



Cite this: *Chem. Commun.*, 2017, 53, 420

Received 14th September 2016,
Accepted 2nd December 2016

DOI: 10.1039/c6cc07481b

www.rsc.org/chemcomm

Vibrational blue shift of coordinated N₂ in [Fe₃O(OAc)₆(N₂)_n]⁺: “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 N₂ 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.

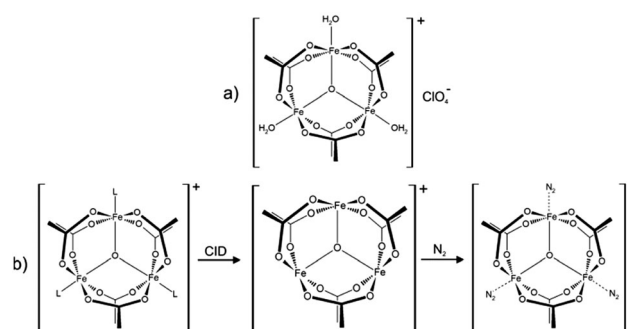
The discovery of transition metal dinitrogen complexes in 1965¹ launched the ever growing research field of N₂ coordination chemistry.^{2,3} One of the main goals is the conversion of N₂ to ammonia under mild conditions⁴ by homogenous catalysis.^{5,6} The basic idea is to weaken N–N bonds upon coordination to enable subsequent functionalization. Precursors to N₂ activation manifest by elongated N–N distances and by red shifting of stretching frequencies^{7,8} with respect to the vibration of the free N₂ molecule. Such bond activation and vibrational red shifts have been interpreted before, *e.g.* in terms of the Dewar–Chatt–Duncanson (DCD) model,⁹ and the Blyholder surface coordination (BSC)¹⁰ model. Theoretical as well as spectroscopic investigations confirmed these interpretations.¹¹ However, FT-IR studies on N₂ coordinated to Lewis acid centers in zeolites¹² and alumina¹³ revealed blue shifted vibrational N₂ stretching frequencies (up to 24 cm⁻¹). 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^{14,15} exhibit blue shifted vibrational CO stretching frequencies (up to 138 cm⁻¹).¹⁶ Gas phase studies of isolated non-classical metal carbonyl clusters^{17,18} proved the intrinsic molecular origin of this effect. The cause for the CO blue shift was a topic of debate¹⁹ and is now understood in terms of an interplay between π back donation and electrostatic effects.²⁰ 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 Spectrometry²¹ (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^{22,23} and at room temperature.²⁴ 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 [Fe₃O(OAc)₆(N₂)_n]⁺ (*n* = 1–3, OAc = CH₃CO₂⁻, *cf.* Scheme 1), which exhibit a significant blue shift (17 cm⁻¹) of N₂ vibrations in the complex with respect to the free N₂ molecule. We examine the N₂ 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 N₂ 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 [Fe₃O(OAc)₆(H₂O)₃](ClO₄) salt. In solution the water molecules exchange with solvent molecules (L = *e.g.* acetonitrile). (b) ESI-MS reveals the formation of [Fe₃O(OAc)₆(L)_n]⁺ (*n* = 0, 1, 2, 3). Collision induced dissociation (CID) of L yields the under-coordinated [Fe₃O(OAc)₆]⁺ complex, which binds 1–3 N₂ molecules at 26 K.

Fachbereich Chemie und Forschungszentrum OPTIMAS, Technische Universität Kaiserslautern, 67663 Kaiserslautern, Germany. E-mail: jlang@chemie.uni-kl.de
† Electronic supplementary information (ESI) available. See DOI: 10.1039/c6cc07481b



