



Cite this: *Phys. Chem. Chem. Phys.*, 2022, 24, 11616

Optical absorption and shape transition in neutral Sn_N clusters with $N \leq 40$: a photodissociation spectroscopy and electric beam deflection study†

Andreas Lehr, * Filip Rivic, Marc Jäger, Martin Gleditsch and Rolf Schäfer

Neutral Sn_N clusters with $N = 6$ –20, 25, 30, 40 are investigated in a joint experimental and quantum chemical study with the aim to reveal their optical absorption in conjunction with their structural evolution. Electric beam deflection and photodissociation spectroscopy are applied as molecular beam techniques at nozzle temperatures of 16 K, 32 K and 300 K. The dielectric response is probed following the approach in S. Schäfer *et al.*, *J. Phys. Chem A*, 2008, **112**, 12312–12319. It is improved on those findings and the cluster size range is extended in order to cover the prolate growth regime. The impact of the electric dipole moment, rotational temperature and vibrational excitation on the deflection profiles is discussed thoroughly. Photodissociation spectra of tin clusters are recorded for the first time, show similarities to spectra of silicon clusters and are demonstrated to be significantly complicated by the presence of multi-photon absorption in the low-energy region and large excess energies upon dissociation which is modelled by the RRKM theory. In both experiments two isomers for the clusters with $N = 8, 11, 12, 19$ need to be considered to explain the experimental results. Triple-capped trigonal prisms and double-capped square antiprisms are confirmed to be the driving building units for almost the entire size range. Three dominating fragmentation channels are observed, *i.e.* the loss of a tin atom for $N < 12$, a Sn_7 fragment for $N < 19$ and a Sn_{10} fragment for $N \geq 19$ with Sn_{15} subunits constituting recurring geometric motifs for $N > 20$. The prolate-to-quasispherical structural transition is found to occur at $30 < N \leq 40$ and is analyzed with respect to the observed optical behavior taking quantum chemical calculations and the Mie–Gans theory into account. Limitations of the experimental approach to study the geometric and electronic structure of the clusters at elevated temperatures due to vibrational excitation is also thoroughly discussed.

Received 10th March 2022,
 Accepted 25th April 2022

DOI: 10.1039/d2cp01171a

rsc.li/pccp

1 Introduction

Group 14 of the periodic table, comprising C, Si, Ge, Sn, Pb (and Fl), is arguably the most significant group of elements in all chemistry. The metals lead and tin are widely known for their use in the production of structural components and alloys, the semiconductors silicon and germanium constitute

the foundation of modern technology and the nonmetal carbon, evidently, states the basis of organic life.¹ It is in particular this nonmetal-to-semiconductor-to-metal transition along the group of these tetrel elements (referring to the most common valency of four) that intrigues engineers and researchers alike. With tin being on the very edge of this transition, it exhibits, as a consequence, two stable allotropes, the gray diamond cubic α -Sn and the white body-centered tetragonal β -Sn, resembling the zero-gap semiconductor² (<286 K) and metal nature of this material, respectively. Albeit standing in the shadows of silicon (and also germanium) dominating the semiconductor industry, tin offers various desirable properties such as being non-toxic, inexpensive and a widespread element. Driven by the endeavor to further miniaturize electronic devices, silicon nanoparticles have already found their way into the microelectronic industry.^{3–6} Also tin nanoparticles are dealt in energy storage applications as promising anode material in next-generation batteries due to their high capacity.^{7–10} Since small clusters are well known for their physical properties being tremendously

Technical University of Darmstadt, Eduard-Zintl-Institut, Alarich-Weiss-Straße 8, 64287 Darmstadt, Germany. E-mail: andreas.lehr@tu-darmstadt.de

† Electronic supplementary information (ESI) available: Geometries, symmetries and energetics of all Sn_N^+ isomers; (relative) ellipsoid semiaxes of all Sn_N isomers; vibrational frequencies of all Sn_N isomers; dissociation and RRKM excess energies of all Sn_N isomers; Boltzmann correction for the fitted isomer mixtures in the electric beam deflection and photodissociation spectroscopy experiments of Sn_8 , Sn_{11} , Sn_{12} and Sn_{19} ; impact of the rotational temperature and computed electric dipole moment on the electric beam deflection profile demonstrated for Sn_{10} ; computed moments of inertia, electric dipole moment components and polarizabilities as well as experimentally extracted electric dipole moments and polarizabilities of all Sn_N isomers; TDDFT fundamental and optical band gaps for all Sn_N clusters. See DOI: <https://doi.org/10.1039/d2cp01171a>



size, composition and charge state-dependent, thus being tunable, and with tin clusters showing similar structural growth patterns and optoelectronic properties as silicon clusters – as to be revealed by this study – a vast range of potential applications is expected to open up.

