, confirming earlier studies. However, they did not cite any of the earlier pioneering studies on the effects of hypoxia in Yfh1-deficient yeast cells (16, 30, 32).
Cellular iron redistribution in Yfh1-deficient yeast

The effect of hypoxia (and thus O₂) on Yfh1-deficient cells helps disentangle how iron homeostasis is regulated. Our studies show that both O₂ and a deficiency of Yfh1 are required for yeast cells to be iron-dysregulated and exhibit the classic phenotype involving loss of ISC activity, accumulation of FeIII-oxhydroxide nanoparticles in mitochondria, increase in ROS, decline of respiratory ability, etc. The model of Figure 1 describes the interplay between these two factors in mechanistic terms, with the respiratory shield playing an essential role.

Effect of hypoxia in Yfh1-deficient cells is not specific to Yfh1-deficiency

The phenotypes of hypoxic/anaerobic cells deficient in Atm1 or Yah1 are similar to that of Yfh1-deficient cells. As with Yfh1-deficient cells, the iron content of mitochondria isolated from aerobic fermenting Atm1-deficient and Yah1-deficient cells is an order-of-magnitude higher than WT mitochondria (31, 33). Atm1-deficient mitochondria from aerobically grown cells are massively overloaded with nanoparticles and do not exhibit an ISC doublet in their MB spectra. Their aconitase activities are near zero and vacuolar FeIII is absent. Thus, iron is dysregulated in aerobic Atm1-deficient cells. The best-studied mechanism of iron homeostasis is thought to involve Atm1 exporting “X-S” which connects the ISC activity of mitochondria with the Af1/2-dependent iron regulon that controls iron import into the cell and mitochondria. Accordingly, the lack of X-S caused by an Atm1 deficiency upregulates the import of Fe into the cell and mitochondria. Yah1-deficient cells grown aerobically show a similar phenotype indicating iron dysregulation.

However, mitochondria isolated from Atm1- and Yah1-deficient mutant cells grown under anaerobic/hypoxic conditions are not overloaded with nanoparticles. Hypoxic Atm1-deficient cells contain cytochromes at WT levels and have significant aconitase activities. Like Yfh1, Atm1 is not required for mitochondrial ISC cluster assembly or heme biosynthesis—and such clusters are observed if cells are grown under anaerobic/hypoxic conditions. Unlike Atm1 and Yfh1, Yah1 appears to be essential for ISC assembly, since the MB spectrum of isolated mitochondria from these cells, when grown hypoxically, do not exhibit a distinct CD as is observed in mitochondria isolated from hypoxic Atm1- or Yfh1-deficient cells.

The results of Miao et al. for mitochondria isolated from Atm1-deficient cells grown under anaerobic conditions (33) cannot be easily explained by assuming that Atm1 exports X-S in the mechanism of Af1/2-dependent iron regulation (55). (Atm1 likely exports X-S for use in cytosolic ISC assembly). The contents and distribution of Fe in mitochondria isolated from anaerobic WT and Atm1-depleted cells are quite similar suggesting that the homeostatic mechanism regulating mitochondrial iron remains operational even in the absence of Atm1, as long as O₂ is excluded or is present at low concentrations. This suggestion needs to be evaluated further.

The effect of hypoxia in rescuing the nanoparticle accumulation phenotype does not appear specific to a specific protein deficiency. Rather, the curative effect of hypoxia seems to be a general response to a deficiency in various proteins which, we suggest, collectively contribute to a major mitochondrial process such as respiration. The respiratory shield again comes to mind because its function to slow O₂ from diffusing into the mitochondrial matrix depends on the concerted functioning of many proteins. Defects in any of these proteins could weaken the shield, leading to a massive accumulation of nanoparticles under aerobic conditions. The same defects would be present under anaerobic conditions, but in that case, a weakened shield might be sufficient to prevent nanoparticle formation and dysregulation when coupled with the lower oxygen levels associated with hypoxic/anaerobic conditions.

If the rate of cellular iron import (Rcell) were regulated the same as the rate of mitochondrial iron import (Rmit), a change in one rate would always be matched by a similar change in the other. Such matching occurs for Yfh1-deficient cells under aerobic conditions (both rates increase), and it may occur under hypoxic conditions (both rates do not increase).

