

The spin state of iron in minerals of Earth's lower mantle

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[1] The spin state of Fe(II) and Fe(III) at temperatures and pressures typical for the Earth's lower mantle is discussed. We predict an extended high-spin to low-spin crossover region along the geotherm for Fe-dilute systems depending on crystal-field splitting, pairing energy, and cooperative interactions. In particular, spin transitions in ferromagnesian silicate perovskite and ferroperricite, the dominant lower mantle components, should occur in a wide temperature-pressure range. We also derive a gradual volume change associated with such transitions in the lower mantle. The gradual density changes and the wide spin crossover regions seem incompatible with lower mantle stratification resulting from a spin transition. **Citation:** Sturhahn, W., J. M. Jackson, and J.-F. Lin (2005), The spin state of iron in minerals of Earth's lower mantle, *Geophys. Res. Lett.*, 32, L12307, doi:10.1029/2005GL022802.

1. Introduction

[2] The chemical composition, the elastic and transport properties, and the thermodynamic parameters of Earth's deep interior are of general importance to geochemical modeling, geodynamic simulation, and interpretation of seismic wave observations [Kellogg *et al.*, 1999; van der Hilst and Kárason, 1999; Trampert *et al.*, 2004]. Most of the minerals and polymorphs expected in Earth's mantle are believed to incorporate low concentrations of Fe(II) and/or Fe(III) of about 10 atomic% or less. The relevance of the Fe spin states in the major lower mantle constituents, ferroperricite $\text{Mg}_{1-x}\text{Fe}_x\text{O}$ with $0.1 \leq x \leq 0.25$ (hereafter referred to as Fp) and ferromagnesian silicate perovskite $\text{Mg}_{1-y}\text{Fe}_y\text{SiO}_3$ with $0.05 \leq y \leq 0.15$ (hereafter referred to as Pv) [Liu, 1975; McCammon *et al.*, 1998], with respect to density, iron partitioning, partial melting, radiative thermal conductivity, and compositional layering in the lower mantle has been emphasized previously [Shannon and Prewitt, 1969; Gaffney and Anderson, 1973; Sherman, 1988, 1991; Sherman and Jansen, 1995; Badro *et al.*, 2003, 2004; Li *et al.*, 2004]. Experimental studies using x-ray emission spectroscopy observed an electronic configuration change of the Fe in Fp with $x = 0.17$ between 60 and 70 GPa [Badro *et al.*, 2003]. In Pv with $y = 0.1$, two changes of the spin state of the iron at 70 GPa and 120 GPa

were interpreted as partial and full spin transitions of Fe(II), respectively [Badro *et al.*, 2004]. Using the same experimental technique on Pv of similar composition, as well as on an Al-bearing Pv, an independent study reported a gradual transformation of the iron from an initial high-spin state toward a lower spin state [Li *et al.*, 2004]. A study using synchrotron Mössbauer spectroscopy on Pv with $y = 0.05$ and $y = 0.1$ could not detect a change in the spin state of Fe(II) up to 120 GPa, however, indications for a changing spin state of Fe(III) around 70 GPa were reported [Jackson *et al.*, 2005]. All these experiments were performed at room temperature. Density-functional calculations of the electronic ground state of $(\text{FeO}_6)^{10-}$ and $(\text{FeMg}_{12}\text{O}_{14})^{2-}$ clusters suggested that the iron in Fp undergoes a pressure-dependent transition from a high-spin to a low-spin configuration [Sherman, 1988, 1991; Sherman and Jansen, 1995]. Density-functional calculations on wüstite [Cohen *et al.*, 1997] suggested that iron in Pv should remain in the high-spin state up to much higher pressures than found in the lower mantle. In both cases, the existence or absence of spin transitions was concluded from the pressure dependence of the electronic ground states. The temperature dependence of the transition pressure was extrapolated into the lower mantle using a Clausius-Clapeyron relation under the assumption that the transition persists at high temperatures [Sherman, 1988; Badro *et al.*, 2004; Li *et al.*, 2004]. Clearly, theoretical and experimental studies have provided valuable and often detailed information on the effect of pressure on the spin state of Fe in oxides and silicates, but the effect of temperature on the population of the spin states has not been considered.

