The Role of Defects in Fe(II)–Goethite Electron Transfer

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ABSTRACT: Despite substantial experimental evidence for Fe(II)–Fe(III) oxide electron transfer, computational chemistry calculations suggest that oxidation of sorbed Fe(II) by goethite is kinetically inhibited on structurally perfect surfaces. We used a combination of ^57Fe Mössbauer spectroscopy, synchrotron X-ray absorption and magnetic circular dichroism (XAS/XMCD) spectroscopies to investigate whether Fe(II)–goethite electron transfer is influenced by defects. Specifically, Fe L-edge and O K-edge XAS indicates that the outermost few Angstroms of goethite synthesized by low temperature Fe(III) hydrolysis is iron deficient relative to oxygen, suggesting the presence of defects from Fe vacancies. This nonstoichiometric goethite undergoes facile Fe(II)–Fe(III) oxide electron transfer, depositing additional goethite consistent with experimental precedent. Hydrothermal treatment of this goethite, however, appears to remove defects, decrease the amount of Fe(II) oxidation, and change the composition of the oxidation product. When hydrothermally treated goethite was ground, surface defect characteristics as well as the extent of electron transfer were largely restored. Our findings suggest that surface defects play a commanding role in Fe(II)–goethite redox interaction, as predicted by computational chemistry. Moreover, it suggests that, in the environment, the extent of this interaction will vary depending on diagenetic history, local redox conditions, as well as being subject to regeneration via seasonal fluctuations.

INTRODUCTION

Ferrous and ferric iron comprise one of the most abundant redox couples, and electron transfer between these two oxidation states controls the cycling and availability of Fe in water, soil, and air.¹,² Over the past decade significant evidence has accumulated to demonstrate interfacial electron transfer between sorbed Fe(II) and Fe(III) in Fe oxides and Fe-containing clay minerals.³–¹⁵ In some cases, electron transfer also appears to be followed by mixing of Fe atoms from the bulk mineral structure with the surrounding fluid (also termed Fe(II)-catalyzed recrystallization).¹⁰,¹⁶–²² While Fe(II)–Fe(III) electron transfer and mixing have been clearly demonstrated, a mechanistic understanding of these reactions remains elusive. Knowledge gaps in our understanding of Fe(II)–Fe(III) reaction mechanisms limit our ability to reliably predict important environmental and geochemical processes, such as cycling of C, N, and P,²⁴–²⁹ water treatment,³⁰ contaminant remediation,³¹–³⁴ metal cycling,³⁵,³⁶ mineral transformations,³⁶ and interpreting the ancient rock record.³⁷

Thus far, Fe(II)–Fe(III) electron transfer has been demonstrated for several Fe oxides, including hematite, goethite, magnetite, ferricydrite, as well as Fe-containing clay minerals.³–¹⁵ Of the Fe minerals, electron transfer between Fe(II) and goethite has been the most extensively studied.

Oxidation of Fe(II) by goethite has been shown to occur over a range of Fe(II) concentrations, amounts of Al-substitution, and in the presence of various anions, such as phosphate, bicarbonate, silicate, and organic matter.³,⁴,⁶,⁸,¹⁰,¹²,¹⁶,¹⁷ Despite abundant experimental evidence for Fe(II)–Fe(III) oxide electron transfer, computational evidence suggests that Fe(II)–goethite electron transfer is not energetically favored on structurally perfect surfaces.³⁸–⁴⁰ For example, density functional theory (DFT) calculations of Fe(II) adsorbed onto charge neutral, stoichiometric goethite (110) surfaces suggest that the oxidation of sorbed Fe(II) by lattice Fe(III) is energetically uphill. Others similarly conclude that charge only minimally delocalizes between Fe(II) and surface Fe(III), with only a minor dependence on the type of Fe(II)–mineral complex formed.³⁸,⁴⁰ Additionally, a recent molecular dynamics study examining the electron transfer kinetics for stable inner and outer-sphere Fe(II) complexes on a wide range of perfect goethite terminations showed that the energetics are at best only thermoneutral, with large activation energies.⁴¹

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Each of these studies, however, suggest or infer that surface defects may underlie the experimentally observed interfacial electron transfer.\textsuperscript{38–40} For example, calculations simulating an oxygen vacancy on goethite (110) showed that electron transfer to the resulting lower-coordinated Fe(III) was more energetically favorable and less kinetically inhibited. This idea is conceptually similar to the notion that defects provide traps in the interior of the solid for excess electrons to accumulate.\textsuperscript{9,42} Given these computational findings, experiments designed to test the prospective role of defects could help advance our mechanistic understanding of Fe(II)–Fe(III) oxide electron transfer.

