Emergent high-spin state above 7 GPa in superconducting FeSe

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The local electronic and magnetic properties of superconducting FeSe have been investigated by $K\beta$ x-ray emission and simultaneous x-ray absorption spectroscopy (XAS) at the Fe K edge at high pressure and low temperature. Our results indicate a sluggish decrease of the local Fe spin moment under pressure up to 7 GPa, in line with previous reports, followed by a sudden increase at higher pressure. The magnetic surge is preceded by an abrupt change of the Fe local structure as observed by the decrease of the XAS preedge region intensity and corroborated by *ab initio* simulations. This pressure corresponds to a structural transition from the Cmma form to the denser Pbnm form with octahedral coordination of iron. Finally, the near-edge region of the XAS spectra shows a change before this transition at 5 GPa, corresponding well with the onset pressure of the sudden enhancement of T_c . Our results emphasize the delicate interplay between structural, magnetic, and superconducting properties in FeSe under pressure.

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FeSe is the simplest form of the Fe-based superconductors (FeSC), yet one of the more fascinating members, as seen in its many overlapping phases and their apparent correlation with the superconducting critical temperature T_c (Fig. 1). However, in spite of intensive research over the past years, the physics of FeSC remains still poorly understood because of the intricate coupling between structural, magnetic, and electronic degrees of freedom. Below 6-7 GPa, FeSe has a PbO-type tetragonal structure (P4/nmm) at ambient temperature which is slightly distorted to an orthorhombic structure (Cmma) [1] below the structural and magnetic transition temperatures (T_s and T_N). The low-temperature orthorhombic phase, denoted here as ortho-I, is characterized by nematic order with two distinct origins: below T_s , it is a second-order structural transition believed to be driven by orbital order [2,3]; while below T_N , it is a simultaneous first-order magnetostructural transition driven by stripe-type spin fluctuations [4–6]. A small magnetic moment with unknown antiferromagnetic (AFM) order exists below T_N , as seen with muon spin relaxation (μ SR) at 0.8–2.4 GPa [7–9] and with nuclear forward scattering until 4 GPa [6]. Transport measurements show that this AFM order exists to at least 6.3 GPa [10]. Above 6–7 GPa, a structural transition is observed [11-14] to a MnP-type orthorhombic phase (Pnma or equivalently Pbnm), denoted here as ortho-II, with the Fe ion site symmetry changing from tetrahedral (T_d) to octahedral (O_h) coordination. This transition was observed with ⁵⁷Fe Mössbauer spectroscopy above 7 GPa as an additional component in the spectrum; however, no magnetic hyperfine splitting is observed at 4.2 K [15]. The superconducting critical temperature in FeSe shows drastic changes with pressure. After an initial increase, T_c dips at the onset of AFM order, which is attributed to a reduction in the density of states due to a reconstruction of the Fermi surface [16,17]. Increasing the pressure further enhances both the magnetic and superconducting orders, with T_N reaching a maximum of 45–55 K [8,10] at \sim 4.2 GPa and T_c plateauing around 20 K below this pressure [10,18]. Above this pressure, magnetic order decreases and coincides with a sharp rise in T_c to a maximum around 37 K at \sim 6.2 GPa [10,12,13,15,19]. After this maximum, T_c has a discontinuous change from a positive to a negative slope, coinciding with the transition to the ortho-II phase [11]. An AFM region in the middle of a superconducting "dome" is quite unique, and the phase diagram shows their intimate connection as two T_c anomalies where the two intersect.

This macroscopic complexity reflects the strong coupling between charge, spin, and structure which in turn affects the superconducting properties. It is therefore of great importance to be able to probe these different degrees of freedom at the local level while exploring the phase diagram in the vicinity of the superconducting regions. X-ray spectroscopy in the hard x-ray range is well suited to that aim while being compatible with high-pressure conditions [20]. In particular, $K\beta$ (3 $p \rightarrow 1s$ transition) x-ray emission spectroscopy (XES) and x-ray absorption spectroscopy (XAS) at the K edge are well-established probes of the local electronic, magnetic, and structural properties. As a primarily atomic probe, XES can access the local magnetic moment of a selected atom regardless of the magnetic order, while XAS provides a view of the local

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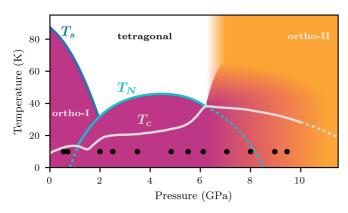


