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# Vibrational blue shift of coordinated N<sub>2</sub> in [Fe<sub>3</sub>O(OAc)<sub>6</sub>(N<sub>2</sub>)<sub>n</sub>]<sup>+</sup>: “non-classical” dinitrogen complexes†

Johannes Lang,\* Jennifer Mohrbach, Sebastian Dillinger, Joachim M. Hewer and Gereon Niedner-Schatteburg

**We present “non-classical” dinitrogen Fe(III) oxo acetate complexes *in vacuo* utilizing Infrared Photodissociation (IR-PD) at cryo temperatures. The IR-PD spectra reveal a blue shift of the N<sub>2</sub> stretching vibration frequencies in the complexes. Density Functional Theory (DFT) calculations confirm the experiments and indicate strengthened N–N bonds due to pronounced σ bonding and a lack of π back donation.**

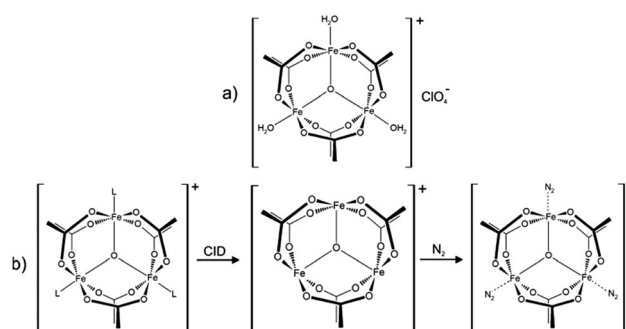
The discovery of transition metal dinitrogen complexes in 1965<sup>1</sup> launched the ever growing research field of N<sub>2</sub> coordination chemistry.<sup>2,3</sup> One of the main goals is the conversion of N<sub>2</sub> to ammonia under mild conditions<sup>4</sup> by homogenous catalysis.<sup>5,6</sup> The basic idea is to weaken N–N bonds upon coordination to enable subsequent functionalization. Precursors to N<sub>2</sub> activation manifest by elongated N–N distances and by red shifting of stretching frequencies<sup>7,8</sup> with respect to the vibration of the free N<sub>2</sub> molecule. Such bond activation and vibrational red shifts have been interpreted before, *e.g.* in terms of the Dewar–Chatt–Duncanson (DCD) model,<sup>9</sup> and the Blyholder surface coordination (BSC)<sup>10</sup> model. Theoretical as well as spectroscopic investigations confirmed these interpretations.<sup>11</sup> However, FT-IR studies on N<sub>2</sub> coordinated to Lewis acid centers in zeolites<sup>12</sup> and alumina<sup>13</sup> revealed blue shifted vibrational N<sub>2</sub> stretching frequencies (up to 24 cm<sup>-1</sup>). This indicates N–N bond strengthening rather than weakening – much beyond these established models. A similar effect has been observed in the case of isoelectronic carbon monoxide: so called “non-classical” metal carbonyl complexes<sup>14,15</sup> exhibit blue shifted vibrational CO stretching frequencies (up to 138 cm<sup>-1</sup>).<sup>16</sup> Gas phase studies of isolated non-classical metal carbonyl clusters<sup>17,18</sup> proved the intrinsic molecular origin of this effect. The cause for the CO blue shift was a topic of debate<sup>19</sup> and is now understood in terms of an interplay between π back donation and electrostatic effects.<sup>20</sup> It is paramount to characterize the geometrical and electronic structures of such complexes in

order to acquire a fundamental insight into the prevailing interactions leading to such non-classical behaviour.

The combination of Electrospray Ionization Mass Spectrometry<sup>21</sup> (ESI-MS) and infrared (IR) laser spectroscopy is suitable for the characterization of isolated coordination complexes with defined stoichiometry. Infrared (Multiple) Photon Dissociation (IR-(M)PD) provides direct access to structural and vibrational information, both under cryogenic conditions<sup>22,23</sup> and at room temperature.<sup>24</sup> The experimental results and dedicated *ab initio* calculations are combined in order to obtain detailed insight into the geometrical structure and intrinsic properties of the isolated molecular ions.

