Vacuoles are acidic organelles that store Fe\textsuperscript{III} polyphosphate, participate in iron homeostasis, and have been proposed to deliver iron to mitochondria for iron–sulfur cluster (ISC) and heme biosynthesis. Vma2\textDelta cells have dysfunctional V-ATPases, rendering their vacuoles nonacidic. These cells have mitochondria that are iron-dysregulated, suggesting disruption of a putative vacuole-to-mitochondria iron trafficking pathway. To investigate this potential pathway, we examined the iron content of a vma2\textDelta mutant derived from W303 cells using Mössbauer and EPR spectroscopies and liquid chromatography interfaced with inductively-coupled-plasma mass spectrometry. Relative to WT cells, vma2\textDelta cells contained WT concentrations of iron but nonheme Fe\textsuperscript{II} dominated the iron content of fermenting and respiring vma2\textDelta cells, indicating that the vacuolar Fe\textsuperscript{III} ions present in WT cells had been reduced. However, vma2\textDelta cells synthesized WT levels of ISCs/hemes and had normal aconitase activity. The iron content of vma2\textDelta mitochondria was similar to WT, all suggesting that iron delivery to mitochondria was not disrupted. Chromatograms of cytosolic flow—through solutions exhibited iron species with apparent masses of 600 and 800 Da for WT and vma2\textDelta, respectively. Mutant cells contained high copper concentrations and high concentrations of a species assigned to metallothionein, indicating copper dysregulation. vma2\textDelta cells from previously studied strain BY4741 exhibited iron-associated properties more consistent with prior studies, suggesting subtle strain differences. Vacuoles with functional V-ATPases appear unnecessary in W303 cells for iron to enter mitochondria and be used in ISC/heme biosynthesis; thus, there appears to be no direct or dedicated vacuole-to-mitochondria iron trafficking pathway. The vma2\textDelta phenotype may arise from alterations in trafficking of iron directly from cytosol to mitochondria.

This article contains supporting information.

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that are deficient in yeast frataxin homolog 1 (Yfh1) (19), yeast adrenodoxin homolog 1 (Yah1) (20), and the inner membrane transporter Atm1 (21). These cells respond to iron regulon activation by importing excess iron and mobilizing vacuolar iron stores.

The iron regulon is activated in vma2Δ cells grown in iron-replete (μM range) media even though their mitochondria are not deficient in proteins involved in ISC assembly (22, 23). This implies that mitochondria cannot assemble sufficient ISCs when the vacuolar lumen is insufficiently acidic, and it highlights an intriguing connection between the two organelles. vma2Δ cells have reduced activities of mitochondrial ISC-containing proteins that are involved in respiration, including aconitase, succinate dehydrogenase, and the Rieske iron–sulfur protein (23). vma-mutant cells grow poorly on respiring media (3, 22, 24, 25). However, supplementing growth media with mM concentrations of iron a) rescues this phenotype, b) deactivates the iron regulon (22, 23), c) increases the activity of ISC-containing enzymes, d) increases the rate of O2 consumption, and e) stimulates respiration (24–26).

Excess supplemented iron provides the additional iron needed for the biosynthesis of iron-rich mitochondria which are in turn needed for respiration. This requirement is uncompromising, even in WT cells, such that cells grown on iron-deficient minimal medium (MM) succumb to autophagy so that they can recycle sufficient cellular iron for use in mitochondriogenesis (27). During autophagy, Smf3 and Fet5/ Fth1 iron exporters on the vacuolar membrane pump iron into the cytosol to satisfy mitochondrial demand. Iron-deficient vma mutants have difficulty undergoing autophagy which hinders respiration (27).

When the iron regulon is activated in ISC-mutant cells, the rate of cellular iron import increases dramatically owing to expression of a high-affinity iron importer (composed of Fet3 and Ftr1) on the plasma membrane. Expression of the high-affinity vacuolar iron exporter (composed of Fet5 and Fth1) also increases. Fet3 and Fet5 are paralogous multicopper oxidases controlled by the iron regulon. As a result, the vacuoles of ISC-mutant cells are nearly devoid of iron, and much of the excess iron that flows into the cytosol ultimately accumulates in mitochondria as Fe3+ oxyhydroxide phosphate–associated nanoparticles (19–21).

vma-mutant cells have difficulty importing extracellular iron because copper-containing Fet3 has difficulty maturing (i.e., metallating with copper) in the Golgi (1, 28). Whether these cells also have difficulty exporting vacuolar iron is unknown, but this seems likely, given the similarities between Fet3 and Fet5. In any event, the concentration of iron in vma-mutant cells is not excessive relative to WT cells even though the iron regulon is activated. vma2Δ cells reportedly contain only ~30% more iron than WT cells (23). In contrast, ISC mutants contain an order-of-magnitude more iron than WT cells.

The mechanism by which a decline in the acidity of vacuoles causes a decline of mitochondrial function, and how excessive iron rescues this decline, has been investigated by Kane (23) and more recently by Hughes (26). Both hypothesize that the flow of iron from vacuoles to mitochondria is disrupted in vma-mutant cells and that excess media iron re-establishes this pathway. Besides sequestering iron, vacuoles store amino acids, including cysteine (29). Hughes et al. (26) found that loss of vacuolar acidity increases nonvacuolar pools of cysteine and other amino acids. One intriguing possibility is that excess cysteine in the cytosol binds iron, thereby blocking import into mitochondria.

vma mutants generate high levels of reactive oxygen species (ROS) and are hypersensitive to oxidative stress (5, 22, 30, 31). They are more easily intoxicated by high levels of transition metals than WT cells. This likely occurs because vma cells have difficulty storing metals in their vacuoles, which otherwise imparts metal resistance to the cell (17, 22, 24, 32, 33). Surprisingly, mitochondrial respiration is not the major source of ROS in vma2Δ cells (22). Rather, excessive ROS is likely associated with iron dyshomeostasis and increased Fenton chemistry (5, 23).

Also surprising is the relationship between vacuolar acidity, cellular iron, mitochondrial function, and ROS damage is related to human aging and longevity. Hughes and Gottschling found that a decline of vacuolar acidity is associated with age-related deterioration of mitochondria (34). Overexpressing V-ATPase subunits suppresses mitochondrial dysfunction in aging cells by hyperacidifying vacuoles. Young yeast cells treated with a V-ATPase inhibitor and mutant cells lacking V-ATPase subunits both have impaired mitochondrial morphology similar to the organelle in aged cells (34).