FIG. 1. Schematic P-T phase diagram of FeSe [10,11]. Tetragonal (white), ortho-I (purple), and ortho-II (orange) structures are described in the text. T_s , T_N , and T_c are the structural, magnetic, and superconducting transitions, with dashed lines representing extrapolations. P-T measurement points for XES/XAS are indicated with circles.

electronic and structural properties. In recent works [21,22], Kumar et al. and Chen et al. have used XES to probe Fe magnetism in FeSe under high pressure. Their results show a decrease of the Fe spin state with a discontinuous change in slope near the ortho- $I \rightarrow$ ortho-II transition. This is interpreted as a high-spin (HS) to low-spin (LS) transition [21] or a state with a smaller magnetic moment [22]. Their experiments were performed with polycrystalline samples including the hexagonal polymorph of FeSe (11% in the sample of Chen et al.) and at room temperature or only up to 8 GPa at 8 K for Kumar *et al.* In this work, we perform a high-pressure study of FeSe single crystals at 10 K until 9.5 GPa using Fe- $K\beta$ XES and XAS at the Fe K edge. Our results demonstrate a LS \rightarrow HS transition in the same pressure region as the ortho- $I \rightarrow$ ortho-II transition and the maximum of T_c . The spin state transition coincides with the structural transition and T_c slope change thus demonstrating the strong interplay between electronic and lattice degrees of freedom, adding to the remarkable properties of FeSe, and more generally to Fe superconductors. Here, we report a LS to HS transition under pressure in a 3d metal compound which challenges the classical crystal-field derived picture of spin transitions in these materials. We assign the low-spin state here to the atypical orbital-selective "Mottness" of the 3d electrons in Fe superconductors.

The XES/XAS experiment was performed on the GALAX-IES beamline [23] at the SOLEIL synchrotron facility. We used high-purity FeSe single crystals from the Institut Néel (Grenoble) grown by chemical vapor transport [24]. Pressure was applied using a membrane-driven diamond-anvil cell equipped with 1.2-mm-thick diamonds with 300- μ m culets. Several FeSe single crystals were loaded in a 150- μ m hole of a CuBe gasket, along with ruby chips for *in situ* pressure measurement [25] and silicone oil as a pressure transmitting medium. XES/XAS were measured with the spectrometer in a transmission geometry using a 1-m-radius spherically bent Ge(620) crystal analyzer and an avalanche photodiode detector arranged in the Rowland circle geometry. The total energy resolution at the Fe $K\beta$ line (\sim 7057 eV, $\theta_B = 79^\circ$) was 1.2 eV FWHM. The XES spectra were measured with 10-keV

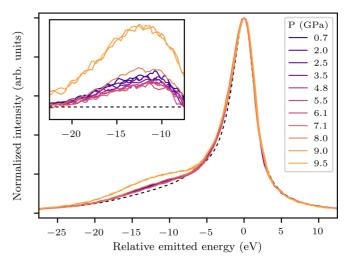


FIG. 2. Fe $K\beta$ x-ray emission in FeSe as a function of pressure measured at 10 K. The inset is a zoom of the satellite region showing the difference with the FeS₂ zero-spin reference [26], shown as a black dashed line in the main plot.

incident x rays. The XAS spectra was measured using the partial-fluorescence yield (PFY) method with the spectrometer fixed to the Fe $K\beta$ line. This technique leads to an intrinsic sharpening effect due to the shallower 3p core hole left in the final state with respect to the deeper 1s level [20]. The energies of our XAS spectra have been calibrated using the low-pressure XAS spectrum of FeSe from Chen *et al.* as a reference.

Our XES spectra measured at 10 K are shown in Fig. 2 for increasing pressure up to 9.5 GPa. The spectra are aligned to the main peak at 7057 eV and normalized to the maximum intensity. There is a weak satellite located around -12 eV from the main line, which is well established as a signature of the local magnetic moment [27]. The satellite can be seen more clearly by subtracting the zero-spin reference [26], FeS₂, as shown in the inset of Fig. 2. The fitted intensity of the satellite from the difference spectra is shown in Fig. 3 as red squares. FeSe exhibits a LS state with a gradually decreasing magnetic moment until 6–7 GPa. The magnetic moment starts to increase between 7 and 8 GPa and then jumps between 8 and 9 GPa to a HS state which plateaus around 9.5 GPa. The low-pressure behavior is consistent with previous results [21,22] and is understood to be due to band-structure effects in the compressed lattice; however, the sudden increase around 8 GPa was previously unobserved. The pressure region of the transformation of this LS → HS transition coincides with the ortho-I \rightarrow ortho-II coexistence region at low temperature [11].