In this work we investigate isolated dinitrogen complexes [Fe<sub>3</sub>O(OAc)<sub>6</sub>(N<sub>2</sub>)<sub>n</sub>]<sup>+</sup> (*n* = 1–3, OAc = CH<sub>3</sub>CO<sub>2</sub><sup>-</sup>, *cf.* Scheme 1), which exhibit a significant blue shift (17 cm<sup>-1</sup>) of N<sub>2</sub> vibrations in the complex with respect to the free N<sub>2</sub> molecule. We examine the N<sub>2</sub> coordination in detail to rationalize the blue shift and N–N bond strengthening. This is, to the best of our knowledge, the first report on the “non-classical” behaviour of isolated N<sub>2</sub> complexes.

We utilize a customized Fourier Transform-Ion Cyclotron Resonance (FT-ICR)-mass spectrometer (Apex Ultra, Bruker Daltonics) equipped with an ESI ion source (Apollo 2, Bruker).



**Scheme 1** (a) Molecular structure of the precursor [Fe<sub>3</sub>O(OAc)<sub>6</sub>(H<sub>2</sub>O)<sub>3</sub>](ClO<sub>4</sub>)<sup>+</sup> salt. In solution the water molecules exchange with solvent molecules (L = *e.g.* acetonitrile). (b) ESI-MS reveals the formation of [Fe<sub>3</sub>O(OAc)<sub>6</sub>(L)<sub>n</sub>]<sup>+</sup> (*n* = 0, 1, 2, 3). Collision induced dissociation (CID) of L yields the under-coordinated [Fe<sub>3</sub>O(OAc)<sub>6</sub>]<sup>+</sup> complex, which binds 1–3 N<sub>2</sub> molecules at 26 K.

Fachbereich Chemie and Forschungszentrum OPTIMAS, Technische Universität Kaiserslautern, 67663 Kaiserslautern, Germany. E-mail: jlang@chemie.uni-kl.de  
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We coupled the ICR cell with a KTP/KTA optical parametric oscillator/amplifier (OPO/A) IR laser system (LaserVision). Optimized minimum energy structures and linear IR absorption spectra were calculated at the B3LYP<sup>25</sup> level of theory using cc-pVTZ basis sets<sup>26</sup> and Stuttgart RSC 1997<sup>27</sup> effective core potential basis sets (Gaussian 09<sup>28</sup>). We present calculations with 15 unpaired alpha electrons yielding a spin multiplicity of 16 with other multiplicities (2–18) found to be significantly less stable (*cf.* Fig. S12, ESI†). We scale the calculated frequencies with two different scaling factors: one scaling factor (0.951) is specifically designed to elucidate N<sub>2</sub> stretching bands in [Fe<sub>3</sub>O(OAc)<sub>6</sub>(N<sub>2</sub>)<sub>n</sub>]<sup>+</sup>. It scales the calculated N<sub>2</sub> stretching frequencies such that a calculated free N<sub>2</sub> stretching vibration frequency matches the experimental value<sup>29</sup> of 2330 cm<sup>-1</sup>. This approach conveniently reveals any effects of Fe–N<sub>2</sub> coordination on N<sub>2</sub> stretching frequencies. A second unspecific scaling factor (0.986) is applied for all other bands below 1800 cm<sup>-1</sup>. It scales the calculated asymmetric carboxylate stretching vibration frequencies of [Fe<sub>3</sub>O(OAc)<sub>6</sub>(N<sub>2</sub>)<sub>1</sub>]<sup>+</sup> to match our own experimental value of 1587 cm<sup>-1</sup>. Unscaled spectra are provided in the ESI† (*cf.* Fig. S4).