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 B3LYP²⁵ level of theory using cc-pVTZ basis sets²⁶ and Stuttgart RSC 1997²⁷ effective core potential basis sets (Gaussian 09²⁸). 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 N₂ stretching bands in [Fe₃O(OAc)₆(N₂)_n]⁺. It scales the calculated N₂ stretching frequencies such that a calculated free N₂ stretching vibration frequency matches the experimental value²⁹ of 2330 cm⁻¹. This approach conveniently reveals any effects of Fe–N₂ coordination on N₂ stretching frequencies. A second unspecific scaling factor (0.986) is applied for all other bands below 1800 cm⁻¹. It scales the calculated asymmetric carboxylate stretching vibration frequencies of [Fe₃O(OAc)₆(N₂)₁]⁺ to match our own experimental value of 1587 cm⁻¹. 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 [Fe₃O(OAc)₆(L)_n]⁺ (L = H₂O, acetonitrile, acetic acid; *n* = 0, 1, 2, 3). Elimination of L by Collision Induced Dissociation (CID) and subsequent coordination of N₂ in the hexapole at cryo temperatures (26 K) leads to the formation of [Fe₃O(OAc)₆(N₂)_n]⁺ (*cf.* Scheme 1b and Fig. S2, ESI†). Note that we observe *n*_{max} = 3, thus “titrating” the three available Fe

coordination sites. The calculated Gibbs energies at various temperatures reveal N₂ binding energies of 11, 9, and 8 kJ mol⁻¹ for the first, second, and third N₂ (26 K; BSSE corrected, *cf.* Fig. S3, ESI†). The N₂ binding Gibbs energies diminish with increasing temperature, vanishing above 80 K.

We recorded IR-PD spectra of cryocooled [Fe₃O(OAc)₆(N₂)_n]⁺ (*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⁻¹ and 1500 cm⁻¹, coinciding with the predicted CH₃ bending modes of the acetate ligands. While the calculated band frequencies around 1423 cm⁻¹ and 1467 cm⁻¹ 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 N₂ coordination (*n* = 1, 2, 3) has no significant influence on the frequency and intensity of the CH₃ 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–N₂ coordination sites.

We find a strong IR-PD band at 1587, 1590, and 1591 cm⁻¹ 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⁻¹ with

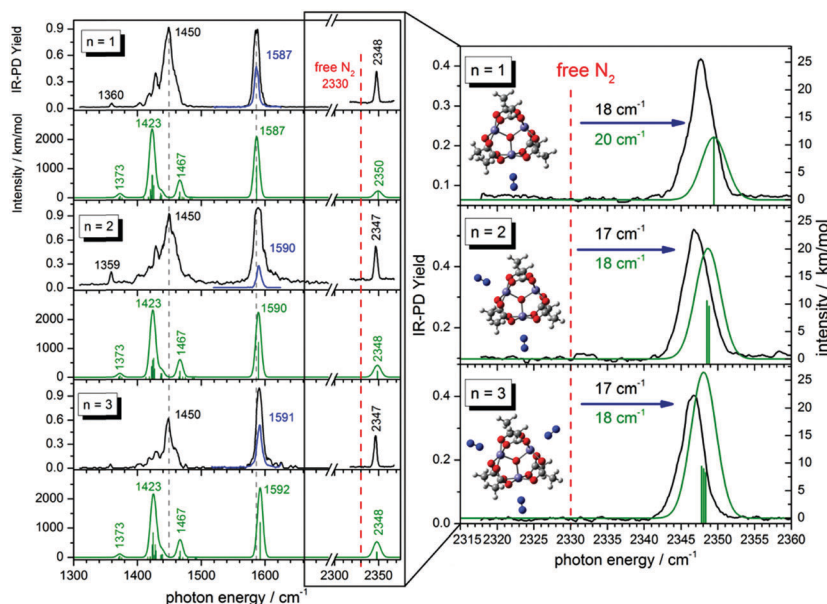


Fig. 1 Left: IR-PD spectra of [Fe₃O(OAc)₆(N₂)_n]⁺ (*n* = 1, 2, 3) at 26 K (black and blue curves) and calculated IR absorption spectra of optimized [Fe₃O(OAc)₆(N₂)_n]⁺ (*n* = 1–3) (green curves) in the range of 1300–2400 cm⁻¹. The blue IR-PD spectrum shows the CO stretching band (recorded with highly 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⁻¹ (below 2300 cm⁻¹). Calculated stick spectra were convoluted with a Gaussian envelope of FWHM = 7 cm⁻¹. Right: A zoom into the N₂ stretching vibration region. Calculated lines were convoluted with a Gaussian envelope of FWHM = 3.5 cm⁻¹. Insets show associated geometry optimized structures.