[3] In this paper, we explore the temperature effect on spin states for minerals in Earth's lower mantle like Fp and Pv: the change from a high-spin to a low-spin state (or vice versa) can occur as a continuous crossover rather than a sudden transition. We will use crystal field theory and an extended Bragg-Williams mean-field theory to model the effects of the host mineral and of the iron-iron interactions on the 3d energy levels of the iron. If the Fe concentration of the mineral is of the order of 10 atomic% or less, the average iron-iron distance becomes large enough to result in negligible spin-spin interactions between neighboring iron atoms, and a continuous crossover or "spin equilibrium" is expected [Gütlich and Goodwin, 2004]. The application of these concepts allows us to express the influence of temperature and discuss the implications for a potential strati-

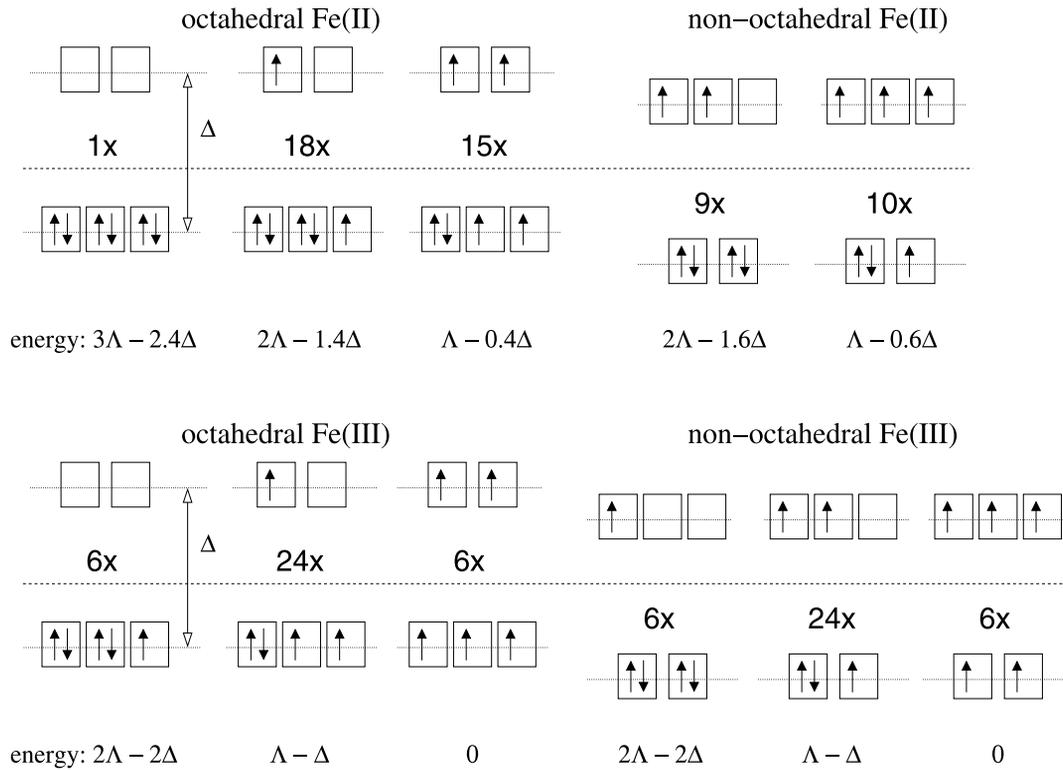


Figure 1. Configurations of the 3d electrons of Fe(II) and Fe(III), their degeneracies, and energies. The lowest energy configurations for octahedral and other symmetries are shown. The dashed lines represent the energy of the 3d electrons in a spherical symmetry.

fication of the lower mantle caused by the spin state of iron in minerals.

2. Interacting Spin Ensemble

[4] Isolated Fe(II) and Fe(III) in a host lattice experience a change in the electronic configuration of the 3d shell that is characteristic to the local environment [Burns, 1993]. The lowest energy configurations for the 3d electrons of Fe(II) and Fe(III) in nondistorted octahedral, tetrahedral, cubic, and dodecahedral environments are shown in Figure 1. The assumption of perfect symmetries allows us to describe the energy levels with only one parameter, $\omega = \Delta - \Lambda$, the difference between the crystal-field splitting energy $\Delta > 0$ and the exchange energy $\Lambda > 0$ (also called the spin-pairing energy) which is mostly independent of the local environment. For reduced symmetries, the energy degeneracy is partially or fully lifted, and a larger number of parameters is required. Values for Δ and Λ are typically 1 to 2 eV [Burns, 1993], and for temperatures of the mantle $\beta\Delta \gg 1$ and $\beta\Lambda \gg 1$ with $\beta = 1/(k_B T)$ as the inverse temperature. However the energy differences of the configurations in Figure 1 are multiples of ω , and each configuration may occur with reasonable probability in a thermally equilibrated material. If the interaction between the iron atoms is negligible, the influence of the host lattice is well described by values of ω . Such an assumption is justified for very dilute systems but will probably fail in iron-rich compounds like FeS or FeO. The minerals that are relevant to the lower mantle are not iron rich, but the influence of weak interactions may still

be noticeable. We will include interaction effects in a simple mean-field model based on the Bragg-Williams theory that was originally developed to describe order-disorder transitions in alloys [Williams, 1935].