Defects arise from a deviation in the perfect composition and/or structure of a mineral. Such features are common in goethite,\textsuperscript{43–47} and have been previously studied in terms of their relationship to growth conditions,\textsuperscript{45,46,48} including non-classical growth pathways such as imperfect oriented attachment.\textsuperscript{49–51} The type and concentration of defects strongly influence goethite properties (e.g., α, dimension of the unit cell,\textsuperscript{43} Neel temperature,\textsuperscript{52} magnetic moment\textsuperscript{45,53–55}) and particle reactivity (e.g., dissolution rates and ion sorption). Defects in the ideal goethite stoichiometry (i.e., α-FeOOH) can take the form of iron or oxygen vacancy, but often can be characterized as an excess of water/hydroxyl relative to the theoretical formula.\textsuperscript{46} Such that the effective formula is α-Fe
\[\text{Fe}_1-\text{y/3} \text{O}_{1-\text{x}} (\text{OH})_{1+x}\] Changes in goethite properties after hydrothermal treatment have been interpreted to be due to removal of defects.\textsuperscript{43,44,56} Conversely, ball-milling has been suggested to add defects to goethite based on observed changes in the Neel temperature.\textsuperscript{45}

While there is clear evidence that defects are often present in goethite, little is known about how defects influence Fe(II)–goethite electron transfer. To date, no experimental data has been presented that specifically evaluates the role of defects in Fe(II)–goethite electron transfer, despite significant speculation about their role in enabling electron transfer and driving Fe(II)-catalyzed Fe oxide recrystallization.\textsuperscript{19,21,38–40} Here we provide the first experimental evidence that shows defects influence the extent of Fe(II)–goethite electron transfer and the composition of the product formed. Our findings indicate that low temperature Fe(III) hydrolysis results in goethite particles that have excess hydroxyl/water content and corresponding Fe vacancies that enable Fe(II)–goethite electron transfer. Hydrothermally treating the goethite particles appears to remove defects, inhibit Fe(II)–goethite electron transfer, and alter the composition of the oxidation product. Our findings suggest that surface defects play an important role in Fe(II)–goethite redox interaction, as predicted by computational chemistry.

## MATERIALS AND METHODS

### Oxide Synthesis

Goethite was prepared from 56Fe-enriched Fe metal ((Isoflex, 99.94% purity), 56Fe goethite) by modifying the Schwertmann and Cornell method, using iron metal as the synthesis’ starting point instead of Fe(NO\textsubscript{3})\textsubscript{3}.\textsuperscript{57} Briefly, 56Fe(0) was dissolved in HCl to obtain 15 mL of an Fe(II) stock (\(\sim 0.6 \text{ M Fe(II)}, \sim 1.8 \text{ M HCl}\)), and the solution was oxidized using 2 mL of 30% (slight excess) H\textsubscript{2}O\textsubscript{2} to produce Fe(III). Then, the pH was raised with 16 mL of 5 M KOH and the resulting precipitate was placed in an oven at 70 \textdegree C for 60 h. 56Fe goethite was washed, centrifuged, freeze-dried, ground with a mortar and pestle, and passed though a 100 mesh sieve. The final mineral is referred to as “as-synthesized” and it is similar to the microgoethite used in our previous work.\textsuperscript{5,6,10,12,16,17} The Brunauer–Emmett–Teller (BET) specific surface area was determined by N\textsubscript{2} sorption at 77 K and found to be 28 to 34 m\textsuperscript{2} g\textsuperscript{-1}. X-ray diffraction (XRD - Rigaku Mini FlexII) patterns showed that the material contains goethite and no other minerals (Supporting Information (SI) Figure S7).