To gain insight into the Fe local properties, we also measured high-resolution XAS at each pressure point after each XES measurement. The series of PFY-XAS spectra taken at the same pressures as XES is shown in Fig. 4, where the inset emphasizes the preedge region which is sensitive to the 3d states. The spectra are normalized and flattened using the software ATHENA [28]. Subsequently, the postedge points are binned to increase statistics. The prepeak feature A in FeSe at low pressure is mainly due to the dipole transitions from Fe 1s to Fe 3d-Se 4p hybrid bands as expected in T_d symmetry, with a smaller contribution from Fe quadrupole

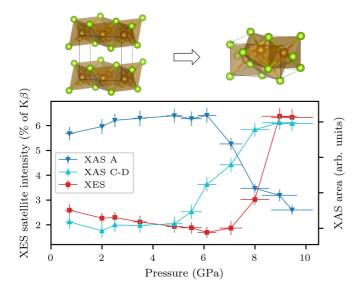


FIG. 3. Top: Schematic of ortho-II to ortho-II structural transition. Bottom: Pressure dependence of XES and XAS spectra at 10 K. Left scale: Intensity of $K\beta$ satellite shown as red squares. Right scale: Area of the Fe preedge feature A shown as blue inverted triangles and area between XAS features C and D shown as cyan triangles (scaled and offset).

 $1s \rightarrow 3d$ transitions [22,29–31]. Fe 3d-Se 4p hybridization is sensitive to the local geometry, therefore the prepeak can be used primarily to study the pressure evolution of the structure (and the spin state as discussed below). To clarify its pressure dependence, it was fitted by a Gaussian line shape after subtraction of a background (error function + linear) to account for the rising edge. This preedge area is shown as blue inverted triangles in Fig. 3 using the right scale and the label "XAS A." As FeSe is compressed the prepeak area slowly increases until 6 GPa, above which it decreases with a much sharper slope until our highest pressure point at 9.5 GPa. The decreased intensity is due to a reduced hybridization as the

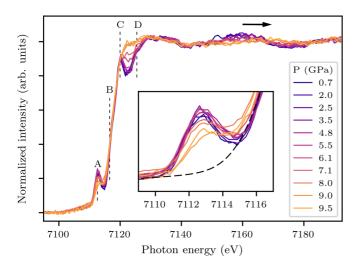


FIG. 4. Partial-fluorescence yield x-ray absorption at the Fe K edge in FeSe as a function of pressure measured at 10 K. The inset is a zoom of the preedge region, with the 0 GPa background (error function + linear) shown with a dashed line.

Fe-Se bond lengths increase by $\approx 5\%$ [13] and the coordination of iron increases during the ortho-I $(T_d) \rightarrow$ ortho-II (O_h) transition [11]. The increased coordination is also supported by the change of the hyperfine splitting at high pressure [15]. The remarkable parallel evolution of the magnetic (XES) and structural (XAS) properties shows the interplay between the electronic and lattice degrees of freedom. The fact that the structural transition precedes the magnetic transition suggests that it is not magnetically driven.

The absorption features B, C, and D can be assigned as Fe dipole $1s \rightarrow 4p$ transitions, where C and D also have significant contribution from Fe 1s to Fe 4p-Se 3d hybrid band transitions [22,29-31]. The features at higher energies are dominated by multiple scattering of the photoelectron with the nearest neighbors. The feature around 7160 eV shows a clear shift (arrow in Fig. 4) supporting the change of local structure we see with the prepeak. A significant spectral change is observed in the region between features C and D, even before the structural and magnetic transitions. The change is shown in Fig. 3 (cyan triangles) by taking the area between C and D. The onset of the increased intensity corresponds remarkably well with the sudden increase in T_c around 5 GPa [10]. The trend is continuous until the ortho-II dominant phase at 9.5 GPa, which puts into question the possible connection between ortho-II and superconductivity since ortho-II is not believed to support superconductivity [11], and a negative $d\rho/dT$ suggests it is a semiconductor [10,15].