The objective of the current study was to better understand the connection between vacuolar acidity, mitochondrial function, and iron homeostasis by using powerful biophysical and bioanalytical probes, namely Mössbauer (MB) and EPR spectroscopies and liquid chromatography interfaced with inductively-coupled-plasma mass spectrometry (LC-ICP-MS). Mitochondria, vacuoles, and the cytosol of vma2Δ cells were isolated and investigated. Our results confirm and extend some but not all relevant previous results, with differences probably due to the background strains and media/conditions used. New insights into the connection between vacuoles and mitochondria are presented, highlighting the potential importance of low-molecular-mass (LMM) iron, copper, and sulfur species in the cytosol of these yeast cells.

Results

A haploid VMA2 deletion strain was generated from a W303 parent to allow for direct comparison to previous biophysical studies that used the same genetic background (10, 11, 20, 21, 35–39). We refer to W303 and the mutant strain as WT and vma2ΔW, respectively. We initially investigated whether the iron content of these two strains differed and whether the pH of the medium influenced that content. WT and vma2ΔW cells were grown in MM containing 2% glucose as the carbon source (fermentation) and supplemented with iron (typically 40 μM 57FeIII citrate). Medium pH was buffered using citric acid and sodium citrate. Under these conditions, vma2ΔW cells grew well at pH values 3, 4, and 5 although growth was inhibited when the medium pH was 6 or 7; in contrast, WT cells grew well in media buffered from pH 3 to 7.
Whole-cell iron concentrations (Table 1) revealed no obvious pH dependence; however, some strain-dependent trends were evident. *vma2Δw* cells contained 30% less iron, 60% less phosphorus, 6× less manganese, 4× less zinc, and 2.5× more copper than WT cells. The increased copper when considered with previous reports that Fet3 in *vma2Δ* cells contained 30% less iron, 60% less phosphorus, 6× less manganese, 4× less zinc, and 2.5× more copper than WT cells. The increased copper when considered with previous reports that Fet3 in *vma2Δ* cells is dysregulated in these mutant cells and that *vma2Δ* cells compensated by importing excessive copper.

MB spectra of both strains were collected to investigate the type of iron present. WT cells exhibited qualitatively similar spectra regardless of pH (Fig. 1), but some subtle pH-dependent trends were evident. At low pH, over 60% of spectral intensity was a magnetic feature originating from nonheme high-spin (NHHS) FeIII. The solid purple line in Figure 1 is a simulation using parameters in Table S1. The FeIII ions are coordinated to anionic polyphosphate chains in vacuoles (10, 11). With increasing pH, the percentage of cellular iron in this form declined to ~30% at pH 7. We conclude that WT vacuoles store less iron as medium pH increases.

Also evident in WT MB spectra was the central doublet (CD) with isomer shift and quadrupole splitting parameters typical of [Fe4S4]2+ clusters and low-spin FeII hemes (simulated by the green line in Fig. 1). MB spectra of isolated mitochondria are dominated by the CD (36–38), which originates mainly from iron-rich respiratory complexes and respiration-related iron-containing proteins, e.g., aconitase. A substantial portion of the CD intensity in whole cells is due to cytosolic [Fe4S4]2+-containing proteins. Percentagewise, the spectral intensity of the CD in whole-cell WT spectra increased from ~15% at low pH values to ~30% at pH 7. This increase likely arose from the percentagewise decrease of vacuolar iron, such that the absolute concentration of the [Fe4S4]2+ clusters and LS FeII hemes that give rise to this spectral feature were fairly constant with changes in pH.

Another ~20% of the intensity of WT cell spectra was a quadrupole doublet (simulated by the blue line in Fig. 1) with isomer shift and quadrupole splitting parameters typical of [Fe4S4]2+ clusters and low-spin FeII hemes (simulated by the green line in Fig. 1). MB spectra of isolated mitochondria are dominated by the CD (36–38), which originates mainly from iron-rich respiratory complexes and respiration-related iron-containing proteins, e.g., aconitase. A substantial portion of the CD intensity in whole cells is due to cytosolic [Fe4S4]2+-containing proteins. Percentagewise, the spectral intensity of the CD in whole-cell WT spectra increased from ~15% at low pH values to ~30% at pH 7. This increase likely arose from the percentagewise decrease of vacuolar iron, such that the absolute concentration of the [Fe4S4]2+ clusters and LS FeII hemes that give rise to this spectral feature were fairly constant with changes in pH.

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The WT EPR spectrum of whole WT cells cultured at pH 5 with 40 μM supplemented FeIII citrate (Fig. 2A) exhibited signals similar to those observed previously (11, 38). Vacuolar FeIII exhibited a prominent g = 4.3 signal. Features at g = 6.5 and 5.5 are typical of high-spin FeIII hemes. A hyperfine-split signal arising from MnII ions was observed at g = 2.00, as was an isotropic radical signal. No signals attributable to CuII ions were observed.

MB spectra of vma2ΔW cells grown in MM at pH 3, 4, and 5 were dominated by an FeII quadrupole doublet (Fig. 3) that accounted for ~60% of spectral intensity. The high-energy line of the CD was also evident as a partially resolved shoulder on the low-energy line of the dominating FeII doublet at ~ + 1 mm/s. Like WT cells, the iron content of vma2ΔW cells was generally invariant in media in which acidity varied by three orders of magnitude. We had predicted that the magnetic feature due to vacuolar FeIII in MB spectra of WT cells would also be present in spectra of vma2ΔW cells grown at pH 3 or 4, given that the acidity of vacuoles lacking functional V-ATPase is affected by the pH of the growth media. This prediction was not realized, possibly for reasons discussed below.

The CD represented ~20% of the spectral intensity for vma2ΔW cells at each pH within this range. Assuming an average cellular iron concentration of 180 μM (Table 1) suggests that ~40 μM iron in these cells arose from [Fe4S4]2+ clusters and LS FeII hemes. For WT cells, with an average iron concentration of ~250 μM and ~20% CD intensity, the corresponding concentration would be ~50 μM. The two concentrations are the same within our uncertainties. We had expected that the concentration of such centers in vma2ΔW cells would have been much lower and that most of the iron in these cells would have been FeIII phosphate–associated nanoparticles as is observed in other ISC-mutant cells in which the iron regulon is activated. However, MB spectra of vma2ΔW cells did not exhibit intense features typical of nanoparticles nor did they provide clear evidence that the iron regulon was activated.