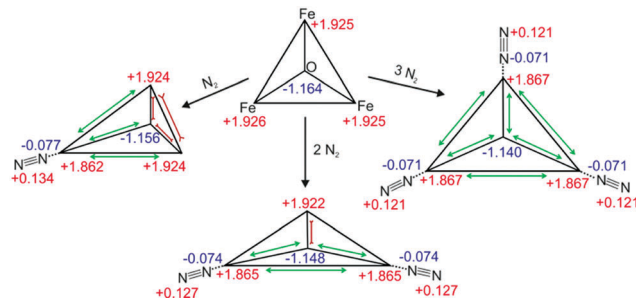


respect to the $n = 2, 3$ bands, and with respect to the corresponding band of the free acetate anion³⁰ at 1591 cm^{-1} . The DFT calculated asymmetric carboxylate stretching frequencies – scaled to match the experiment at $n = 1$ – reproduce well the reduced redshift of the experiments on $n = 2, 3$. The coordinated carboxylate groups thus sense 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 cm^{-1} revealing the IR active 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 N_2 molecule confirms our assignment of these IR-PD bands to the 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 cm^{-1} ; *cf.* Fig. S17, ESI†) and the same blue shift with respect to the free 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 ($17\text{--}18\text{ cm}^{-1}$) with respect to the (IR inactive) stretching frequency of the free $^{14}\text{N}_2$ molecule (2330 cm^{-1}).²⁹ These blue shifts indicate N–N bond strengthening upon coordination. Our DFT calculations predict this blue shift remarkably well ($18\text{--}20\text{ cm}^{-1}$). Multiple coordinated N_2 molecules have almost identical stretching frequencies. There seems to be no coupling between the N_2 molecules adsorbing at distinct, equivalent Fe sites. In contrast to the carboxylate asymmetric stretching bands the N_2 stretching bands shift slightly to lower frequencies with increasing n (IR-PD spectrum: 1 cm^{-1} ; DFT: 1.5 cm^{-1}). However, in all three cases ($n = 1\text{--}3$) the N_2 stretching band is blue shifted with respect to the free N_2 .

The DFT calculations reveal subtle distortions of the triangular Fe_3O -core upon coordination of N_2 (*cf.* Scheme 2 and Table S2, ESI†). Each N_2 molecule binds end on to the respective Fe atom. In general, the coordination of N_2 enlarges Fe–Fe distances and Fe– $\text{O}_{\text{central}}$ bond lengths (by 0.03 \AA resp. 0.01 \AA) of those bonds which involve the N_2 coordinating Fe-center. All the other Fe–Fe distances and Fe– $\text{O}_{\text{central}}$ bond lengths shorten by approx. 0.03 \AA or 0.01 \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 Fe_3O -core (*cf.* Scheme 2). The Fe centers gain $0.058\text{--}0.064\text{ e}$ electron density per coordinating N_2 . The non-coordinated N atoms donate $0.121\text{--}0.134\text{ e}$ electron density in about equal parts to the coordinating N and Fe atoms. Free, non-polar N_2 molecules thus polarize and oxidize significantly upon Fe coordination, both magnitudes decreasing with n – as does the blue shift of the N_2 stretching bands. Note that the N–N bond length contracts slightly upon coordination (-0.002 \AA). The high positive charge on the Fe centers ($1.862\text{--}1.922\text{ e}$) diminishes their electron donating capability and thus inhibits the π back donation. Such charge effects seem to increase the “non-classical” effect in metal carbonyl complexes.³¹