### Electron Transfer Experiments

All experiments were carried out in an anaerobic glovebox with N\textsubscript{2}/H\textsubscript{2} atmosphere (93/7%), and all solutions were purged at least 2 h with N\textsubscript{2} prior to transfer into the glovebox. Fe(II) stock solutions were prepared inside the glovebox by reacting 57Fe metal (Cambridge Isotope, 96.93% purity) with 1 M HCl overnight. The resulting solution was filtered to remove any residual Fe(0) and diluted with deionized (DI) water to the desired concentration (\(\sim 100 \text{ mM Fe(II), } \sim 0.1 \text{ M HCl}\)).

Batch reactors were prepared by adding 10 mL of 25 mM HEPES (4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid), pK\textsubscript{a} 7.55\textsuperscript{58} buffer adjusted to pH 7.5 ± 0.05 plus 25 mM KBr electrolyte to a 20 mL glass vial and adding Fe(II) stock to reach an initial 57Fe(II) concentration of \(\sim 1 \text{ mM}\). The reaction was started by adding 20.0 ± 0.2 mg of 56Fe goethite and the reactors were placed on a end-over-end rotator in the absence of light. The aqueous phase was filtered (0.2 µm) and acidified with trace metal grade HCl for subsequent Fe(II) and total Fe analysis using the 1,10-phenanthroline method.\textsuperscript{59}

### Extraction

Two additional reactors were prepared by reacting 56Fe goethite with 57Fe(II), and the reacted solids were centrifuged and then extracted to remove the sorbed Fe species. The first reactor was extracted with a 0.4 M HCl (15 min). For the second reactor, the reacted solids were subjected to a sequential extraction procedure using HEPES buffer (pH 7.5, 1 h) followed by 1 M CaCl\textsubscript{2} (pH 7, 4 h) and 1 M NaH\textsubscript{2}PO\textsubscript{4} (pH 5, 18 h). A 30 min wash step with DI water was carried out after the CaCl\textsubscript{2} and NaH\textsubscript{2}PO\textsubscript{4}. The extracted solids were analyzed by Mössbauer spectroscopy and the extracted aqueous phase were analyzed for Fe(II) and total Fe.

### Hydrothermal Treatment

Goethite was subjected to a hydrothermal treatment to anneal defects. A suspension of the as-synthesized 56Fe goethite in deionized water was placed into a digestion bomb, and kept in an oven at 150 \textdegree C for 44 h. The digestion bomb was allowed to cool down and the solids were centrifuged and freeze-dried (referred to as hydrothermally treated goethite). The batch of hydrothermally treated goethite was split in two, and part of it was reacted with 1 mM 57Fe(II) as discussed above. The other part was crushed with mortar and pestle to restore defects (referred to as ground goethite). Again, the batch was split, and part was reacted with 1 mM 57Fe(II), while the other aliquot was hydrothermally treated again under the same conditions, and then reacted with 1 mM 57Fe(II) (referred to as hydrothermally treated again goethite).

### Mössbauer Spectroscopy

For Mössbauer spectroscopy, solids were collected on a 0.2 µm nitrocellulose filter and then sealed between two pieces of Kapton tape to avoid air oxidation. Mössbauer spectra were collected at 77 K on a spectrometer supplied by Web Research, Inc. (Edina, MN) and equipped with closed-cycle cryostat (CCS-850 System, Janis Research Co., Wilmington, MA). We acquired spectra in...
transmission mode using a constant acceleration drive system and a $^{57}$Co source. The velocity scale was calibrated using a 7 μm α-Fe(0) foil. We fit the spectra using the software Recoil (Ottawa, Canada).$^{60}$

Selected samples were reanalyzed after atmosphere exposure. After initial Mössbauer analysis, these samples were stored for a month in normal atmosphere and then reanalyzed by Mössbauer spectroscopy. For comparison, one sample was kept inside the anaerobic glovebox for the same time span.