[5] Assume that a total number of N iron atoms in the crystal can occupy energetically different spin states described by an index i and that the fraction of iron atoms in such a state is given by η_i with $\sum_i \eta_i = 1$. A combinatorial analysis and the use of Stirling's formula provides for the entropy of the ensemble

$$S = -k_B N \sum_i \eta_i \ln \frac{\eta_i}{g_i}, \quad (1)$$

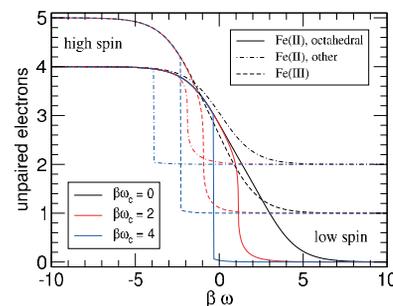


Figure 2. The average number of unpaired electrons for increasing values of the reduced coupling parameter $\beta\omega_c$ between low-spin atoms. Phase transitions occur for $\beta\omega_c > 2$, which would require quite large values of ω_c 0.31...0.52 eV for temperatures in Earth's lower mantle.

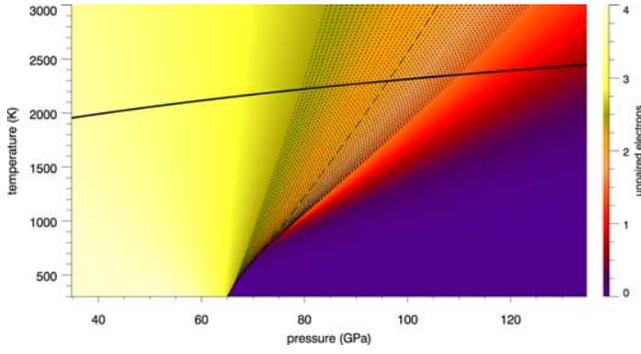


Figure 3. The average number of unpaired electrons for $\text{Mg}_{0.83}\text{Fe}_{0.17}\text{O}$ in the pressure-temperature sector. The dotted contours show the region between 1.5 and 2.5 unpaired electrons. This region clearly widens with increasing temperature. In particular, the iron spin state varies smoothly along a lower mantle geotherm [Brown and Shankland, 1981] given by the solid line. Parameters used for this calculation are given in the text.

where g_i is the degeneracy. In the spirit of the mean-field approach, the interaction energy is spatially averaged and the internal energy of the spin ensemble is approximated by

$$U = -N \sum_{ij} J_{ij} \eta_i \eta_j + N \sum_i E_i \eta_i, \quad (2)$$

where J_{ij} represents the coupling between an iron atom in spin state i and a nearest neighbor shell of iron atoms in the spin state j . The second term incorporates the crystal-field splitting, where E_i are the energy levels given in Figure 1. The equilibrium occupation of the spin states is determined from the requirement that the Helmholtz free energy $F = U - TS$ is stationary (minimal) with respect to the probabilities η_i . For a total number of s spin states, this is equivalent to a set of s equations $\partial F / \partial \eta_i = 0$ that are coupled by the normalization condition $\sum_i \eta_i = 1$, and therefore $s - 1$ independent equations are obtained. With the assumption $J_{ij} = J_{ji}$ we write

$$\beta(E_i - E_1) = 2\beta \sum_j (J_{ij} - J_{1j}) \eta_j - \ln \left(\frac{g_i \eta_i}{g_1 \eta_1} \right). \quad (3)$$

[6] With decreasing iron concentration $J_{ij} \rightarrow 0$, and we obtain the solution $\eta_i = g_i \exp(-\beta E_i) / Z$ with $Z = \sum_j g_j \exp(-\beta E_j)$, the spin-equilibrium with a continuous crossover between the different spin states.