Other evidence suggests the opposite. The “triple mutant” strain in which Yfh1, Mrs3, and Mrs4 have all been deleted (ΔΔΔ) exhibits behavior suggesting independent regulation of Rcell and Rmit. The iron that accumulates as nanoparticles in Yfh1-deficient mitochondria is imported via Mrs3/4 transporters on the inner mitochondrial membrane. The ΔΔΔ mutant does not accumulate iron in mitochondria but does accumulate iron in the cell (16). This suggests that ΔΔΔ cells are iron-dysregulated (probably due to low [FS]) even though their mitochondria are not, implying independent regulation between cellular iron import and mitochondrial iron import. Under hypoxic conditions, the ΔΔΔ strain grows slowly in unsupplemented media, but faster in high-iron media as more iron enters the mitochondria through low-affinity importers such as Rim2 (56). This behavior is reminiscent of the ΔΔΔΔrs3/4 double-deletion strain in which the respiratory shield was strengthened by an increased Rmit under iron-replete conditions (19).

In our current model, we presumed that Rcyt is regulated by mitochondrial ISC production (i.e., [FS]) while Rmit is regulated by [FM]. This allowed us to explain why mitochondria import massive amounts of iron in Yfh1-, Atm-, Yah1-mutant cells under aerobic but not hypoxic conditions. Under aerobic conditions, both [FS] and [FM] are low and [MP] is high (the mitochondria are filled with nanoparticles), such that Rcyt and Rmit are both activated. [FS] is low because the lack of Yfh1 slows the reaction FM → FS; [FM] is low because the respiratory shield is weakened (due to low [FS]) and the increased O₂ that diffuses into the mitochondria reacts quickly with FM to generate MP. Under hypoxic conditions, the shield weakens but is sufficiently strong to slow the rate of O₂ diffusion into the matrix (since external [O₂] is low). This prevents [FM] from declining (converting to MP) such that Rmit is not activated.
Redox state of vacuolar iron depends on O2 levels in WT cells but on metabolic state in mutant cells

Under both aerobic and hypoxic growth conditions, current and previous studies suggest that mitochondria from fermenting or respiring WT cells are similar in terms of MB spectra (dominated by the CD with some NHHS FeIII). Under aerobic conditions, vacuolar iron is FeIII, while under hypoxic conditions, vacuolar iron in WT cells is FeII (perhaps mobilized from vacuoles to cytosol). This implies an independence between vacuolar and mitochondrial iron.

The corresponding situation was different for the mutant; here, under both aerobic and hypoxic growth, vacuolar iron remained FeIII when cells respired but was reduced to FeII when cells fermented. Mutant mitochondria under hypoxic conditions were qualitatively indistinguishable from WT mitochondria, whereas mutant mitochondria under aerobic conditions were almost certainly overloaded with nanoparticles (we did not isolate mitochondria under this condition in this study but whole-cell MB showed an intense nanoparticle doublet indistinguishable from spectra of mitochondria previously isolated from Yfh1-deficient cells). This implies that redox state (and perhaps mobilization) of vacuolar iron is controlled independently of the regulatory process that controls mitochondrial iron.

The proposed regulatory independence of vacuolar versus mitochondrial iron is also evident in aerobic WT cells grown under [IRON] = 1 versus 40. Under both nutrient conditions, the growth rate of cells is essentially invariant, and mitochondria from both cells are “normal”. By contrast, vacuoles are largely devoid of iron when cells are grown in 1 μM Fe and largely filled when grown in 40 μM Fe. This implies that filling vacuoles (due to μvac in the model) is regulated by something besides or in addition to [FS] activity in mitochondria. In the model, μvac was regulated by both [FC] and [FS]. This dual regulation can also explain why vacuoles are devoid of iron in Yfh1-deficient cells grown under aerobic conditions, in that the increased flow of Fe into mitochondria (due to the decline of [FMJ]) results in low [FC], which then promotes the exodus (or inhibits the import) of iron from/to vacuoles.

Synchronized versus independent iron regulation

Our results and modeling provide some evidence for regulatory independence among the major sites of regulation in the trafficking of iron in yeast cells; these include the site of cellular iron import, mitochondrial iron import, vacuolar iron import/export, and vacuolar redox state.