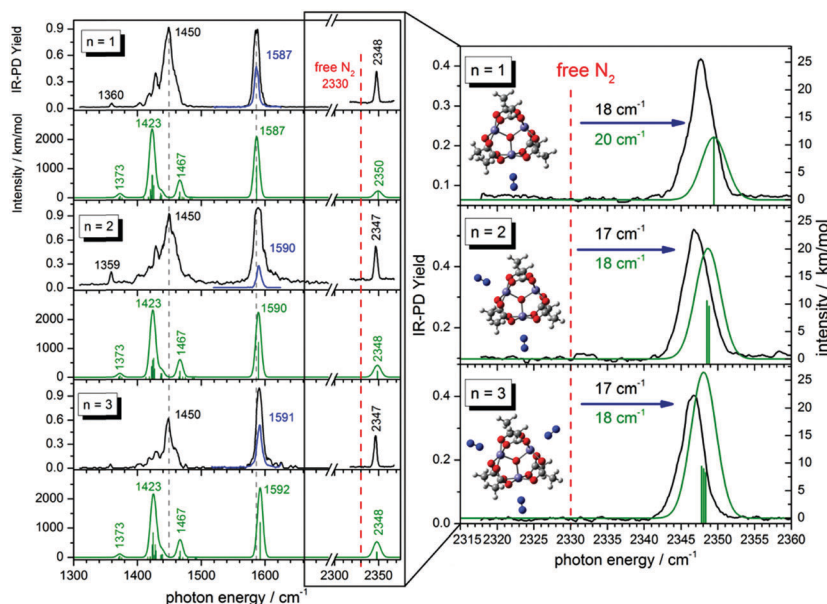
Upon spraying the sample solution and recording mass spectra in the ICR-cell, we observe a series of isotopic peaks matching convincingly with simulated isotopic patterns (*cf.* Fig. S1, ESI†). We assign those peaks to [Fe<sub>3</sub>O(OAc)<sub>6</sub>(L)<sub>n</sub>]<sup>+</sup> (L = H<sub>2</sub>O, acetonitrile, acetic acid; *n* = 0, 1, 2, 3). Elimination of L by Collision Induced Dissociation (CID) and subsequent coordination of N<sub>2</sub> in the hexapole at cryo temperatures (26 K) leads to the formation of [Fe<sub>3</sub>O(OAc)<sub>6</sub>(N<sub>2</sub>)<sub>n</sub>]<sup>+</sup> (*cf.* Scheme 1b and Fig. S2, ESI†). Note that we observe *n*<sub>max</sub> = 3, thus “titrating” the three available Fe

coordination sites. The calculated Gibbs energies at various temperatures reveal N<sub>2</sub> binding energies of 11, 9, and 8 kJ mol<sup>-1</sup> for the first, second, and third N<sub>2</sub> (26 K; BSSE corrected, *cf.* Fig. S3, ESI†). The N<sub>2</sub> binding Gibbs energies diminish with increasing temperature, vanishing above 80 K.

We recorded IR-PD spectra of cryocooled [Fe<sub>3</sub>O(OAc)<sub>6</sub>(N<sub>2</sub>)<sub>n</sub>]<sup>+</sup> (*n* = 1, 2, 3, Fig. 1, black traces) and conducted DFT simulations to obtain their linear IR absorption spectra (Fig. 1, green traces). The DFT calculations reveal optimized minimum structures as depicted in the insets of Fig. 1. We observe several bands between 1300 cm<sup>-1</sup> and 1500 cm<sup>-1</sup>, coinciding with the predicted CH<sub>3</sub> bending modes of the acetate ligands. While the calculated band frequencies around 1423 cm<sup>-1</sup> and 1467 cm<sup>-1</sup> agree well with the IR-PD spectrum, the calculated intensities differ significantly. Switching the DFT functional from B3LYP to PBE0 yields a much better match of IR intensities but significant deviations of calculated and observed vibrational frequencies (*cf.* Fig. S5, ESI†). Neither of these empirical functionals predicts both entities correctly. In the following we utilize the B3LYP results for further discussion.

The amount of N<sub>2</sub> coordination (*n* = 1, 2, 3) has no significant influence on the frequency and intensity of the CH<sub>3</sub> bending bands (neither in the IR-PD experiments nor in the DFT calculations). This finding likely originates from the spatial separation of the affected methyl groups from the Fe–N<sub>2</sub> coordination sites.

We find a strong IR-PD band at 1587, 1590, and 1591 cm<sup>-1</sup> in the cases of *n* = 1, 2, 3. We assign this band to carboxylic CO stretching bands of the six coordinated acetate ligands. The *n* = 1 CO stretching band is red shifted by ≈ 3–4 cm<sup>-1</sup> with



**Fig. 1** Left: IR-PD spectra of [Fe<sub>3</sub>O(OAc)<sub>6</sub>(N<sub>2</sub>)<sub>n</sub>]<sup>+</sup> (*n* = 1, 2, 3) at 26 K (black and blue curves) and calculated IR absorption spectra of optimized [Fe<sub>3</sub>O(OAc)<sub>6</sub>(N<sub>2</sub>)<sub>n</sub>]<sup>+</sup> (*n* = 1–3) (green curves) in the range of 1300–2400 cm<sup>-1</sup>. The blue IR-PD spectrum shows the CO stretching band (recorded with highly attenuated laser power to avoid saturation effects). The calculations were performed at the B3LYP/cc-pVTZ (H,C,N,O) and Stuttgart 1997 ECP (Fe) level of theory. The multiplicity is 16 and frequencies are scaled with 0.951 (0.986) above 2300 cm<sup>-1</sup> (below 2300 cm<sup>-1</sup>). Calculated stick spectra were convoluted with a Gaussian envelope of FWHM = 7 cm<sup>-1</sup>. Right: A zoom into the N<sub>2</sub> stretching vibration region. Calculated lines were convoluted with a Gaussian envelope of FWHM = 3.5 cm<sup>-1</sup>. Insets show associated geometry optimized structures.