After the first mass spectrometric detection of small tin clusters by simple vaporization in 1952,¹¹ it took several years until their stabilities have been probed by means of determining atomization enthalpies.^{12–14} The time-resolved laser-induced fluorescence spectroscopy of the Sn₂ dimer in 1982 provided accurate insights into the geometric and electronic structure^{15,16} and, besides early extended Hückel studies,^{17,18} opened up a playground for quantum chemists to study the low-lying excited states, equilibrium geometric constants and potential energy surfaces for small clusters up to Sn₆ employing extensive multi-reference configuration interaction methods.^{19–27} While on the theoretical side density functional theory (DFT) has become today's routine for investigating the geometries and electronic structure of larger clusters,^{28–51} there have only been a few experimental studies validating the manifold of proposed geometries. Mass spectrometric investigations alone^{52,53} or involving photoionization and photofragmentation^{54–59} are usually not capable of obtaining structural information. Cationic tin cluster geometries have been successfully examined by ion mobility^{42,60–62} and low-energy surface collision measurements^{63,64} revealing a prolate growth pattern, similar to observations made for silicon and germanium clusters⁶¹ before quasispherical structures become favorable again for larger clusters. To study anionic clusters photoelectron spectroscopy has been proven one of the most powerful tools,^{65–71} but also ion mobility, trapped ion diffraction and collision induced dissociation experiments have been conducted.^{39,41,45} When talking about charged tin clusters, it is worth mentioning that for cationic species the abnormally high melting temperatures above the bulk value have been discovered for the first time.^{36,72–77} Also, the icosahedral stannaspherene cluster Sn₁₂^{2–} which was found by photoelectron spectroscopy^{78–81} has acquired huge interest due to being the scaffold for paramagnetic superatoms^{82–84} as well as its relation to Zintl ions and (inter-)metalloid compounds.^{85–98} Neutral clusters, however, are more difficult to probe experimentally. There are only two kinds of studies addressing the geometry discrimination of neutral tin clusters, one which measures the formation energies of tin cluster distributions consisting of hundreds of atoms calorimetrically^{99,100} and one which focuses on the structure elucidation of Sn_N clusters with N = 6–20 based on measuring the Stark effect in combination with photoionization and quantum chemical calculations.³⁷ The results presented here take up the findings of the latter study on the geometric evolution of small tin clusters, extend the size range and place special focus on the interplay between geometric structure and the optical absorption. Apart from a standalone study of a Sn₁₄ cluster,⁵¹ the optoelectronic characteristics lack entirely both experimental and theoretical investigations making this the first such study employing a combined photodissociation spectroscopy and electric beam deflection approach supported by demanding quantum chemical calculations.

2 Experimental methods

An extensive presentation of the methodology and experimental details are described for both experimental approaches elsewhere.^{101–105} Therefore, only a brief overview is given here. In general, neutral and charged Sn clusters are generated by a pulsed (10 Hz) and He buffer gas perfused laser vaporization source which can be cooled down *via* a cryogenic nozzle ($T_{\text{nozzle}} \geq 16$ K) prior to the expansion into high vacuum and the subsequent formation of the molecular beam.