The extreme possibilities for regulating multiple sites of iron trafficking in cells are global synchronization, regulated by a common sensed species, or complete independence, achieved by separate regulatory processes operating locally on different reactions. A spectrum of intermediate situations in which there is some degree of independence, and some degree of synchronization is also possible. On a molecular level, independence can arise from variations in the strength or kinetics of binding of the same transcription factor to promoter sites of the regulated genes. It may arise from the involvement of multiple transcription factors in controlling regulation. Transcription factors Aft1, Aft2, Yap5, and Cth2 are all involved in iron regulation but are only partially understood at the cellular or systems level. Perhaps, these and other unknown factors are responsible for the degree of regulatory independence that we observed.

Advantages of mathematical modeling

Understanding the development of the frataxin-deficient phenotype requires quantitative mathematical models to integrate myriad interrelated events. Our model represents a quantitative hypothesis regarding the mechanism of iron trafficking and regulation in a yeast cell. Some experiments reported here were used to test the model and determine whether it has predictive power. The model of Figure 1 evolved from earlier textual statements (30), as well as pictorial (6, 31, 33) and mathematical models (35, 36).

Extending this approach to iron metabolism in human cells is a long-term objective. Such a model, with sufficient complexity and realism, may help develop new therapies and better treatments of Friedreich’s ataxia.

Conclusions

The major conclusions of this study are

- The first event to occur as Yfh1 levels decline is a reduction of vacuolar FeIII and an increase in NHHS FeII. Following this is a decline of ISC and rise of mitochondrial nanoparticles.
- Both hypoxic WT and Yfh1-deficient cells lack vacuolar FeIII and nanoparticles; they are dominated by NHHS FeII followed by ISCs.
- Aerobic Yfh1-deficient cells contained higher Mn concentrations than WT cells.
- Most iron in aerobic Yfh1-deficient cells is nanoparticles. Mutant cells grown on higher nutrient iron contained fewer nanoparticles.
- Hypoxic Yfh1-deficient cells are not iron dysregulated.
- The redox state of vacuolar iron is sensitive to the O2 level during cell growth.
- The redox state of vacuolar iron and perhaps the rates of vacuolar iron import/export are independent of the iron regulation/dysfunction status of the mitochondria or whole cells.
- Mitochondria from Yfh1-deficient cells grown hypoxically do not accumulate nanoparticles but contain ISCs. Yfh1 stimulates ISC assembly but is NOT essential for it.
- A developed mathematical model of the kinetics of iron trafficking and regulation in growing yeast cells could explain semiquantitatively the behavior of five different cellular states.
- According to the model, under healthy conditions, respiration slows the diffusion of O2 into the matrix. Yfh1 deficiency slows the rate of ISC assembly, allowing more O2 to enter the matrix and reacts with the FeII pool to generate nanoparticles. O2 also inhibits ISC assembly, creating a vicious cycle that culminates in the diseased state.
**Cellular iron redistribution in Yfh1-deficient yeast**

- Assuming that global cellular regulation is controlled by the
  ISC activity in the mitochondria was less able to reproduce
  observed behavior than assuming local independent regulation.

**Experimental procedures**

**Yeast strains and cell growth**

ERyfh1 cells were constructed from the YPH499 parent strain by promoter swap in which the YFH1 open reading frame was placed under control of the GAL1 promoter. The plasmid pGEV4-LEU2 was inserted and GALA4 was deleted, thereby creating a strain in which the YFH1 open reading frame was placed under control of an estradiol regulated promoter (MATa ura3-52, lys2-801 (amber) ade2-101 (ochre) trpl-A63 leu2-A1 cyh2 Δgal4::KanMX6 HIS3MX6-pGAL1-YFH1[pGEV-LEU2] (6, 41, 57). Cells were maintained on MM agar plates without leucine to select for the pGEV plasmid and with 5 μM estradiol to maintain Yfh1 expression. For some batches, cells were inoculated into liquid minimal media lacking leucine but containing estradiol at various concentrations. For other batches, complete synthetic medium (Sunrise Science) medium lacking leucine was used. For hypoxic growths, 50 ml of aerobically grown cultures were used to inoculate larger cultures that had been supplemented with ergosterol (20 mg/L), Tween-80 (1 ml/L), and antifoam B (Sigma Aldrich, 100 ppm final concentration) (31, 33). Cultures were vigorously bubbled with a 95:5 N2:O2 or 3:1 N2:air gas mixture to render them hypoxic. Whole cell cultures were harvested by centrifugation at 5000g for 5 min. Cells were washed 3x in a solution of 100 μM EDTA, and then 3x in distilled, deionized water. Cells were packed into either a MB cup or EPR tube by centrifugation and frozen in LN2.