Scheme 2 Calculated geometries and natural charge distributions of the Fe_3O -core and the coordinated 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 $2.451\text{--}2.501\text{ \AA}$ ($n = 1\text{--}3$). At such distances the π back donation is doomed to weakness due to scant π orbital overlap. Instead, it stands to reason that σ -donation of the N_2 molecule constitutes the driving force of attraction in the Fe– 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 N_2 stretching vibrations upon coordination. To rationalize this effect in a fairly perspicuous way we refer to the molecular orbital (MO) diagram of N_2 (*cf.* Scheme S1, ESI†): all bonding MOs are populated. When ruling out short range π 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 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

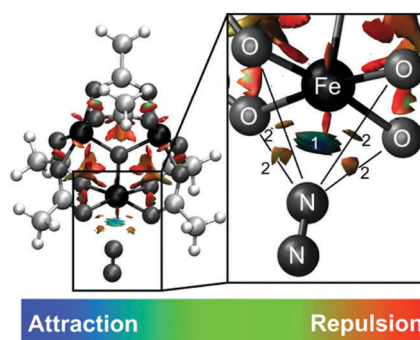


Fig. 2 NCI plot of geometry optimized $[\text{Fe}_3\text{O}(\text{OAc})_6(\text{N}_2)_1]^+$ (*cf.* Fig. S10 for $n = 2, 3$, ESI†). The NCI plot reveals attractive interaction between the Fe center and the coordinated N atom (1) and repulsive interaction between of the O atoms and the N atom (2).



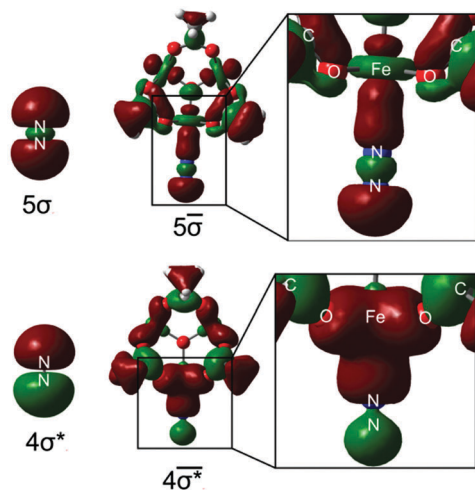


Fig. 3 Left: $4\sigma^*/5\sigma$ molecular orbitals of free N_2 . Right: Selected molecular orbitals of $[Fe_3O(OAc)_6(N_2)_n]^+$ involving the former $4\sigma^*(N_2)$ and $5\sigma(N_2)$ orbitals. $4\sigma^*(N_2)$ electron density delocalizes into the whole complex, thus strengthening the N–N bond.

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 $5\bar{\sigma}$ 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 $5\bar{\sigma}$ 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.³² Inspection of all other delocalized MOs in the $[Fe_3O(OAc)_6(N_2)_n]^+$ complex reveals a total lack of π 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 σ 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.

This work was supported by the German Research Foundation DFG within the Transregional Collaborative Research Center