[7] Depending on parameters like $\beta\omega$ and βJ_{ij} the free energy can become minimal for many solution vectors. The deepest minimum determines the physically meaningful solution, and the associated probabilities η_i can change suddenly, e.g., as a function of pressure. Therefore mathematically sharp spin transitions may occur. This behavior will be studied on an ensemble of Fe(II) atoms with an octahedrally symmetric environment. The crystal-field splitting is caused by the interaction of a given Fe atom with the surrounding atoms including other Fe atoms. Low-spin Fe(II) shows a smaller Fe-O distance R than high-spin Fe(II) corresponding to a larger value for the crystal-field

splitting. If we associate this energy difference with the coupling energy between low-spin iron atoms in the lattice we may write $\omega_c = a(x)\Delta(1 - R_L^5/R_H^5)$, where $\Delta \propto R^{-5}$ is the crystal-field splitting in the high-spin state and $a(x)$ depends on the iron concentration x . The term in parenthesis is of order unity for reasonable values of $R_L/R_H \approx 0.9 \dots 0.95$ [Burns, 1993]. The same R^{-5} dependence leads to $a(x) = x^{5/3}$, and we arrive at the estimate $\omega_c \approx x^{5/3}\Delta$.

[8] The effect of cooperative interactions was modelled by $J_{11} = \omega_c$, and all other coupling parameters set to zero. A numerical solution of this model represented by the average number of unpaired electrons calculated from the η_i values is shown in Figure 2. An increase in the coupling causes a decrease in the crossover region and leads to a mathematically sharp transition for $\beta\omega_c > 2$. With parameter pairs (Δ , T) typical for Earth's lower mantle we obtain for the iron concentrations (1 eV, 1800 K): 0.33, (1.5 eV, 1800 K): 0.26, (1 eV, 3000 K): 0.44, (1.5 eV, 3000 K): 0.35. For the much lower Fe concentrations in lower mantle minerals, sharp transitions between spin states seem therefore unlikely.

3. Geophysical Implications: Spin Crossover in Ferroperriclite

[9] Ferroperriclite, $\text{Mg}_{1-x}\text{Fe}_x\text{O}$, is a suitable mineral for a demonstration of the implications of the arguments raised in this paper. The ferrous iron atoms occupy a crystallographic site with octahedral symmetry. We will use a third-order Birch-Murnaghan equation of state (EOS) with $K_{0T} = (K_{0S} + T(\partial K_S / \partial T)) / (1 + \alpha\gamma T)$, $K_{0S} = 166$ GPa, and $\partial K_S / \partial P = 4.2$ [Kung et al., 2002] (values for $x = 0.17$). The effect of temperature will be modeled by $\partial K_S / \partial T = -0.019$ GPa/K [Sinogeikin et al., 2000], the Grüneisen parameter $\gamma = 1.35$ for the investigated range $0.67 \leq V/V_0 \leq 0.89$ [Speziale et al., 2001], and the thermal expansion coefficient $\alpha = (2.763 \cdot 10^{-3} \text{ K}^{-1} + 1.439 \cdot 10^{-8} T \cdot \text{K}^{-2})$ [Fiquet et al., 1999] (values for MgO). The results of $(\text{FeMg}_{12}\text{O}_{14})^{2-}$ cluster calculations [Sherman, 1991] indicate that the volume dependence of the crystal-field splitting is given by $\Delta \propto V^{-\xi}$ with $\xi = 1.5$, and therefore $\omega = \Delta(V) - \Lambda = \Delta_0 \{ (V_0/V)^\xi - (V_0/\bar{V})^\xi \}$, where \bar{V} is the volume at 65 GPa and 300 K, the reported spin transition in ferroperriclite

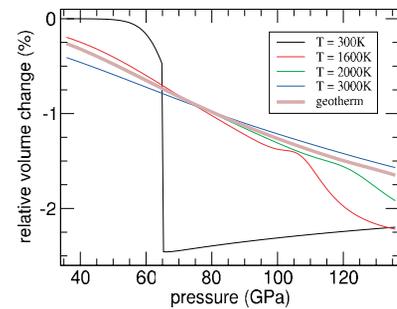


Figure 4. The relative volume change for $\text{Mg}_{0.83}\text{Fe}_{0.17}\text{O}$ with increasing pressure for several isotherms and a lower mantle geotherm [Brown and Shankland, 1981]. The volume change was calculated under the assumption that the equation of state does not change across the spin crossover. Parameters used for this calculation are given in the text.