To further understand the structural and electronic transitions, the XAS spectra were simulated using the FDMNES code [32]. A cluster radius of 12 Å (10 Å) for the low pressure (high pressure) reached full convergence. Spin-orbit coupling was included by relativistic effect corrections and the Fermi level was set self-consistently [33]. The crystal structures were taken from Ref. [1] for the low-pressure ortho-I phase and from Ref. [14] for the high-pressure ortho-II phase. To simulate the pressure-induced spin-state transition, the calculations in both phases were carried out using either a LS $(3d_{\uparrow}^33d_{\downarrow}^3)$ or a HS $(3d_{\uparrow}^53d_{\downarrow}^1)$ ground-state configuration.

The results are shown in Fig. 5 along with the experimental spectra measured at 0.7 and 9.0 GPa. We do not expect to yield an accurate description of the d electronic structure in FeSe without including correlations—more accurate approaches show that many-body effects lead to a renormalization of the dbands and changes of the d density of states close to the Fermi edge [34]—but the K edge should be well described since it mostly connects to a single-particle picture. We find that the calculations for both the low-pressure and high-pressure phases compare well to the experimental spectra in the preedge and near-edge regions. We notice, however, that the intensity of the preedge for the high-pressure phase is overestimated. The preedge intensity is strongly dependent on minor distortions of the metal site which favor or disfavor dipolar transitions to ligand p states in addition to the weaker quadrupolar transition to the d states. On the contrary, the preedge shape is mostly related to the orbitals' energy splitting and spin state, therefore it is a much better fingerprint of the local 3d electronic properties. The preedge shape matches very well with a high-spin (low-spin) configuration for the high-pressure (low-pressure) phase. The less-pronounced prepeak in ortho-II

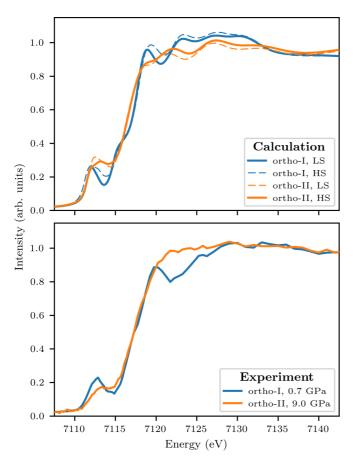


FIG. 5. Comparison of calculated and measured XAS spectra at the Fe K edge in FeSe. Top panel: Calculated XAS spectra for the ortho-I phase (Cmma) and the ortho-II phase (Pnma) in low-spin (LS) and high-spin (HS) configurations. Thick lines represent the best agreement with experiment. Bottom panel: Experimental XAS spectra measured at 10 K.

is consistent with high-spin Fe^{II} in an octahedral environment [35]. This agrees with our XES results that there is a transition to a high-spin state above 7 GPa at 10 K in FeSe.

The main outcome of our study is the original observation of a high-spin state above 7 GPa as FeSe adopts the ortho-II phase. This follows the initial decrease of the local moment in the ortho-I phase which has been reported elsewhere [21,22] and confirmed here. Both XAS and XES indicate that the magnetic surge at high pressure is consistent with a pressure-induced LS to HS transition. The electronic change is likely resulting from the change of Fe site symmetry: in the ortho-I phase, Fe occupies a tetrahedral (T_d) site that becomes increasingly distorted with pressure, whereas Fe sits in an octahedral (O_h) site in the ortho-II phase [11–13]. At first glance, the spin state of FeSe is surprising. According to ligand field theory, one would expect Fe to be high spin in the ortho-I phase since tetrahedral coordination normally favors a high-spin state. However, the Fe-based superconductor's electronic structure was also shown to be controlled by the correlated and specific Mottness of d electrons with band-dependent correlations yielding e_g states less correlated than t_{2g} states [36]. We suggest that the T_d low-spin state is stabilized because it minimizes the on-site electron correlations by filling preferentially the e_{g} bands. At high pressure, correlation becomes less effective as the bandwidth broadens which eventually allows a high-spin configuration as the Fe symmetry turns O_h . In line with recent band-structure calculations [37], we can further argue that the exceptional stability of the magnetism at high pressure in FeSe could be aided by the increased three-dimensional character of the compressed lattice and change of the Fermi surface.

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^[1] S. Margadonna, Y. Takabayashi, M. T. McDonald, K. Kasperkiewicz, Y. Mizuguchi, Y. Takano, A. N. Fitch, E. Suard, and K. Prassides, Chem. Commun., 5607 (2008).