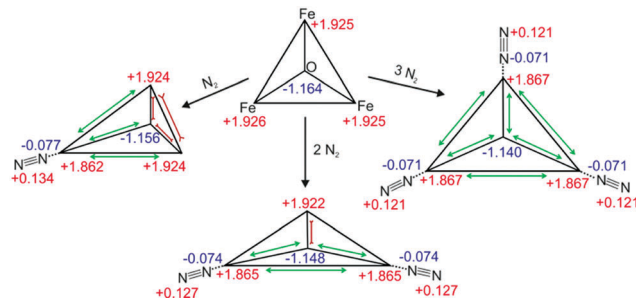


respect to the  $n = 2, 3$  bands, and with respect to the corresponding band of the free acetate anion<sup>30</sup> at  $1591\text{ cm}^{-1}$ . The DFT calculated asymmetric carboxylate stretching frequencies – scaled to match the experiment at  $n = 1$  – reproduce well the reduced redshift of the experiments on  $n = 2, 3$ . The coordinated carboxylate groups thus sense  $\text{N}_2$  coordination through their common Fe centers. We provide an illustrative visualization of the calculated displacement vectors of these modes in Fig. S6 of the ESI.† Note that the  $n = 1$  and 2 coordinations lift the sixfold degeneracy of these asymmetric stretching bands of the six coordinated acetate ligands. The recorded bands (slightly broadened to  $\text{FWHM} \approx 7\text{ cm}^{-1}$ ) may well contain the predicted splittings of  $\approx 0.5\text{ cm}^{-1}$  – yet unresolved (*cf.* Fig. S7, ESI†).

We observe sharp bands ( $\text{FWHM} \approx 3.5\text{ cm}^{-1}$ ) above  $2300\text{ cm}^{-1}$  revealing the IR active  $\text{N}_2$  stretching vibrations of  $[\text{Fe}_3\text{O}(\text{OAc})_6(\text{N}_2)_n]^+$  (*cf.* the displacement vector visualization of these modes in Fig. S8, ESI†). Isotope labeling of the  $\text{N}_2$  molecule confirms our assignment of these IR-PD bands to the  $\text{N}_2$  stretching frequencies. We observe a red shift of the  $^{15}\text{N}$ – $^{15}\text{N}$  vibration band relative to the  $^{14}\text{N}$ – $^{14}\text{N}$  vibration band ( $79\text{ cm}^{-1}$ ; *cf.* Fig. S17, ESI†) and the same blue shift with respect to the free  $\text{N}_2$  in both isotopomers.

In all cases ( $n = 1, 2, 3$ ) the solitary  $^{14}\text{N}_2$  stretching bands of the experimental IR-PD spectra shift to the blue ( $17$ – $18\text{ cm}^{-1}$ ) with respect to the (IR inactive) stretching frequency of the free  $^{14}\text{N}_2$  molecule ( $2330\text{ cm}^{-1}$ ).<sup>29</sup> These blue shifts indicate N–N bond strengthening upon coordination. Our DFT calculations predict this blue shift remarkably well ( $18$ – $20\text{ cm}^{-1}$ ). Multiple coordinated  $\text{N}_2$  molecules have almost identical stretching frequencies. There seems to be no coupling between the  $\text{N}_2$  molecules adsorbing at distinct, equivalent Fe sites. In contrast to the carboxylate asymmetric stretching bands the  $\text{N}_2$  stretching bands shift slightly to lower frequencies with increasing  $n$  (IR-PD spectrum:  $1\text{ cm}^{-1}$ ; DFT:  $1.5\text{ cm}^{-1}$ ). However, in all three cases ( $n = 1$ – $3$ ) the  $\text{N}_2$  stretching band is blue shifted with respect to the free  $\text{N}_2$ .