**Synchrotron X-ray Absorption and Magnetic Circular Dichroism Spectroscopy.** For X-ray absorption spectroscopy, suspensions of goethite were dropped onto indium foil in an anaerobic glovebox with $N_2/H_2$ atmosphere (93/7%), dried, then pressed into the foil and the excess solid was removed. The indium foil was attached to the copper sample manipulator with silver paint. XAS and XMCD measurements were performed using Beamline 6.3.1.1 at the Advanced Light Source (Berkeley, CA). Fe L$_{2,3}$-edge and O K-edge XAS were recorded at room temperature in total electron yield mode; Fe L$_{2,3}$-edge XAS has an effective probing depth of 50 Å. Fe L$_{2,3}$-edge XAS spectra were obtained by measuring two XAS spectra with a fixed degree of circular polarization of $\sim$0.7 and with opposing magnetization directions by reversing the applied field of 1.8 T at each energy point. The XAS spectra were normalized to incident beam intensity, and the XMCD spectra was obtained as the difference between the two spectra.

**Density Functional Theory Computations.** We used the FDMNES code$^{61}$ to individually calculate the O K-edge XANES spectra of the two oxygen sites present in goethite, explicitly taking into account their respective local coordination environments in the bulk structure. The Green’s formalism approach, within the limit of the muffin-tin approximation, was used. The Fermi energy has been determined self-consistently using an aggregate of radii of 7 Å. A cluster of 7 Å radii has also been used to perform the final state calculations. The Hedin-Lundquist potentials$^{62}$ were used to model the exchange-correlation. Dipoles, core-hole contributions and spin–orbit coupling were taken into account. Because the experimental spectra of goethite is made of two components with unknown relative position and intensity, we have used a non-negative least-squares algorithm to refine the position and intensity of the two calculated O$^{2-}$ and OH$^-$ component spectra, and have found the best linear combination that would fit the experimental spectra. The resulting linear combination fit of the measured O K-edge spectra with the computed component spectra for O$^{2-}$ and OH$^-$ was used to quantify relative contributions of these two oxygen species.

**RESULTS AND DISCUSSION**

**Fe(II)—Goethite Electron Transfer.** To explore if defects influence Fe(II)—goethite electron transfer, we ran a series of Fe isotope labeled experiments with as-synthesized and hydrothermally treated goethite particles. Similar to our previous work,$^{3,5,6,10–12,14}$ we took advantage of the element and isotope specificity of $^{57}$Fe Mössbauer spectroscopy to track if Fe(II)—goethite electron transfer occurs. Here, we treated Mössbauer-invisible $^{56}$Fe goethite with a sequence of hydrothermal treatments. Experimental conditions: $[^{56}\text{Fe}]=2$ g L$^{-1}$, 25 mM HEPES/25 mM KBr at pH 7.5.

After reaction with $^{57}$Fe(II), there is, however, a marked difference between the as-synthesized goethite and the hydrothermally treated goethite. The Fe(III) sextets capture a smaller portion of the spectral area of the hydrothermally treated compared to the as-synthesized goethite (as shown by blue sextets in Figure 1a and b). To test if the change in spectral area distribution upon hydrothermal treatment was reversible, we ground the hydrothermally treated goethite particles and reacted them with $^{57}$Fe(II). The resulting spectrum shows a marked increase in the amount of area captured by the sextet (and less hatched area) (Figure 1c). A second hydrothermal treatment returned the spectrum of Fe(II)-reacted solids to one similar to that of the first hydrothermal treatment (Figure 1d). To test that the change in spectral area trend observed in Figure 1 was not particular to a goethite synthesis batch, we ran duplicates of each treatment using two separate batches of goethite (SI Figure S1). The solids revealed two prominent Fe(III) sextets consistent with goethite and suggest that substantial oxidation of $^{57}$Fe(II) occurred and formed $^{57}$goethite on both the as-synthesized goethite and the hydrothermally treated goethite (SI Table S1).

Oxidation of sorbed Fe(II) by goethite is consistent with our previous work.$^{3,6,8,10,12}$

Figure 1. Mössbauer spectra of $^{57}$Fe(II) reacted with $^{56}$Fe goethite after sequential hydrothermal/grinding treatments. Experimental conditions: $[^{56}\text{Fe}]=2$ g L$^{-1}$, 25 mM HEPES/25 mM KBr at pH 7.5.
similarity of the duplicate experiments conducted with two separately synthesized goethite batches provides strong evidence that hydrothermal treatment and grinding are reproducibly altering the goethite in a way that influences how it reacts with Fe(II).