^[2] S.-H. Baek, D. V. Efremov, J. M. Ok, J. S. Kim, J. van den Brink, and B. Büchner, Nat. Mater. 14, 210 (2015).

^[3] P. Massat, D. Farina, I. Paul, S. Karlsson, P. Strobel, P. Toulemonde, M.-A. Méasson, M. Cazayous, A. Sacuto, S. Kasahara, T. Shibauchi, Y. Matsuda, and Y. Gallais, Proc. Natl. Acad. Sci. USA 113, 9177 (2016).

^[4] Q. Wang, Y. Shen, B. Pan, Y. Hao, M. Ma, F. Zhou, P. Steffens, K. Schmalzl, T. R. Forrest, M. Abdel-Hafiez, X. Chen, D. A. Chareev, A. N. Vasiliev, P. Bourges, Y. Sidis, H. Cao, and J. Zhao, Nat. Mater. 15, 159 (2016).

^[5] P. S. Wang, S. S. Sun, Y. Cui, W. H. Song, T. R. Li, R. Yu, H. Lei, and W. Yu, Phys. Rev. Lett. 117, 237001 (2016).

^[6] K. Kothapalli, A. E. Böhmer, W. T. Jayasekara, B. G. Ueland, P. Das, A. Sapkota, V. Taufour, Y. Xiao, E. Alp, S. L. Bud'ko, P.

C. Canfield, A. Kreyssig, and A. I. Goldman, Nat. Commun. 7, 12728 (2016).

^[7] M. Bendele, A. Amato, K. Conder, M. Elender, H. Keller, H.-H. Klauss, H. Luetkens, E. Pomjakushina, A. Raselli, and R. Khasanov, Phys. Rev. Lett. 104, 087003 (2010).

^[8] M. Bendele, A. Ichsanow, Y. Pashkevich, L. Keller, T. Strässle, A. Gusev, E. Pomjakushina, K. Conder, R. Khasanov, and H. Keller, Phys. Rev. B 85, 064517 (2012).

^[9] R. Khasanov, Z. Guguchia, A. Amato, E. Morenzoni, X. Dong, F. Zhou, and Z. Zhao, Phys. Rev. B 95, 180504 (2017).

^[10] J. P. Sun, K. Matsuura, G. Z. Ye, Y. Mizukami, M. Shimozawa, K. Matsubayashi, M. Yamashita, T. Watashige, S. Kasahara, Y. Matsuda, J. Q. Yan, B. C. Sales, Y. Uwatoko, J. G. Cheng, and T. Shibauchi, Nat. Commun. 7, 12146 (2016).

^[11] V. Svitlyk, M. Raba, V. Dmitriev, P. Rodière, P. Toulemonde, D. Chernyshov, and M. Mezouar, Phys. Rev. B 96, 014520 (2017).