The DFT calculations reveal subtle distortions of the triangular  $\text{Fe}_3\text{O}$ -core upon coordination of  $\text{N}_2$  (*cf.* Scheme 2 and Table S2, ESI†). Each  $\text{N}_2$  molecule binds end on to the respective Fe atom. In general, the coordination of  $\text{N}_2$  enlarges Fe–Fe distances and Fe– $\text{O}_{\text{central}}$  bond lengths (by  $0.03\text{ \AA}$  resp.  $0.01\text{ \AA}$ ) of those bonds which involve the  $\text{N}_2$  coordinating Fe-center. All the other Fe–Fe distances and Fe– $\text{O}_{\text{central}}$  bond lengths shorten by approx.  $0.03\text{ \AA}$  or  $0.01\text{ \AA}$ . Natural Population Analysis (NPA) of  $[\text{Fe}_3\text{O}(\text{OAc})_6(\text{N}_2)_n]^+$  ( $n = 0, 1, 2, 3$ ) reveals the dependence upon  $n$  of local charge densities within the triangular  $\text{Fe}_3\text{O}$ -core (*cf.* Scheme 2). The Fe centers gain  $0.058$ – $0.064\text{ e}$  electron density per coordinating  $\text{N}_2$ . The non-coordinated N atoms donate  $0.121$ – $0.134\text{ e}$  electron density in about equal parts to the coordinating N and Fe atoms. Free, non-polar  $\text{N}_2$  molecules thus polarize and oxidize significantly upon Fe coordination, both magnitudes decreasing with  $n$  – as does the blue shift of the  $\text{N}_2$  stretching bands. Note that the N–N bond length contracts slightly upon coordination ( $-0.002\text{ \AA}$ ). The high positive charge on the Fe centers ( $1.862$ – $1.922\text{ e}$ ) diminishes their electron donating capability and thus inhibits the  $\pi$  back donation. Such charge effects seem to increase the “non-classical” effect in metal carbonyl complexes.<sup>31</sup>



Scheme 2 Calculated geometries and natural charge distributions of the  $\text{Fe}_3\text{O}$ -core and the coordinated  $\text{N}_2$  in  $[\text{Fe}_3\text{O}(\text{OAc})_6(\text{N}_2)_n]^+$  ( $n = 0, 1, 2, 3$ ).

Non covalent interaction (*cf.* Fig. 2 for  $n = 1$  and Fig. S11 for  $n = 2, 3$ , ESI†) analysis reveals a rather strong attractive, but non covalent interaction between the Fe atoms and the coordinated N atoms at rather long Fe–N coordination distances of  $2.451$ – $2.501\text{ \AA}$  ( $n = 1$ – $3$ ). At such distances the  $\pi$  back donation is doomed to weakness due to scant  $\pi$  orbital overlap. Instead, it stands to reason that  $\sigma$ -donation of the  $\text{N}_2$  molecule constitutes the driving force of attraction in the Fe– $\text{N}_2$  coordination in  $[\text{Fe}_3\text{O}(\text{OAc})_6(\text{N}_2)_n]^+$ . Besides the obvious attractive Fe–N interaction, we identify repulsive interactions between the carboxylate O atoms and the coordinated N atoms.

Simple considerations as *e.g.* by the DCD and BSC models do not explain the observed blue shift of the  $\text{N}_2$  stretching vibrations upon coordination. To rationalize this effect in a fairly perspicuous way we refer to the molecular orbital (MO) diagram of  $\text{N}_2$  (*cf.* Scheme S1, ESI†): all bonding MOs are populated. When ruling out short range  $\pi$  back donation into empty anti-bonding MOs (inhibited by the net positive charge of the coordinating complex), there is a way to increase the formal N–N bond order (and thus strengthen the N–N bond and increase the stretching frequency): this is a depopulation of the antibonding  $4\sigma^*$  orbital into appropriate Fe centered acceptor orbitals. We have elucidated the prevailing coordination and bonding by inspection of computed molecular orbitals in  $[\text{Fe}_3\text{O}(\text{OAc})_6(\text{N}_2)_n]^+$ . Indeed, we found that the  $4\sigma^*$  MO of the  $\text{N}_2$  unit overlaps efficiently with MOs located at the Fe centers and at the carboxylate oxygen atoms of the  $[\text{Fe}_3\text{O}(\text{OAc})_6]^+$  subunit (*cf.* Fig. 3 for  $n = 1$  and