EPR spectra of vma2ΔW cells at pH 5 (Fig. 2, B–C) supported the MB spectra and provided new insights. The intensity of the g = 4.3 signal due to vacuolar FeIII was strongly reduced relative to in WT spectra, consistent with the absence of the magnetic feature due to vacuolar FeIII in corresponding MB spectra. Minor features attributed to high-spin hemes at g = 6.5 and 5.5 were comparable in intensity to those of WT, suggesting normal levels of cellular hemes in the mutant cells. The hyperfine-split signal due to MnII ions was dramatically reduced in vma2ΔW EPR spectra, consistent with the 6-fold decline of Mn concentration relative to in WT cells (Table 1). No signals attributable to S = ½ CuII ions were observed, despite the 2.5-fold increase in Cu concentration in...
the mutant cells. This suggests that the additional Cu ions in \(vma2\Delta_{W}\) cells were in the reduced diamagnetic Cu\(^{1}\) state.

We assessed whether the dominating Fe\(^{II}\) species in \(vma2\Delta_{W}\) cells arose from an accumulation of LMM iron complexes in the cytosol. The cytosol was isolated from WT and \(vma2\Delta_{W}\) cells, and the concentration of iron in these cytosol isolates was determined (Table 1). The iron concentration of cytosol from both strains was high (200, 270 \(\mu\)M), similar to values reported previously (38). Cytosol solutions were passed through a 10-kDa cutoff membrane, and the metal contents of flow-through solutions (FTSs) were also determined. We previously found that LMM iron species constituted ca. 70% of cytosolic iron at 40 \(\mu\)M iron supplementation (38). Here, we observed that ~30% of cytosolic iron from cells grown with 1 \(\mu\)M Fe was present in an LMM form (Table 1). The concentration of LMM iron in the cytosol was higher than expected by an order-of-magnitude. We suspected that iron-rich vacuoles may have burst during cytosol isolation and metals from the ruptured organelles leached into the cytosol. However, vacuoles contain little iron at 1 \(\mu\)M and 40 \(\mu\)M iron-supplemented WT cells were similar. Thus, vacuole-bursting seems unable to explain the high concentrations.

FTSs were subjected to LC-ICP-MS chromatography. The resulting traces (Fig. 4) revealed unresolved LMM iron-detected peaks for each strain. The dominant LMM iron species in WT cytosol FTS migrated with an apparent mass of ~600 Da (all masses quoted in the text are apparent), whereas the dominant LMM iron species in \(vma2\Delta_{W}\) cytosol FTS migrated with a mass of ~800 Da. Both species were present in the FTS of both strains, and their collective intensities were similar. We conclude that the LMM iron in the cytosolic FTS of \(vma2\Delta_{W}\) and WT cells are similar though not identical in terms of species and concentration. We caution readers that about half of the LMM iron in the cytosolic fraction adsorbed onto the column.

We considered that much of the NHHS Fe\(^{II}\) in \(vma2\Delta_{W}\) cells was located in the mitochondria, and so this organelle was isolated from both WT and \(vma2\Delta_{W}\) strains (grown on fermenting MM at pH 5). MB spectra of isolated mitochondria from each strain (Fig. 5) were dominated by the CD (ca. 80% intensity). A minor NHHS Fe\(^{II}\) doublet (15%–20%) was also evident. Aconitase activities in these mitochondrial lysates were similar (WT = 2.6 ± 1.2 units/mg protein, \(vma2\Delta_{W}\) = 2.3 ± 1.2 units/mg protein; n = 2). Little if any Fe\(^{III}\) oxyhydroxide nanoparticles were present in \(vma2\Delta_{W}\) mitochondria, unlike mitochondria from strains in which ISC assembly is defective. In summary, the iron contents of mitochondria isolated from WT and \(vma2\Delta_{W}\) cells, as evaluated by MB spectroscopy and by the activity of an ISC-containing enzyme, were similar. The low-intensity NHHS Fe\(^{II}\) doublet observed in WT and \(vma2\Delta_{W}\) mitochondria indicate that mitochondrial Fe\(^{II}\) alone cannot account for the dominating Fe\(^{II}\) intensities observed in \(vma2\Delta_{W}\) whole-cell MB spectra.

We wondered whether the metabolic state of \(vma2\Delta_{W}\) cells (fermenting versus respiring) affected the iron content of the cells. To address this, WT and \(vma2\Delta_{W}\) cells were grown on a respiring carbon source (glycerol/ethanol) in MM buffered at pH 5 and supplemented with 40 \(\mu\)M \(^{57}\)Fe\(^{III}\) citrate. In this case, \(vma2\Delta_{W}\) cells required prior growth in glucose-based media before switching to respiring MM. About 65% of the intensity of the resulting MB spectrum of WT cells (Fig. 6B) arose from vacuolar high-spin Fe\(^{III}\) (10, 11); another ca. 25% arose from the CD. The higher intensity observed for the CD confirms that respiration-related ISC proteins are upregulated during respiration (36). In contrast, the MB spectrum of respiring \(vma2\Delta_{W}\) cells (Fig. 6D) lacked the magnetic feature arising from vacuolar Fe\(^{III}\) but exhibited an Fe\(^{II}\) doublet representing ~40% of spectral intensity. Over half of the spectral intensity.
Mössbauer and LC-ICP-MS of vma2Δ yeast cells

arose from the CD combined with unresolved iron in the middle of the spectrum which fit to a doublet with the parameters of nanoparticles. The cellular concentrations of [Fe₃S₄]- and LS Fe³⁺ heme-containing species in respiring WT and vma2Δ W cells were again similar, as observed for fermenting cells. In summary, the iron content of respiring vma2Δ W cells was similar to that of WT cells, though with a dominance of Fe²⁺ instead of vacuolar high-spin Fe³⁺. The simplest interpretation is that vacuolar Fe³⁺ in WT cells are converted to NHHS mononuclear Fe²⁺ and Fe³⁺ nanoparticles in fermenting and respiring vma2Δ W cells.

We considered that some of the dominant NHHS Fe²⁺ species in iron-replete vma2Δ W cells were present in vacuoles, and so two batches of vacuoles from vma2Δ W cells were isolated and FTSs were prepared. LC traces of these solutions exhibited LMM iron peaks that were either of similar intensity as peaks from WT vacuoles — or less intense (data not shown). However, these studies used a column that was not treated such that adsorption of iron was considerable. We considered that vma2Δ W vacuoles contained additional iron that adsorbed onto the column, but the iron concentration of those vacuoles was not unusually high.