The reversible, reproducible changes in the Mössbauer spectra indicate that hydrothermal treatment and grinding are influencing the product formed from $^{57}$Fe(II) interaction with goethite. To fit the hatched area in the Mössbauer spectra we tried a variety of approaches. We concluded that the best method to capture the hatched area was to include an Fe(II) doublet and a broad, collapsed sextet consistent with our and other’s previous approach (for more details see SI Figure S2).10,63 Small Fe(II) doublets comprising less than 10% of the total area have been previously observed in spectra of goethite reacted with $^{57}$Fe(II),8,10,12 but little is known about the composition of the broad, collapsed sextet and we have, in our previous work, been careful to not interpret it beyond that it was likely Fe(III).10 To check whether that the collapsed feature was influenced by buffer—Fe interactions,64 we ran a control experiment without buffer (SI Figure S4). There was no difference between the spectra of the buffered and unbuffered samples indicating that the buffer—Fe interactions were not responsible for the collapsed feature.

To investigate the composition of the collapsed feature, we subjected the as-synthesized goethite particles reacted with $^{57}$Fe(II) to different extraction procedures and collected Mössbauer spectra afterward (Figure 2). A mild HCl extraction (0.4 M) removed both the Fe(II) doublet and the broad, collapsed sextet from the Mössbauer spectra. All of the Fe(II) that had initially sorbed was recovered in the HCl extract (SI Table S2). Surprisingly, the aqueous extract from the HCl treatment contained only Fe(II), suggesting that the collapsed feature was at least partially due to Fe(II). To minimize the change in pH during extraction, we also extracted goethite reacted with $^{57}$Fe(II) with CaCl$_2$ and NaH$_2$PO$_4$ which only decreased the pH to 7 and 5, respectively. Similar to the HCl extraction, the milder extractions removed most of the collapsed feature and recovered similar amounts of Fe(II) (for additional discussion see SI Figure S5).

As an alternative approach to evaluate whether the collapsed feature contains Fe(II), we oxidized a sample of goethite reacted with $^{57}$Fe(II) to see if the collapsed feature disappeared. One month of air exposure resulted in complete removal of the collapsed feature and the Fe(II) doublet from the spectra, providing an additional line of evidence that the collapsed feature contained some Fe(II) (Figure 2). The spectra of the oxidized goethite, however, looks slightly different than the spectra of the HCl-extracted goethite, in which a third sextet appeared. The third sextet was identified as akaganéite, a mineral typically formed by the hydrolysis of Fe(III) salts in the presence of Cl$^-$.65 Regardless of the precise identity of the collapsed feature, the oxidation and extraction data combined provide compelling evidence that the collapsed feature contains some Fe(II) and is more likely a mixed Fe(II)−Fe(III) phase rather than a pure Fe(III) phase as we previously thought.10 While we cannot rule out that it is a pure Fe(II) phase, the Mössbauer parameters of the collapsed sextet are more consistent with a mixed Fe(II)−Fe(III) compound (further discussed in the SI).66 Importantly, the presence of a mixed Fe(II)−Fe(III) phase rather than an Fe(III) phase suggests that some of $^{57}$Fe(II) sorbed on goethite was not oxidized by the goethite.
goethite sextets, it would indicate that all of the sorbed $^{57}$Fe(II) was oxidized to $^{57}$goethite. Complete oxidation of Fe(II) did not occur in any of the samples, but instead the amount of Fe(II) oxidized to form goethite varied between 63 and 87% (as estimated from the relative area of the sextets). What is most striking in Figure 3, however, is how reversible and reproducible the hydrothermal and grinding treatment is in influencing the extent of oxidation of the goethite-sorbed Fe(II). The as-synthesized goethite oxidized $\approx$87% of the sorbed Fe(II) to form goethite, whereas hydrothermally treating the goethite particles resulted in only $\approx$66% of the sorbed $^{57}$Fe(II) becoming oxidized to goethite, suggesting that electron transfer was inhibited by hydrothermal treatment and that the composition of the oxidation product changed. Grinding the hydrothermally treated goethite largely restored the extent of electron transfer ($\approx$ 81%), and hydrothermally treating this goethite sample again inhibited the extent of electron transfer ($\approx$63%) (Figure 3).