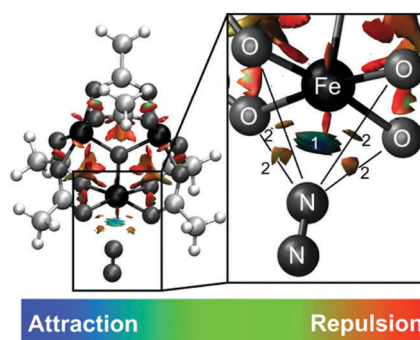


Fig. 2 NCI plot of geometry optimized  $[\text{Fe}_3\text{O}(\text{OAc})_6(\text{N}_2)_1]^+$  (*cf.* Fig. S10 for  $n = 2, 3$ , ESI†). The NCI plot reveals attractive interaction between the Fe center and the coordinated N atom (1) and repulsive interaction between of the O atoms and the N atom (2).



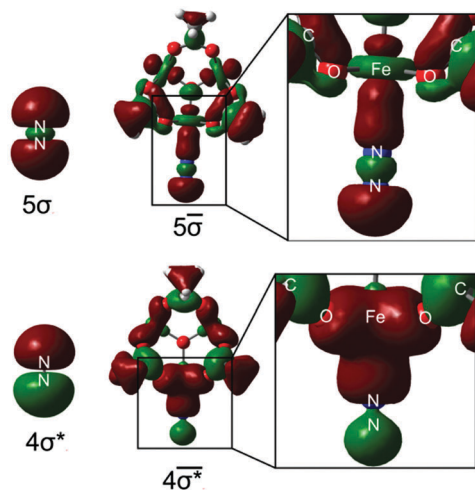


Fig. 3 Left:  $4\sigma^*/5\sigma$  molecular orbitals of free  $N_2$ . Right: Selected molecular orbitals of  $[Fe_3O(OAc)_6(N_2)_n]^+$  involving the former  $4\sigma^*(N_2)$  and  $5\sigma(N_2)$  orbitals.  $4\sigma^*(N_2)$  electron density delocalizes into the whole complex, thus strengthening the N–N bond.

Fig. S13 for  $n = 2, 3$ , ESI†). The electrons of the anti-bonding  $4\sigma^*(N_2)$  orbital delocalize into a  $4\sigma^*$  MO of the whole complex, whereby the  $4\sigma^*(N_2)$  orbital polarizes towards the Fe center. Note that the node plane of  $4\sigma^*(N_2)$  (between the N atoms) shifts somewhat towards the Fe center. The depletion of the anti-bonding electron density along the N–N bond increases the net bond order of  $N_2$  and blue shifts the  $N_2$  stretching vibration. The bonding  $5\sigma(N_2)$  contributes to a  $5\sigma$  MO of the complex. This is expected to result in N–N bond weakening counteracting the effect of  $4\sigma^*(N_2)$  delocalization. However,  $5\sigma(N_2)$  receives a partial  $4\sigma^*(N_2)$  character to form the  $5\sigma$  MO *via* hybridization. The strengthening effect of  $4\sigma^*(N_2)$  electron density delocalization thus seems to be the critical factor for the “non-classical” behavior of  $[Fe_3O(OAc)_6(N_2)_n]^+$ . The involvement of the  $4\sigma^*(N_2)$  orbital in the M– $N_2$  bonding scheme as well as  $4\sigma^*/5\sigma$  hybridization has been suggested in the context of X-ray absorption studies of  $N_2$  adsorbed on metal surfaces.<sup>32</sup> Inspection of all other delocalized MOs in the  $[Fe_3O(OAc)_6(N_2)_n]^+$  complex reveals a total lack of  $\pi$  back donation from the  $[Fe_3O(OAc)_6]^+$  unit to empty  $\pi^*(N_2)$  orbitals. Considering the high charge on the Fe center and the long Fe– $N_2$  distance (see above) this seems reasonable. The  $N_2$  coordination and thus the “non-classical” behavior of  $[Fe_3O(OAc)_6(N_2)_n]^+$  originate from  $\sigma$  bonding effects.

Our fundamental insight into the class of “non-classical”  $N_2$ –Fe complexes is remarkable in view of the industrial use of bulk iron for  $N_2$  activation and hydrogenation. It might help to advance a general understanding of dinitrogen chemistry beyond established coordination models.

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SFB/TRR 88 “Cooperative effects in homo and heterometallic complexes” (3MET).

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