Cells grown in media containing low concentrations of iron do not store much of the metal in their vacuoles. Thus, the contribution of vacuolar iron to MB spectra can be largely eliminated by growing cells under these conditions. The MB spectrum of respiring WT cells grown in MM supplemented with 1 μM ⁵⁷Fe citrate (Fig. 6A) was noisy because the cells contained only 180 μM Fe. However, the S/N was sufficient to conclude that the vacuolar Fe³⁺ feature was essentially absent (<15% of spectral intensity). The corresponding spectrum of vma2Δ W cells, containing 220 μM Fe (Fig. 6C), also lacked the vacuolar Fe³⁺ feature and also exhibited an intense CD (even stronger than for WT cells). This indicates that vma2Δ W mitochondria were fully able to generate [Fe₄S₄] clusters (and LS Fe³⁺ hemes). We conclude that vma2Δ W mitochondria were able to import cytosolic iron and were able to use it for ISC assembly even though vacuoles were largely devoid of iron. This behavior was unexpected, given previous results which indicated a strong need for functional acidic vacuoles to deliver iron to mitochondria during mitochondriogenesis associated with respiration, as well as with the reported rescue of growth of vma-mutant cells with high (mM) concentrations of nutrient iron (22, 24, 26). In summary, these experiments performed with only 1 μM added iron gave essentially the same result in vma2Δ W cells, showing a normal ISC pool in cells where vacuoles did not store significant iron.

To further gauge the vacuolar iron content of WT and vma2Δ W cells, we cultured such cells in high (mM) concentrations of nutrient iron and monitored growth rates. As yeast lack a means of exporting iron, vacuolar iron storage typically provides resistance to excessive iron levels. WT cells grew rapidly until 20 to 30 mM Fe³⁺ citrate had been added to the growth medium, beyond which the growth rate declined (Fig. 7, triangles). vma2Δ W cells grew slower than WT in medium containing up to 10 mM iron and grew poorly in media containing higher iron concentrations (Fig. 7, circles). We suspect that under these high-iron conditions, vma2Δ W cells were unable to sequester toxic iron in their vacuoles in a nontoxic form.

The shapes of the plots of Figure 6 suggested that we could simulate growth rates using Equation 1:

\[ \alpha = \alpha_{\text{cyt}} \left( \frac{1}{1 + \left( \frac{[Fe]_\text{cyt}}{K_\alpha} \right)^{5/2}} \right) + \alpha_{\text{vac}} \]  

which assumes two contributions to the growth rate, one that is sensitive to excessive iron in the media (\( \alpha_{\text{cyt}} \)) and another that is not (\( \alpha_{\text{vac}} \)). The observed experimental growth rate \( \alpha \) was the slope of the ln(OD600) versus time in the exponential growth region. This empirical equation may only apply to the range of media iron concentrations investigated. It implies that

Figure 6. Mössbauer spectra of respiring WT (A–B) and vma2Δ W (C–D) cells grown on media supplemented with 1 (A and C) and 10 (B and D) μM ⁵⁷Fe³⁺ citrate and 10 μM copper sulfate and at pH 5.

Figure 7. Plot of the exponential growth rate \( \alpha \) of fermenting WT (red triangles) and vma2Δ W (blue circles) versus the concentration of Fe³⁺ citrate added to the growth medium. Points are the averages of 2–3 independent batches; bars represent standard error. The solid red line is a simulation using Equation 1 and \( \alpha_{\text{cyt}} = 0.185 \) h⁻¹, \( \alpha_{\text{vac}} = 0.176 \) h⁻¹, and \( K_\alpha = 0.15 \) mM. The solid blue line is the same except \( \alpha_{\text{vac}} = 0 \).
excess cytosolic iron is toxic, whereas iron sequestered in the vacuole is not, as least in WT cells. The red line in Figure 7 simulates the growth rate of WT cells in which vacuoles sequester iron from the cytosol. The blue line simulates the growth of vma2Δw cells in which vacuoles are presumed unable to sequester iron in a benign form such as FeIII polyphosphate.

vma2Δw cells grew more slowly than WT cells, even in media that was not supplemented with iron at high concentrations. Our MB studies indicate that these cells accumulated FeIII ions. We considered that the FeIII species in vma2Δw cells might be susceptible to Fenton chemistry and that this caused ironmentation on the WT and enriched samples used to generate spectra in Figure 2 with formation on other elements, and so we also monitored Cu, media had no noticeable effect on iron in WT cells. In contrast, the presence of high or low phosphate in the growth media may have been limited by phosphate rather than iron. In vma2Δw cells, the formation of nanoparticles suggests that in vma2Δw cells inhibi vma2Δw cells on fermenting MM (pH 5) supplemented with 1 mM cysteine. The MB spectra of such cells were unaffected by cysteine supplementation (Fig. 3B) except for a modest decline in the CD (22% in the spectrum of unsubsupplemented vma2Δw cells down to 14% for the same cells supplemented with cysteine). Hughes et al. (26) suggested that cysteine supplementation caused “bioavailable” iron to decline and n vacuolar cysteine to accumulate. The MB spectrum of the cysteine-supplemented vma2Δw sample did not show an increase in FeIII nanoparticles, a form of iron considered to be bio unavailable.

Mn and Zn chromatograms of vma2Δw and WT cytosol (Fig. 10, A–B) were like those described previously (38), including a single Zn peak at 700 Da and a single Mn peak at 200 Da. Phosphorus chromatograms (Fig. 10, C–F) exhibited a major peak at 400 Da and a shoulder at 600 Da. The intensities of Mn, Zn, and P peaks were significantly lower in vma2Δw traces, consistent with the lower concentration of these elements in vma2Δw cells compared to WT cells. The lack of a CuII-based EPR signal in vma2Δw cells is consistent with diamagnetic CuI binding to metallothioneins. Also present were ca. 2 low-intensity Cu peaks in the range of 500 to 700 Da; their intensities were similar for both vma2Δw and WT FTSs (Fig. 8, insets).

LMM sulfur–based LC peaks were also detected in cytosolic FTSs from both WT and vma2Δw cells (Fig. 9). The dominant sulfur peak in the FTS from both cells (Vw = 20 ml) originated from the sulfur-containing MES buffer used early on in isolating the cytosol. The right shoulder of this peak coeluted with numerous sulfur standards including but not limited to cysteine (Fig. 9F). In vma2Δw cytosol FTS, the intensity of the shoulder was greater relative to that of WT FTS. The sulfur peak at an apparent mass of 100 Da originated from the PMSF added to the buffer during isolation; the sulfur peak at an apparent mass of 50 Da originated from the DTT used earlier during isolation.