Similar to the as-synthesized goethite, one month of air exposure of the hydrothermally treated goethites resulted in removal of the collapsed feature and the Fe(II) doublets from the spectra, consistent with the collapsed feature containing some Fe(II) (SI Figure S6). The hydrothermally treated goethites reacted with $^{57}$Fe(II) also lost substantially more spectral area than the ground samples (as-synthesized and ground goethite), providing additional evidence that hydrothermally treating goethite leads to less oxidation of sorbed Fe(II) to goethite.

**Bulk Characterization of Goethite Particles.** It appears that hydrothermal treatment and grinding reversibly and reproducibly influences the extent of electron transfer from sorbed Fe(II) to goethite as well as the composition of the oxidation product. Given previous evidence that goethite synthesis conditions such as temperature alter the defect content in goethite, we hypothesized that hydrothermally treating the goethite particles at 150 °C annealed defects present in the goethite synthesized at 70 °C. We further hypothesized that grinding the particles added defects back to the hydrothermally treated particles. To evaluate if there were any changes in the bulk goethite particles before and after treatments, we characterized the particles with XRD, BET analysis, and microscopy. XRD spectra of treated solids confirmed that hydrothermal treatment did not transform goethite into any other mineral (SI Figure S7), and cell dimensions were unchanged after the goethite sequential hydrothermal/grinding treatments (SI Table S3). Schwertmann and collaborators observed a slight change in the a-axis dimension upon hydrothermal treatment, however, their original material was a highly defective goethite. Measurements revealed a small, but progressive loss in BET area (from 28 to 19 m² g⁻¹, SI Table S3), consistent with previous observations. In our work, TEM images revealed no substantial difference between as-synthesized and hydrothermally treated goethite, but SEM images revealed slight changes of the surface (SI Figure S8 and S9). Hydrothermally treated goethite appeared to have more perfectly formed ends when compared to the original mineral suggesting the hydrothermal treatment may have removed some surface defects (SI Figure S9).

Because hydrothermal treatment resulted in smaller BET surface area and amounts of Fe(II) sorbed, we considered the alternative hypothesis that less Fe(II)—goethite electron transfer could be simply due to less surface area and less surface-bound Fe(II) available to be oxidized. To explore whether the amount of sorbed Fe(II) influenced the extent of electron transfer, we plotted the percent relative area of the Fe(III) sextets versus sorbed Fe(II) per m² (Figure 4). We binned the data into ground goethites which includes as-synthesized and ground (solid markers) and hydrothermally treated goethites which includes both rounds of hydrothermal treatment (open markers). Both ground and hydrothermally treated goethites converge to $\sim$5.5 μmoles Fe(II) sorbed per m², revealing that the sorption of Fe(II) was not influenced by the treatment applied to the mineral. However, it is remarkable that, when we compare a hydrothermally treated and a ground goethite that have the same amount of Fe(II) sorbed per m², there is a marked difference ($\approx$20%) in the extent of electron transfer. Figure 4 provides compelling evidence that the changes we observed in Fe(II)—goethite electron transfer between our four treatments were not due to changes in surface area or amount of Fe(II) sorbed.

**Surface Characterization of Defects on Goethite Particles.** While bulk characterization of the treated goethite particles is informative, it is unlikely we would be able to detect specific changes in goethite surface structure with these techniques. To detect the presence of defects and changes in surface structure in the treated goethites, we collected X-ray absorption spectroscopy (XAS) and X-ray magnetic circular dichroism (XMCD) at the Fe L-edge and XAS at the O K-edge. By collecting total electron yield data, these techniques probe a depth no greater than 5 nm and are mainly sensitive to the upper few Angstroms. Fe L-edge XMCD spectra for all three samples is shown in Figure 5, while the illustrative O K-edge spectra and DFT-based spectral analysis are shown in Figure 6.