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

Notes and references

- M. D. Fryzuk, *Chem. Commun.*, 2013, **49**, 4866–4868.
- J. L. Crossland and D. R. Tyler, *Coord. Chem. Rev.*, 2010, **254**, 1883–1894.
- N. Khoenkhoen, B. de Bruin, J. N. H. Reek and W. I. Dzik, *Eur. J. Inorg. Chem.*, 2015, 567–598.
- S. F. McWilliams and P. L. Holland, *Acc. Chem. Res.*, 2015, **48**, 2059–2065.
- D. V. Yandulov and R. R. Schrock, *Science*, 2003, **301**, 76–78.
- H. Tanaka, K. Arashiba, S. Kuriyama, A. Sasada, K. Nakajima, K. Yoshizawa and Y. Nishibayashi, *Nat. Commun.*, 2014, **5**, 1–11.
- N. Lehnert and F. Tuczek, *Inorg. Chem.*, 1999, **38**, 1659–1670.
- N. Lehnert and F. Tuczek, *Inorg. Chem.*, 1999, **38**, 1671–1682.
- J. Chatt and L. A. Duncanson, *J. Chem. Soc.*, 1953, 2939–2947.
- G. Blyholder, *J. Phys. Chem.*, 1964, **68**, 2772–2777.
- F. Stuy and F. Tuczek, *J. Comput. Chem.*, 2006, **27**, 1278–1291.
- K. Hadjiivanov and H. Knözinger, *Catal. Lett.*, 1999, **58**, 21–26.
- R. Wischert, C. Coperet, F. Delbecq and P. Sautet, *Chem. Commun.*, 2011, **47**, 4890–4892.
- H. Willner and F. Aurbke, *Angew. Chem., Int. Ed.*, 1997, **36**, 2402–2425.
- A. J. Lupinetti, G. Frenking and S. H. Strauss, *Angew. Chem., Int. Ed.*, 1998, **37**, 2113–2116.
- P. K. Hurlburt, J. J. Rack, J. S. Luck, S. F. Dec, J. D. Webb, O. P. Anderson and S. H. Strauss, *J. Am. Chem. Soc.*, 1994, **116**, 10003–10014.
- J. Velasquez, B. Njagic, M. S. Gordon and M. A. Duncan, *J. Phys. Chem. A*, 2008, **112**, 1907–1913.
- A. Fielicke, G. von Helden, G. Meijer, B. Simard and D. M. Rayner, *J. Phys. Chem. B*, 2005, **109**, 23935–23940.
- A. S. Goldman and K. Krogh-Jespersen, *J. Am. Chem. Soc.*, 1996, **118**, 12159–12166.
- G. Bistoni, S. Rampino, N. Scafuri, G. Ciancaleoni, D. Zuccaccia, L. Belpassi and F. Tarantelli, *Chem. Sci.*, 2016, **7**, 1174–1184.
- J. B. Fenn, *Angew. Chem., Int. Ed.*, 2003, **42**, 3871–3894.
- N. Heine and K. R. Asmis, *Int. Rev. Phys. Chem.*, 2014, **34**, 1–34.
- J. Jašík, J. Žabka, J. Roithová and D. Gerlich, *Int. J. Mass Spectrom.*, 2013, **354–355**, 204–210.
- J. Lang, M. Gaffga, F. Menges and G. Niedner-Schatteburg, *Phys. Chem. Chem. Phys.*, 2014, **16**, 17417–17421.
- A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- T. H. Dunning, *J. Chem. Phys.*, 1989, **90**, 1007–1023.
- M. Dolg, H. Stoll, H. Preuss and R. M. Pitzer, *J. Phys. Chem.*, 1993, **97**, 5852–5859.
- M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. A. Petersson, H. Nakatsuji, M. Caricato, X. Li, H. P. Hratchian, A. F. Izmaylov, J. Bloino, G. Zheng, J. L. Sonnenberg, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, J. A. Montgomery, Jr., J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, N. Rega, J. M. Millam, M. Klene, J. E. Knox, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, R. L. Martin, K. Morokuma, V. G. Zakrzewski, G. A. Voth, P. Salvador, J. J. Dannenberg, S. Dapprich, A. D. Daniels, Ö. Farkas, J. B. Foresman, J. V. Ortiz, J. Cioslowski and D. J. Fox, *Gaussian 09, Revision E.01*, Gaussian, Inc., Wallingford CT, 2009.
- K. P. Huber and G. Herzberg, *Constants of Diatomic Molecules*, Van Nostrand, New York, 1979.
- J. D. Steill and J. Oomens, *J. Phys. Chem. A*, 2009, **113**, 4941–4946.
- A. M. Ricks, J. M. Bakker, G. E. Douberly and M. A. Duncan, *J. Phys. Chem. A*, 2009, **113**, 4701–4708.
- A. Nilsson and L. G. M. Pettersson, *Surf. Sci. Rep.*, 2004, **55**, 49–167.