Hughes et al. (26) recently suggested that increased concentrations of cytosolic cysteine in vma2-mutant cells inhibited the vacuole-to-mitochondria iron trafficking pathway. To examine this, we grew vma2Δw cells on fermenting MM (pH 5) supplemented with 1 mM cysteine. The MB spectra of such cells were unaffected by cysteine supplementation (Fig. 3B) except for a modest decline in the CD (22% in the spectrum of unsubsupplemented vma2Δw cells down to 14% for the same cells supplemented with cysteine). Hughes et al. (26) suggested that cysteine supplementation caused “bioavailable” iron to decline and n vacuolar cysteine to accumulate. The MB spectrum of the cysteine-supplemented vma2Δw sample did not show an increase in FeIII nanoparticles, a form of iron considered to be bio unavailable.
elements in mutant cells and isolated cytosol (Table 1). Notably, only the P peaks at 400 Da declined in intensity for vma2Δ W cytosol; the peak at 600 Da remained the same. The peak at 70 Da comigrated with an AMP standard.

We were puzzled that many of the results obtained using vma2Δ W cells differed from published results on what was ostensibly the same mutant. However, the strain used to generate the mutant in those reports was BY4741, whereas ours was W303. We obtained the mutant in the BY4741 strain (to be called vma2Δ B) and collected MB spectra of those cells grown at different pHs in fermenting MM supplemented with 40 uM ⁵⁷Fe citrate. vma2Δ B cells grown at low pH exhibited spectra that were virtually indistinguishable from WT cells, including an intense magnetic feature due to vacuolar Fe³⁺ (Fig. 11E versus Fig. 11B). At higher pH, vma2Δ B cells exhibited spectra dominated by a broad quadrupole doublet due to Fe³⁺ nanoparticles (Fig. 11C–D). Also, the percent absorbance was higher than in WT spectra, similar to that observed for ISC-mutant cells with an activated iron regulon (19–21). The spectra of vma2Δ B cells grown at intermediate pH contained a stronger-than-WT NHHS Fe II doublet (Fig. 11D), with parameters similar to those used to simulate the FeII doublet in vma2Δ W cells.

We attributed the differences between our results and those in the literature to differences between the W303 and BY4741 background strains. Notably, our experiments had not indicated an iron trafficking defect in the vma2Δ W cells. Prompted by a reviewer’s comment, we sought to determine by Western blot of whole-cell extracts whether the iron regulon in vma2Δ W cells was activated by monitoring the relative abundance of the high-affinity iron importer Fet3. Cytoplasmic Pgk1 was a positive control. We did not observe a significant difference in Fet3 abundance between WT W303 and vma2Δ W cells (Fig. 12, top row, B and C respectively), indicating that the iron regulon is not activated. In contrast, Fet3 was more abundant in vma2Δ B versus WT BY4741 cells (Fig. 12, top row, E versus D respectively), supporting previous studies indicating that the iron regulon is activated in vma2Δ B cells grown in yeast extract peptone adenine dextrose medium (YPAD) media (22, 23).

Discussion

As described in the Introduction, there is significant evidence for an iron trafficking pathway from vacuoles to mitochondria in yeast. Vacuoles store iron that is needed for mitochondrial biogenesis especially when cells shift from fermentation to respiration. Nonacidic vacuoles with defects in V-ATPase have been reported to generate mitochondrial defects in ISC synthesis and iron homeostasis. Our initial objective was to probe this relationship and defects in vma2Δ W cells by

**Figure 10.** Zn-, Mn-, and P-detected chromatograms of cytosol FTS from WT (A–C) and vma2Δ W (D–F) cells. Each displayed Zn (A and D), Mn (B and E), and P (C and F) trace is the average of two traces from independent batches. FTS, flow-through solution.

**Figure 11.** Mössbauer spectra of fermenting BY4741 (A–B) and vma2Δ B (C–E) cells grown in media buffered at the following pH values. A, 6; B, 3; C, 5; D, 4; E, 3.
MB and EPR spectroscopies and LC-ICP-MS chromatography. However, analyzing our results was more challenging than anticipated.

One result was clear: most of the iron in fermenting *vma2Δ* cells was nonheme high-spin FeII. In W303 cells, approximately the same percentage of iron was HS FeIII, previously assigned to vacuolar iron. The overall iron concentrations in WT and *vma2Δ* cells and the spectral contributions of the CD (due to [Fe4S4]2+ clusters and LS FeII hemes) were also similar. We conclude that vacuolar FeIII ions in WT cells are reduced to the FeII state in *vma2Δ* cells. We suspect but are not certain that the FeII remains in the organelle. We have observed a similar FeIII → FeII reduction of vacuolar iron in yeast cells grown on adenine-deficient media in which metabolism has been altered (39). Some vacuolar iron in iron-replete respiring *vma2Δ* cells was reduced to FeII (and some may have formed FeIII nanoparticles).

The reduction of FeIII to FeII as the acidity of vacuoles declines can be partially or wholly explained by the conclusions of Raguzzi et al. (40) and Singh et al. (41) using insights from Schäfer and Buettner (42). These researchers hypothesized that the glutathione redox couple \( \{GSSG + 2H^+ + 2e^- \rightleftharpoons 2GSH\} \) controls the redox status of vacuoles in accordance with the Nernst equation:

\[
E_{GSH}^{\text{pH}(H^+)} = E_{GSH}^{\text{pH}(H^+)} - \frac{59}{2} \log \left( \frac{[GSH]^2}{[GSSG] \left[\frac{[H^+]}{[H^+]_0}\right]^2} \right)
\]

The standard half-cell reduction potential at pH 7 is \( E_{GSH}^{\text{pH}(H^+)} = -240 \) mV, whereas at pH 6, it is \( E_{GSH}^{\text{pH}(H^+)} = -180 \) mV (i.e., more oxidizing under acidic conditions). On this basis, they correctly predicted that the iron in WT vacuoles should be FeIII. For each pH increase of 1 (a 10-fold drop in acidity), the effective reducing power of glutathione increases by 59 mV. Assuming pH 6 to 7 in *vma2Δ* vacuoles, the thermodynamic reduction potential for the FeIII/FeII couple in vacuoles should be in the range of the values listed above.

Interestingly, the same analysis predicts that the FeII should reoxidize to FeIII in *vma2Δ* cells grown at lower pH, which was not observed; the dominant iron in these cells remained FeIII for pH values as low as 3. Perhaps the pH of *vma2Δ* vacuoles is less sensitive to media pH than we have assumed. Other factors may also be involved (43). The redox state of vacuolar iron is likely influenced by the potential ligands in the organelle. For instance, vacuolar iron is coordinated by polyphosphate anions, and the concentration of these chains is diminished in *vma2Δ* vacuoles (16).