The Fe L-edge XAS probes the structure and valence of surface iron, whereas the XMCD is selective for the subset of surface iron that is magnetically ordered and is able to discriminate Fe valence and local coordination. As expected, the Fe L-edge XAS of all goethite samples measured shows features consistent with goethite Fe(III); Fe(II) is not detected (SI Figure S10). The corresponding XMCD information, however, is more revealing (Figure 5). The as-synthesized goethite has a weak magnetic moment at the surface (i.e., from the magnetic dichroism signal intensity), and the shape of the XMCD signal is consistent with octahedrally coordinated Fe(III). The
presence of this magnetic moment indicates that the surface is magnetic, likely due to Fe vacancies that disrupt the antiferromagnetic symmetry that would otherwise exist between perfect atomic planes of goethite. The XMCD signal of the hydrothermally treated goethite contains no distinguishable feature, indicating that the magnetic moment is absent, which is consistent with the known bulk antiferromagnetic structure and the hypothesis that hydrothermal treatment yields a more stoichiometric and crystalline material. After regrinding, the surface magnetic moment, and thus XMCD trace, is largely restored, consistent with reintroduction of Fe vacancies at the surface.

The corresponding O K-edge XAS spectra are illustrated in Figure 6, and analyzed in more detail in SI Figures S11 and S12. The spectral region of primary interest is the pre-edge region between 530 and 535 eV, which is comprised of two $1s \rightarrow 3d$ excitation doublets, one at lower energy corresponding to $O^{2-}$ in the goethite surface, and one at higher energy corresponding to $OH^−$ in the surface. Using DFT, we computed the specific expected shapes of these two doublets and used these theoretical components to perform linear combination fitting (LCF) of the experimentally measured O K-edge spectra for the three goethite samples. We then determined the $OH^−/O^{2−}$ ratio for the surfaces of each of three goethites. Additionally, the ratio of the measured total integrated Fe L-edge and O K-edge XAS intensities were used to estimate the Fe/O ratios in each of the three surfaces. Here we used the hydrothermally treated goethite as a normalization standard to the bulk ratio of 0.5 under the assumption that this surface is the most stoichiometric within the set. The complete surface compositional results are given in Table 1.

The O K-edge spectra for as-synthesized, hydrothermally treated and ground goethites show that the surfaces of the as-synthesized and ground goethite are more hydrous ($OH^−$ rich relative to $O^{2−}$) than the hydrothermally treated goethite (Figure 6 and Table 1). Note that while the bulk ideal $OH^−/O^{2−}$ ratio is 50%, the excitation cross sections of the two component spectra are not necessarily equivalent such that a ratio of their integrated intensities could be expected to also correspondingly be 50%; we chose not to normalize these ratios to the ideal value because the trends between samples remain the same in either case. Surfaces of as-synthesized and ground goethite bear comparably lower Fe/O ratios relative to the hydrothermally treated goethite (Table 1), again consistent with the prevalence of Fe vacancies in these two surfaces relative to the hydrothermally treated surface.

Collectively, the observed surface compositional characteristics are consistent with the known behavior of goethite stoichiometry to vary as $\alpha$-$Fe_{1-y}O_{1-\alpha} (OH)_{1+y}$. Due to excess protons, goethite grown at subhydrothermal temperatures, e.g. 70 °C as used here, tends to be both Fe deficient and correspondingly OH-rich. In contrast, goethite grown hydrothermally tends to be more crystalline and stoichiometric.

### Table 1. Summary of Surface Compositional Analyses of Goethite from X-ray Absorption Spectroscopy at the Fe L-Edge and O K-Edge, With the Latter Further Quantified in Terms of $O^{2−}$ and $OH^−$, Using Linear Combination Fitting of the Experimental Spectra with DFT-Based O K-Edge Doublets

<table>
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<th>as-synthesized</th>
<th>hydrothermally treated</th>
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<td>Fe/O</td>
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<td>$OH^−$</td>
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*To normalize Fe/O ratios based on integrated intensities of separately collected Fe L-edge and O K-edge spectra, all values were scaled proportionally by the factor needed to achieve the idealized 0.50 value for the hydrothermally treated goethite.
Combined, the results from Mössbauer spectroscopy, XAS, and XMCD suggest that particles with fewer defects are less prone to oxidize Fe(II). We propose that the key surface defects are Fe vacancies, which provide sites into which Fe(II) can strongly bind and transfer electrons to lattice Fe(III), propagating a goethite-like surface. Our findings suggest that surface defects play a role in Fe(II)–goethite redox reaction, as predicted by computational chemical modeling.38–40