**MB and EPR spectroscopies**

MB spectra were collected on a MS4 WRC spectrometer (SEE Co, Edina MN) at 5 to 6 K and 0.05 T. The magnetic field was applied parallel to the gamma radiation. An α-iron foil was used for RT calibration. X-band EPR spectra were collected using a Bruker Elexsys E500 spectrometer.

**Mitochondria isolations**

Cells were harvested during exponential phase from a custom-built 24 L glass/titanium bioreactor. For hypoxic growths, two bioreactors were grown in parallel. Upon harvesting, liquid culture was pumped into a refrigerated anaerobic glove box for collection into centrifuge bottles. Filled bottles were removed from the box, centrifuged, and returned to the box where pellets were combined. Mitochondria were purified anaerobically from pellets as described (19). Organelles were frozen in MB cups, EPR tubes, or Eppendorf tubes for later analysis.

**Metal analysis and aconitase assays**

Metal analyses were performed as described (58) using 5% trace metal–grade nitric acid; 2.5% H2O2 was added to samples as needed. Aconitase assays were performed essentially as described (58). Isolated mitochondria were lysed, and their protein contents were determined by the BCA assay. Approximately, 16 to 32 μg of mitochondrial protein in a lysis solution was added to reaction buffer containing 20 mM NaCl, 20 mM Tris-HCl, and 1 mM cis-aconitic acid (final concentrations). The decline in absorbance of the resulting solution at 240 nm was monitored for 6 min, and the slope was used to calculate activity (1 unit = 1 μmol cis-aconitic acid/min).

**Western blot**

The blot was performed essentially as described (13). Crude mitochondria were isolated as above except without the ultracentrifugation step. The frataxin primary antibody was mixed in the TBST:milk solution at a concentration ratio of 1:100. The ratio for the porin antibody was 1:500. Samples were incubated 1 h at RT and then overnight at 4 °C. Secondary antibody was prepared at a concentration ratio of 1:500 for frataxin and 1:2500 for porin, then incubated 2 h at RT and monitored by chemiluminescence (13).

**Data availability**

All data are contained within the manuscript and SI.

**Supporting information**—This article contains supporting information (35).

**Author contributions**—S. F., J. D. W., and R. E. S. investigation; S. W. V. formal analysis; J. D. W. methodology; A. D. resources; A. D., P. A. L. conceptualization. P. A. L. funding; P. A. L. supervision; P. A. L. writing-original draft; S. F., J. D. W., S. M. V., A. D., P. A. L., and R. E. S writing-review and editing.

**Funding and additional information**—This work was supported by the National Science Foundation, United States (MCB-1817389), the National Institutes of Health, United States (GM127021 to P. A. L. and GM107542 to A. D.), and the Robert A. Welch Foundation, United States (A1170). The content is solely the responsibility of the authors and does not necessarily represent the views of the National Institutes of Health, the NSF, or the Welch Foundation.

**Conflict of interest**—The authors declare that they have no conflicts of interest with the contents of this article.

**Abbreviations**—The abbreviations used are: α, exponential growth rate of cells; CD, central quadrupole doublet; CIA, cytosolic iron not associated with FC; ΔΔΔΔ, strain of yeast in which Yfh1, Mrs3, and Mrsc4 have been deleted; EPR, electron paramagnetic resonance; EST, estradiol; F2, vacuolar FeII; F3, vacuolar FeIII; FC, cytosolic iron; FM, mitochondrial FeII pool; FS, iron-sulfur clusters and FeII hemes, many used in respiration; ISC, iron–sulfur cluster; MB, Mössbauer; MP, mitochondrial nanoparticles; N, nutrient iron (FeIII citrate); NHHS, non-heme high spin; ODE, ordinary differential equation; Reg±(S), regulatory functions exhibiting positive or negative feedback, sensing species S; ROS, reactive oxygen species; Yfh1, Yeast frataxin homolog 1; VP, vacuolar nanoparticles; WT, wildtype.
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Cellular iron redistribution in Yfh1-deficient yeast
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