Diab and Kane predicted that *vma2Δ* cells should contain high levels of FeII ions (23). These researchers suggested that this form of iron is toxic to the cell because it participates in Fenton chemistry \( \text{Fe}^{II} + \text{H}_2\text{O}_2 \rightarrow \text{Fe}^{III} + \text{OH}^- + \text{OH}^- \). However, we treated *vma2Δ* cells with H2O2 but found no evidence using MB spectroscopy that the FeII was altered in peroxide-treated cells. Reducing equivalents in the cell may have quickly reduced FeIII back to FeII such that FeII remained dominant in the treated cells despite being used in Fenton chemistry. Upregulation of the cytosolic peroxidase Tsa2 in *vma2Δ* cells may have also diminished the effects of H2O2 (22).

Many of our results differed from expectations based on previous studies of *vma2Δ* cells. To examine whether there might be a strain dependence, we collected MB spectra of *vma2Δ* cells grown at different media pH values. Resulting spectra were closer to our expectations, with WT-looking vacuolar FeIII in *vma2Δ* cells grown at low pH and NHHS FeIII and nanoparticles for cells grown at high pH. BY4741 and W303 cells have different plasma membrane potentials and different sensitivity to alkali-metal salts (44). W303 cells are larger than BY4741 cells, their overall protein concentration is lower, and their vacuoles occupy a greater fractional cell volume (45). We found that *vma2Δ* cells were less sensitive to media pH and exhibited milder phenotypes associated with nonacidic vacuoles such that mitochondria were largely unaffected under the conditions used. We speculate that this reduced sensitivity might reflect larger vacuoles (a lower surface-to-volume ratio and therefore a smaller interface with the cytosol) in W303 cells, but further studies are required to verify this.

We have compared our results mainly to those of Kane and Hughes, but other studies report results regarding *vma* mutants that are more similar to ours. Ohya et al. (46) found that O2 consumption rates in *vma* mutant cells (from parent strains other than W303 or BY4741) grown on glucose were “identical” to those of WT cells. They also reported no significant change in F-ATPase and succinate dehydrogenase activities, consistent with our results and with our suggestion of strain-dependent differences in *vma*-mutant phenotypes.

Hughes et al. (26) suggested that cytosolic accumulation of cysteine drives the low iron and oxidative stress response in V-ATPase–deficient cells. We observed a LMM sulfur species in isolated cytosol from WT and *vma2Δ* cells and found that its intensity increased in *vma2Δ* cells relative to that of traces of WT FT5s. This shoulder to the MES buffer peak approximately comigrated with cysteine and other LMM sulfur species, including GSH and methionine. The increased shoulder intensity in *vma2Δ* isolated cytosol is consistent with the

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**Figure 12. Western blot of fermenting WT and *vma2Δ* whole-cell extract from both W303 and BY4741 background strains.** Top row, anti-Fet3. Bottom row, anti-Pgk1. A, image of prestained protein ladder with masses indicated. B–C, w303 WT and *vma2Δ*, respectively, cultured in MM supplemented with 40 μM iron. D–E, BY4741 WT and *vma2Δ*, respectively, cultured in YPAD. All bands in each row were run on the same gel. MM, minimal medium; YPAD, yeast extract peptone adenine dextrose medium.
findings of Hughes et al., though further confirmation of elevated cytosolic cysteine levels is required.

\( \textit{vma2Δ}_W \) cells grew slower than WT cells and were more easily intoxicated by high concentrations of iron in the medium. We simulated these differences by assuming that \( \textit{vma2Δ}_W \) cells are unable to sequester iron (or unable to sequester it in a benign form). However, the situation is undoubtedly more complicated. The toxicity of imported iron may also be associated with the oxidation state of that iron, with \( \text{Fe}^{III} \) being more toxic than \( \text{Fe}^{II} \).

\( \textit{vma2Δ}_W \) cells and cytosol contained less Mn than in WT cells and cytosol; a similar decline has been observed in \( \textit{PPN1-UP} \) cells (47) which have elevated levels of the polyphosphatase \( \text{Ppn1} \). In both strains, cells contain low levels of polyphosphate relative to WT. Mn binds vacuolar polyphosphate (12), suggesting that these mutant cells may have responded to a decreased capability to store Mn in vacuoles by slowing Mn import.

The decline of phosphorus in \( \textit{vma2Δ}_W \) cells and cytosol may be related to changes in the activity/expression levels of \( \text{Pho84} \) and \( \text{Pma1} \), both of which are located on the plasma membrane (14). \( \text{Pho84} \) is a symporter that imports protons and phosphate ions into the cytosol; \( \text{Pma1} \) exports protons from the cytosol. The activity of \( \text{Pma1} \) decreases in \( \textit{vma} \) mutants (7), which along with the loss of V-ATPase activity increases the acidity of the cytosol. We speculate that this might decrease the activity of \( \text{Pho84} \) and thus reduce the import of phosphate into \( \textit{vma} \) cells. \( \text{Pho84} \) may also be involved in manganese and zinc import (48), and a decline in activity could have contributed to the observed decline of Mn and Zn concentrations in \( \textit{vma2Δ}_W \) cells.

The increased concentration of Cu in our \( \textit{vma2Δ}_W \) cells and cytosol confirms earlier reports that \( \textit{vma} \)-mutant cells contain high levels of copper and that copper is dysregulated in these cells (17, 24). A similar accumulation may be driving the reported decline in \( \text{Ctr1} \) expression in \( \textit{vma2Δ}_B \) cells (22). Most LMM Cu in both WT and \( \textit{vma2Δ}_W \) cytosol is probably bound to metallothionein \( \text{Cup1} \). Higher copper concentrations in \( \textit{vma2Δ}_W \) cells likely afforded increased levels of Cu-bound \( \text{Cup1} \). The inability of nonacidic vacuoles to sequester copper may have increased the concentration of cytosolic Cu, which in turn induced expression of \( \text{CUP1} \). Another possibility is that Cu-bound \( \text{Cup1} \) is mobilized to vacuoles for degradation and that this process is inhibited in nonacidic vacuoles, thereby causing a buildup of Cu-bound \( \text{Cup1} \) in the cytosol. A similar peak was tentatively assigned to \( \text{Cup1} \) in chromatograms of vacuolar lysates (12).

Does iron trafficking in \( \textit{Saccharomyces cerevisiae} \) include a direct or dedicated vacuole-to-mitochondria pathway? A direct pathway would involve contact of the two organelles (kiss-and-run) and would be independent of cytosol. A dedicated pathway would involve the synchronized export of a particular iron species from the vacuole into the cytosol, and as a second step, the import of that species exclusively into mitochondria. If the iron exported from vacuoles entered a cytosolic pool of iron that had various trafficking destinations, including but not limited to mitochondria, we would not regard that pathway as being direct or dedicated.