- [12] G. Garbarino, A. Sow, P. Lejay, A. Sulpice, P. Toulemonde, M. Mezouar, and M. Nunez-Regueiro, Europhys. Lett. 86, 27001 (2009).
- [13] S. Margadonna, Y. Takabayashi, Y. Ohishi, Y. Mizuguchi, Y. Takano, T. Kagayama, T. Nakagawa, M. Takata, and K. Prassides, Phys. Rev. B 80, 064506 (2009).
- [14] R. S. Kumar, Y. Zhang, S. Sinogeikin, Y. Xiao, S. Kumar, P. Chow, A. L. Cornelius, and C. Chen, J. Phys. Chem. B 114, 12597 (2010).
- [15] S. Medvedev, T. M. McQueen, I. A. Troyan, T. Palasyuk, M. I. Eremets, R. J. Cava, S. Naghavi, F. Casper, V. Ksenofontov, G. Wortmann, and C. Felser, Nat. Mater. 8, 630 (2009).
- [16] T. Terashima, N. Kikugawa, A. Kiswandhi, D. Graf, E.-S. Choi, J. S. Brooks, S. Kasahara, T. Watashige, Y. Matsuda, T. Shibauchi, T. Wolf, A. E. Böhmer, F. Hardy, C. Meingast, H. v. Löhneysen, and S. Uji, Phys. Rev. B 93, 094505 (2016).
- [17] T. Terashima, N. Kikugawa, S. Kasahara, T. Watashige, Y. Matsuda, T. Shibauchi, and S. Uji, Phys. Rev. B 93, 180503 (2016).
- [18] K. Miyoshi, K. Morishita, E. Mutou, M. Kondo, O. Seida, K. Fujiwara, J. Takeuchi, and S. Nishigori, J. Phys. Soc. Jpn. 83, 013702 (2014).
- [19] D. Braithwaite, B. Salce, G. Lapertot, F. Bourdarot, C. Marin, D. Aoki, and M. Hanfland, J. Phys.: Condens. Matter 21, 232202 (2009).
- [20] J.-P. Rueff and A. Shukla, Rev. Mod. Phys. 82, 847 (2010).
- [21] R. S. Kumar, Y. Zhang, Y. Xiao, J. Baker, A. Cornelius, S. Veeramalai, P. Chow, C. Chen, and Y. Zhao, Appl. Phys. Lett. **99**, 061913 (2011).
- [22] J. M. Chen, S. C. Haw, J. M. Lee, T. L. Chou, S. A. Chen, K. T. Lu, Y. C. Liang, Y. C. Lee, N. Hiraoka, H. Ishii, K. D. Tsuei, E. Huang, and T. J. Yang, Phys. Rev. B 84, 125117 (2011).
- [23] J.-P. Rueff, J. M. Ablett, D. Céolin, D. Prieur, T. Moreno, V. Balédent, B. Lassalle, J. E. Rault, M. Simon, and A. Shukla, J. Synchrotron Radiat. 22, 175 (2015).

- [24] S. Karlsson, P. Strobel, A. Sulpice, C. Marcenat, M. Legendre, F. Gay, S. Pairis, O. Leynaud, and P. Toulemonde, Supercond. Sci. Technol. 28, 105009 (2015).
- [25] A. Dewaele, M. Torrent, P. Loubeyre, and M. Mezouar, Phys. Rev. B 78, 104102 (2008).
- [26] J. F. Lin, H. Watson, G. Vankó, E. E. Alp, V. B. Prakapenka, P. Dera, V. V. Struzhkin, A. Kubo, J. Y. Zhao, C. McCammon, and W. J. Evans, Nat. Geosci. 1, 688 (2008).
- [27] G. Peng, F. M. F. de Groot, K. Hämäläinen, J. A. Moore, X. Wang, M. M. Grush, J. B. Hastings, D. P. Siddons, W. H. Amstrong, O. C. Mullins, and S. P. Cramer, J. Am. Chem. Soc. 116, 2914 (1994).
- [28] B. Ravel and M. Newville, J. Synchrotron Radiat. 12, 537 (2005).
- [29] J. M. Chen, S. C. Haw, J. M. Lee, S. A. Chen, K. T. Lu, M. J. Deng, S. W. Chen, H. Ishii, N. Hiraoka, and K. D. Tsuei, J. Chem. Phys 137, 244702 (2012).
- [30] M. Bendele, C. Marini, B. Joseph, L. Simonelli, P. Dore, S. Pascarelli, M. Chikovani, E. Pomjakushina, K. Conder, N. L. Saini, and P. Postorino, J. Phys.: Condens. Matter 25, 425704 (2013).
- [31] B. Joseph, A. Iadecola, L. Simonelli, Y. Mizuguchi, Y. Takano, T. Mizokawa, and N. L. Saini, J. Phys.: Condens. Matter 22, 485702 (2010).
- [32] S. A. Guda, A. A. Guda, M. A. Soldatov, K. A. Lomachenko, A. L. Bugaev, C. Lamberti, W. Gawelda, C. Bressler, G. Smolentsev, A. V. Soldatov, and Y. Joly, J. Chem. Theory Comput. 11, 4512 (2015).
- [33] O. Bunău and Y. Joly, J. Phys.: Condens. Matter 21, 345501 (2009).
- [34] M. Aichhorn, S. Biermann, T. Miyake, A. Georges, and M. Imada, Phys. Rev. B 82, 064504 (2010).
- [35] T. E. Westre, P. Kennepohl, J. G. DeWitt, B. Hedman, K. O. Hodgson, and E. I. Solomon, J. Am. Chem. Soc. 119, 6297 (1997).
- [36] L. de' Medici, G. Giovannetti, and M. Capone, Phys. Rev. Lett. 112, 177001 (2014).
- [37] D. S. Parker, Sci. Rep. 7, 3388 (2017).