ENVIRONMENTAL IMPLICATIONS

Here, we provide the first experimental evidence that defects influence the extent of Fe(II)–goethite electron transfer and the composition of the product formed. Our findings indicate that low temperature Fe(III) hydrolysis, a commonly used method for synthesizing goethite, results in goethite particles that have excess hydroxyl/water content and corresponding Fe vacancies that enable Fe(II)–goethite electron transfer. Hydromagnetically treating the goethite particles appears to remove defects, inhibit Fe(II)–goethite electron transfer, and alter the composition of the oxidation product. The clear role of defects in enabling Fe(II)–goethite electron transfer resolves the previous discrepancy between multiple experimental observations of Fe(II)–goethite electron transfer3,4,6,8,10,12,16,17 and computational calculations that suggest Fe(II)–goethite electron transfer is not energetically feasible on structurally perfect surfaces.38–40

Our experimental evidence that defects enable Fe(II)–goethite electron transfer raises the question of whether defects influence Fe(II)–catalyzed recrystallization as we and others have proposed that Fe(II)–Fe(III) interfacial electron transfer is an integral step in Fe(II)–catalyzed recrystallization.7,9,16,70,71 While there has been speculation that mineral surface and structural defects control Fe(II)–catalyzed recrystallization19,21 experimental data to evaluate this claim is lacking. If Fe(II)–Fe(III) oxide electron transfer controls the extent of Fe oxide recrystallization than our results support the hypothesis that defects play a role in Fe(II)–catalyzed recrystallization. We further speculate that the removal of defects to form a more perfect surface may be the energetic driving force for Fe(II)–catalyzed recrystallization that has continued to elude us. We caution, however, that alternative mechanisms of recrystallization such as solid-state diffusion or pore/void/intergranular diffusion16,19 have not been ruled out and neither of these mechanisms are likely to be as strongly influenced by electron transfer.

The role of defects in Fe redox chemistry also provides valuable insights into the behavior of Fe oxides in reducing environments. If defects in the form of Fe deficient and OH-rich surfaces enable Fe(II)–goethite electron transfer, then it seems reasonable to suggest that oxidative sorption of Fe(II) at the surface would fill Fe vacancies and anneal some surface defects. Consistent with our suggestion that oxidative sorption of Fe(II) anneals surface defects, we have previously observed less oxidation of sorbed Fe(II) by hematite at high concentrations of Fe(II) as well as reduced extents of hematite recrystallization.72 Annealing of goethite by oxidative sorption of Fe(II) followed by inhibition of electron transfer may also explain the recent report of decreased goethite recrystallization rates over time.20 Our hypothesis that oxidative sorption of Fe(II) anneals surface defects is in agreement with results that show addition of Fe(II) inhibits rates of microbial Fe(III) reduction.5,7,19 How defects will impact heterogeneous redox process such as contaminant reduction rates3,35,58,75–78 and the observed paradoxical oxidation of As(III) at the Fe(II)–Fe oxide interface, however, remains open to experimental investigation.8,79,80

Our work also shows that electron transfer between Fe(II) and goethite is sensitive to diagenetic temperature and can be altered by relatively small changes in the structure. We note that we were only able to observe these changes with surface-sensitive techniques (i.e., XMCD and oxygen XAS). Environmental cycles that include temperature fluctuations, changes in the activity of water, and redox changes can likely reinitialize electron transfer between Fe(II) and goethite, and possibly restart recrystallization by controlling the defect content at the goethite/water interface. The small, subtle changes needed to influence Fe(II)–goethite interaction suggest that in the environment, the extent of this interaction, which also likely underlies Fe(II)–catalyzed recrystallization trace element release and incorporation,18,35,70 will be coupled to diagenetic history, local redox conditions, and be subject to regeneration via seasonal fluctuations.34,37

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.est.7b05772.

Details on Mössbauer fitting, discussion on HCl-extracted and oxidized samples as well as additional figures of fitted Mössbauer spectra, Mössbauer spectra of duplicates and extracted samples, SEM and TEM pictures and XRD spectra of the as-synthesized and treated sample (PDF)

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Notes
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