We have not systematically distinguished these possibilities, but our results disfavor direct or dedicated pathways. The vacuoles are nearly empty in cells grown in iron-deficient or iron-limited media (e.g., 1 \( \mu \text{M} \text{ Fe}^{III} \)), yet MB spectra of such \( \textit{vma2Δ}_W \) cells afforded a CD as intense as for equivalent WT cells. This suggests that vacuoles are unnecessary for iron to enter mitochondria and be used to generate ISCs. We tentatively conclude, pending further investigations, that cytosolic iron can enter mitochondria directly without first entering and exiting vacuoles and that there is no direct/ dedicated vacuole-to-mitochondria trafficking pathway as defined above.

So how does the non acidity of vacuoles negatively impact ISC metabolism in mitochondria? We suggest, following Hughes (26), that the effect indirectly involves vacuoles and directly involves the cytosol. Cysteine or a derivative thereof may be blocked from entering nonacidic vacuoles, and its buildup in the cytosol may inhibit iron traffic into mitochondria. Likewise, a hyperacidic cytosol, caused by the absence of V-ATPase to pump cytosolic protons into vacuoles and/or a decline in \( \text{Pma1} \) activity, may also hinder iron trafficking. In support of this, we observed a shift in the LMM iron species in \( \textit{vma2Δ}_W \) cytosol, from dominance of a species with an apparent mass of 600 Da to one with an apparent mass of 800 Da.

Finally, we have had difficulty throughout this study reconciling the milder phenotype of the \( \textit{vma2Δ}_W \) strain with the more dramatic phenotype of \( \textit{vma2Δ}_B \). Both mutant cells grow slowly and with difficulty under respiring conditions, but \( \textit{vma2Δ}_W \) mitochondria are indistinguishable (by our methods) from WT mitochondria. In contrast, \( \textit{vma2Δ}_B \) mitochondria show difficulty generating ISCs, the iron regulon is clearly activated, and nanoparticles and ROS are likely generated. Those cells recover only under conditions in which the media is supplemented with excessive iron.

We suggest that \( \textit{vma2Δ}_W \) mitochondria suffer from the same originating problem as \( \textit{vma2Δ}_B \) but to a milder degree. We hypothesize that the fundamental problem is a slow rate of iron import from the cytosol to mitochondria, due to the hyper acidity of this cellular region and/or to the inhibitory effect of high cysteine concentrations. The effect of slow iron import into mitochondria on the growth of \( \textit{vma2Δ}_W \) cells would not be evident from our studies because we impatiently waited for them to grow to sufficient absorbance at 600 nm prior to harvesting them for MB spectroscopy. By doing this, the ultimate iron content is expected to be like that of WT cells (apart from the reduction of vacuolar \( \text{Fe}^{III} \) → \( \text{Fe}^{II} \)). For \( \textit{vma2Δ}_B \) cells, the rate of iron traffic from the cytosol to the mitochondria might be so slow that the respiratory shield weakens and a vicious cycle ensues, leading to nanoparticles, ROS, and an activated iron regulon, similar to that observed in \( \textit{MRS3/4ΔΔ} \) cells, which can only import iron slowly into mitochondria under iron-limited growth conditions (49, 50). In \( \textit{vma2Δ}_W \) cells, this effect might be muted.
Experimental procedures

Generation of the vma2Δ strain

Studies were conducted with the haploid strain of *S. cerevisiae*, W303 (MAT *α ade2-1, his3-3-11, 15 leu2-3,112, trp1-1, ura3-1). The VMA2 open reading frame was deleted in W303 by the Longtine method (51). Primers were synthesized including VMA2 gene specific sequences (lower case) and plasmid specific sequences (upper case): F1 (agagtagacagcatacaggaataaatattcgagg/CGGATTTCCCCGTATTTAA); R1 (gacaataaaaaagctttctctgaaacgtcctc/GAA TTCGAGCTCGTATTTAACC).

A PCR was performed on template plasmid pF6A-His3MX6 (51) carrying the *S. pombe his5+* gene, and a 1403-bp DNA fragment was amplified as expected. The DNA product (about 5 μg) was used to transform W303 yeast using the lithium acetate procedure (52), and several colonies were selected for histidine prototrophy and carried forward. The correctness of the VMA2 deletion was verified as follows. Genomic DNA was extracted from the transformants, and a PCR, using forward primer I (ctcatgacgcagtgatgcg) and reverse primer N (ATGTA-GATGTAAGAACCTATAC), was performed. The 1 primer resides in the 5' region of VMA2 gene outside the region of recombination. The N primer resides in the TEF terminator region of the pF6A-His3MX6 plasmid. The derived PCR product was approximately 688 bp in size and was present in the knockouts but not in the controls (Fig. S2), confirming that the VMA2 flanking region and His3MX6 module including the *S. pombe his5+* gene were correctly juxtaposed in the genome of the putative knockouts. The knockout strains had the expected auxotrophies and prototrophies and grew more slowly than the parent on rich or defined media.

Cell growth

W303 WT (MAT *α ade2-1 his3-11,15 leu2-3,112 trp1-1 ura3-1) and W303 vma2Δ (MAT *α ade2-1 his3-11,14 leu2-3,112 trp1-1 ura3-1 Δvma2::His3MX6) cells were obtained from frozen 15% glycerol stocks on YPAD (2% (w/v) glucose, 1% (w/v) yeast extract, 2% (w/v) peptone, 0.01% (w/v) adenine, 2% (w/v) bactoagar) plates. BY4741 WT (MATα his3Δ1 leu2Δ0 met15Δ0 ura3Δ0) and congenic BY4741 (Δvma2::KanMX6) cells were maintained similarly.

Individual colonies were inoculated in precultures of MM (20 g/l glucose, 5 g/l ammonium sulfate, 1.7 g/l yeast nitrogen base without ammonium sulfate and ferric chloride (MP Bio), 100 mg/l leucine, 50 mg/l adenine, 20 mg/l histidine, 20 mg/l uracil, 48 mg/l tryptophan, 10 μM copper sulfate) and either 1 or 40 μM ferric citrate. For two batches, the medium was supplemented with 1 mM cysteine. Media were buffered with a 0.05 M citric acid/sodium citrate buffer in varying ratios. For pH 3, 4, 5, 6, and 7, the citric acid-to-sodium citrate ratio was 13.3:1, 1:9:1, 1:1.4, 1:4.4, and 0:1, respectively. Minor adjustments were made with NaOH to achieve the desired media pH, as monitored using an Accumet AB15 pH meter.