[Badro *et al.*, 2003] (values for $x = 0.17$). The volume and the crystal-field splitting at ambient conditions are V_0 and $\Delta_0 = 1.35$ eV, respectively [Goto *et al.*, 1980]. The coupling parameter will be estimated by $\omega_c \approx x^\xi \Delta(V)$ and $\omega_{c0} \approx 0.09$ eV for $x = 0.17$ and ambient conditions. For the calculations, we used the slightly more conservative value of $\omega_{c0} = 0.1$ eV.

[10] In Figure 3, we show the results obtained with the presented mean-field theory for the number of unpaired electrons (shown as colored contours) and thus the spin state of Fe(II) in Fp in pressure-temperature space. At 300 K, our calculations predict that the change from the high-spin state (4 unpaired electrons, light yellow) to the low-spin state (0 unpaired electrons, blue) is sharp at 65 GPa, in agreement with recent x-ray emission spectroscopic measurements [Badro *et al.*, 2003]. However, under realistic lower mantle conditions, temperatures are much higher. Traversing the 65 GPa isobar with increasing temperature, one can see from Figure 3 that the spin transition becomes broader. A plausible lower mantle geotherm [Brown and Shankland, 1981] has been plotted in Figure 3 and reveals that under typical lower mantle temperatures, the number of unpaired electrons decreases over a pressure interval of more than 30 GPa (equivalent to a depth range of ≈ 700 km). In other words, there is no abrupt spin transition in Fp under lower mantle pressures and temperatures. The width of the spin-crossover region depends on the chosen parameter set. We tested our model by calculating the logarithmic derivatives $\partial \ln w / \partial \ln p$ of the width w with respect to each parameter p . The results in the format (parameter:derivative) are ξ : -1.7 , K_{0S} : 1.39 , $\partial K_S / \partial P$: 1.87 , $\partial K_S / \partial T$: 0.61 , α_1 : 0.38 , α_2 : -0.074 , γ : -0.32 , Δ_0 : -1.2 , ω_{c0} : -0.23 , transition pressure at 300 K: 0.8 and show that the model is robust in predicting a wide spin-crossover region.

[11] We also calculated the volume reduction with respect to the EOS using parameters determined well below the spin-transition pressure of 65 GPa. The ensemble of Fe atoms exerts an internal pressure given by $-\partial F / \partial V$, the volume derivative of the free energy defined by equations (1) and (2). The externally applied pressure corrected by this internal pressure gives rise to a slightly different volume via the EOS chosen for this material. The relative volume change due to this effect is shown in Figure 4. At 300 K, the volume reduction is about 2.3% and occurs within a narrow pressure range. However, the volume reduction along the geotherm is smaller and is a smoothly varying function with pressure.

[12] The behavior of Pv is more complex, as it may contain a significant fraction of ferric iron under lower mantle conditions [e.g., McCammon, 1997]. Even though the iron may occupy octahedrally and 8–12 coordinated sites, most of the iron likely resides in the 8–12 coordinated site, which is expected to show the smaller crystal-field splitting energy Δ [Burns, 1993]. The pressure derivative of the crystal-field splitting, $d\Delta/dP = \xi\Delta/K$, is generally smaller for Pv than for Fp because of its higher bulk modulus K and the smaller Δ . This will lead to a larger pressure range of a spin crossover in Pv than in Fp even at 300 K. Our model was constructed for highly symmetrical environments. For distorted environments, which might be expected in Pv and possibly also in Fp, the degeneracy of the energy levels of possible spin states is lifted, and the

spin crossover region is broadened even more. A gradual change in the spin state of iron in Pv to pressures above 100 GPa has recently been observed at 300 K using x-ray emission spectroscopy [Li *et al.*, 2004] and synchrotron Mössbauer spectroscopy [Jackson *et al.*, 2005].

[13] Independently of the character of the transition at 300 K, our calculations predict that the high-spin to low-spin crossover regions and the associated volume changes in Pv and Fp occur over a very broad pressure region at typical lower mantle temperatures. Our results imply that any chemical and physical changes associated with this crossover region, such as iron partitioning and elastic properties [e.g., Kobayashi *et al.*, 2004], would also be gradual. It is also possible that the oxidation state of iron in lower mantle minerals plays an important role in chemical changes associated with this large crossover region of several hundred kilometers depth. Furthermore, the hitherto unknown electronic properties of Fe in post-silicate perovskite [Murakami *et al.*, 2004] may lead to additional complexities at the base of the lower mantle.

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