BY4741 cells for Western Blot were cultured in YPAD media (20 g/l glucose, 10 g/l yeast extract, 20 g/l peptone, 100 mg/l adenine). Cell cultures were incubated at 30 °C and 180 rpm shaker speed. Absorbance was monitored at 600 nm using a Genesy6 spectrophotometer (Thermofisher). Fifty to 100 ml of precultures was transferred to 1- to 2-L media when absorbance at 600 nm was 0.7 ± 0.1. Cells were harvested from 1 to 2 L when culture was at exponential phase (absorbance at 600 nm = 0.8 ± 0.2). Cells were pelleted via centrifugation at 5000g for 5 min using a Sorvall LYNX6000 superspeed centrifuge (ThermoFisher). Cells were washed twice with 1 mM EDTA and then twice with deionized water (5 ml of wash per gram wet cell pellet). After each wash, cells were pelleted as above and the supernatant was discarded. Finally, pellets were suspended in a minimal volume of double-distilled water and pelleted into MB cups by centrifugation at 6400 RPM for 10 min using a Beckman Coulter Optima L-90K ultracentrifuge (SW32Ti rotor); samples were frozen in liquid N2 for later analysis. EPR samples were pelleted after washing by centrifugation at 5000g for 5 min into EPR tubes.

For H2O2 studies, whole-cell MB samples were thawed and incubated in 1 ml 5% H2O2 for 15 min or in 200 μl of 0.0034% H2O2 (initial concentrations) overnight. Cell pellets were reacquired for MB spectroscopy via centrifugation at 5000g for 5 min.

Respiring cell growth

Cell cultures were initially grown in 50 mL of fermenting MM as above. Once absorbance at 600 nm reached 0.6, the cells were pelleted and resuspended in 1.00 L of respiring MM (same as fermenting MM pH 5 except that 3% (v/v) glycerol and 1.5% (v/v) ethanol replaced glucose). Cells were harvested as described above.

Varying phosphate cell growth

Cell cultures were grown in fermenting CSM media at 30 °C and 180 RPM. CSM media consisted of 20 g/l glucose, 1.6 g/l CSM, 5.6 g/l yeast nitrogen base minus phosphates (MP Bio), 100 mg/l NaCl, 60 mg/l adenine, 100 mg/l leucine, 48 mg/l tryptophan, 10 μM copper sulfate, and 40 μM ferric citrate. The medium was supplemented with 0.5 and 21 mM monopotassium phosphate. All such studies were conducted at pH 5 using a 0.05 M citric acid/sodium citrate buffer. Cells were harvested as described above.

Mössbauer and EPR spectroscopics

Mössbauer spectra were collected at 5 K and 0.05 T on a MS4 WRC spectrometer (SEE Co, Minneapolis, MN) calibrated at RT using an α-iron foil. The applied field was parallel to the gamma radiation. EPR spectra were collected in an X-band Elexsys EPR spectrometer (Bruker) using the following conditions: 9.3789 GHz microwave frequency, 25 dB attenuation, 0.6325 mW power, 100 kHz modulation, 10G modulation amplitude, 5000G sweep width, 120 s sweep time, 5 to 10 scans.

Mitochondria studies

Mitochondria were isolated anaerobically from cells harvested from 24 L of fermenting cell cultures (pH 5), as described (35). Mitochondrial protein concentration was determined using the BCA assay. Mitochondria were pelleted via centrifugation at 12,000g for 10 min.
Mössbauer and LC-ICP-MS of vma2Δ yeast cells

Aconitase was assayed as described (53, 54) but with minor changes. Mitochondria were handled within an anaerobic glove box (MBraun Labmaster 130) containing ca. 5 ppm O₂. A pellet of mitochondria containing 80 mg of protein was diluted to 250 μl with lysis buffer (degassed 4.7 mM Triton X-100, 50 mM NaCl, and 50 mM Tris-HCl). Two hundred microliters of the resulting solution was mixed with 200 μl of 50 mM NaCl, 50 mM Tris-HCl, and 50 μl of double-distilled water in a 1-mm pathlength quartz cuvette. The sample was sealed with a rubber septum and removed from the box. Fifty microliters of 10 mM cis-aconitic acid (Sigma-Aldrich) was added through the septum, and the sample was mixed and immediately monitored at 240 nm for 360 s using a Hitachi U-3310 spectrophotometer. The molar absorptivity of cis-aconitic acid at 240 nm (ε$_{240}$) was 4.88 mM$^{-1}$ cm$^{-1}$ (54).

Cytosol and vacuole isolation

The cytosol was isolated from fermenting cells (pH 5) and filtered through a 10-kDa cutoff membrane as described (55). Filtered FTS was injected into a treated size-exclusion (SEC) Superdex Peptide 10/300 Gl column (GE Life Sciences) connected to an Agilent 1260 Bioinert quaternary pump (G5611A) and Agilent 7700x ICP-MS. The mobile phase was 20 mM ammonium acetate pH 6.5 flowing at 0.6 ml/min. Vacuoles were isolated as described (12).

Western blot

Western blot was completed using the whole-cell extract as described (12, 55). An anti-Fet3 antibody from rabbit was used in a 1:1000 dilution. The antibody was generated as described (28).

Elemental analysis

Isolated cytosol samples (100 μl) or double-distilled water-washed whole cells (ca. 100 mg) suspended in fresh double-distilled water were digested with 500 μl of undiluted (70%) trace metal-grade HNO$_3$ (Fisher) at 70 °C for ca. 16 h in plastic falcon tubes sealed using electrical tape. Samples were cooled to RT, and 250 μl of 35% hydrogen peroxide was added. Samples were resealed and incubated at 65 °C for 90 min. Samples were diluted to 10 ml with double-distilled water, affording ca 3.5% and 0.86% (w/w) final concentrations of trace metal–grade nitric acid and hydrogen peroxide, respectively. A cell pellet packing efficiency of 0.7 and whole-cell pellet density of 1.1029 g/ml (56) were used in calculations.

Data availability

All data except those described as “Data not shown” in the text are contained within the manuscript and SI.

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Author contributions—J. E. K. performed most of the experiments and analyzed all results; S. W. V. collected the Mössbauer and electron paramagnetic resonance spectra and maintained those instruments; T. Q. N. performed some experiments and training; A. D. performed all genetic manipulations; P. A. L. conceived of the project, offered advice, analyzed results, and wrote most of the article. All authors edited the article and approved of its publication.

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Abbreviations—The abbreviations used are: CD, central quadrupole doublet; FTS, flow-through solution; ISC, iron sulfur cluster; LC-ICP-MS, liquid chromatography interfaced with inductively-coupled-plasma mass spectrometry; LMM, low-molecular-mass; MB, Mössbauer; NHHS, nonheme high spin; ROS, reactive oxygen species; WT, wild type; YPAD, yeast extract peptone adenine dextrose medium.

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