2.1 Electric beam deflection

In this experiment the molecular beam is formed by expansion into the high vacuum through a double-skimmer. Afterwards, it passes a mechanical shutter based on a hard drive disk^{106,107} which allows one to determine the velocity v_x of the clusters in flight direction x .¹⁰⁸ Two collimating slits shape the molecular beam to the rectangular profile of 0.4 mm width and 2.0 mm height. Here, the second collimating slit is located directly in front of the electric deflection unit which is based on the two-wire geometry to maintain a nearly constant gradient in the direction of deflection z .¹⁰⁹ The deflection is probed by a scanning slit aperture such that the intensities of the mass signals are measured in dependence of the position along the inhomogeneous field. The clusters are ionized by an F₂ excimer laser (7.89 eV) directly behind the scanning slit and detected in a time-of-flight mass spectrometer (TOF-MS). To avoid multi-photoionization and photofragmentation a plane-concave lense widens the profile of the F₂ excimer laser along the direction of flight. To be able to interpret the beam deflection profiles, mass spectra with and without applied field are recorded alternately. Regarding the intensities of the beam deflection profiles shown in this work, these mass spectra pairs are recorded at every position in random order. In order to compare the beam deflection profiles from several experimental runs, the measured deflection d_z has to be displayed independently of the cluster velocity v_x which varies slightly in every run. Therefore, the measured deflection is converted into the mean projection of the dipole moment on the field direction $\bar{\mu}_{\text{el},z}$ of the Sn_N clusters. In eqn (1) m_{Sn} is the mass of the tin atom, dE_z/dz the electric field gradient and l_1 and l_2 the lengths of the electric field and field free region, respectively.

$$\bar{\mu}_{\text{el},z} = \frac{Nm_{\text{Sn}}v_x^2d_z}{\frac{1}{2}l_1^2 + l_1l_2} \left(\frac{dE_z}{dz} \right)^{-1} \quad (1)$$

2.2 Photodissociation spectroscopy

After the expansion into high vacuum through a skimmer and thereby forming the molecular beam, an electric deflector is used to remove charged particles. The optical response is probed by longitudinal photodissociation spectroscopy employing an optical parametric oscillator combined with several harmonic generators, which covers the photon energy range of $\hbar\omega = 1.9$ –5.4 eV. After the interaction with the dissociation laser, the neutral species are photoionized either by a Xe flash lamp (<10.3 eV) or an F₂ excimer laser (7.89 eV) and are detected by a TOF-MS.



The photon fluence is kept low to avoid multiphoton dissociation processes while recording photoionization mass spectra. Optical absorption spectra are measured by monitoring the mass signal depletion I/I_0 as a function of the photon energy $\hbar\omega$ and applying the Beer–Lambert law to calculate the dissociation cross section $\sigma(\hbar\omega)$ assuming an one-photon absorption process. In eqn (2) a perfect overlap between the dissociation laser and the molecular beam has to be guaranteed and the photon fluence ϕ needs to be determined in a separate experiment.

$$\sigma(\hbar\omega) = -\frac{1}{\phi} \ln \frac{I}{I_0} \quad (2)$$

All dissociation cross sections shown here have to be regarded as lower limits to the true absorption cross sections. This is because competing non-dissociative relaxation pathways, such as fluorescence, cannot be detected. In particular, for excitation energies smaller than or comparable to the threshold energy for dissociation, no dissociation is detectable during the experimental time scale in the range of a few 100 μs ,^{110,111} and thus no light absorption can then be observed. Therefore, the dissociation behavior must be taken into account when discussing the determined absorption cross sections, which is further deepened in Section 4.3.

3 Computational methods

3.1 Genetic algorithm

Suitable structural candidates of the Sn_N clusters with $N = 6\text{--}20$, 25, 30, 40 are identified by means of an unbiased global optimization of the configurational space. This is accomplished with the help of the German Improved Genetic Algorithm (GIGA)¹¹² at a plane-wave DFT level of theory using Quantum Espresso v6.4.1.^{113,114} In this process, the PBE exchange–correlation (xc) functional^{115,116} and the ultrasoft RRKJ pseudopotential^{117,118} (36 core electrons for tin) with a plane-wave kinetic energy cutoff of 400 eV were employed. The box size of the unit cell is determined dynamically such that no duplicate clusters are closer than 16 Å and self-interaction can be neglected. Spin polarization was implemented and a small Methfessel–Paxton smearing¹¹⁹ as well as non-linear core corrections¹²⁰ were applied. The pool started with 15 up to 25 randomly generated structures for Sn_6 up to Sn_{20} . In order to render the search for symmetric structures more efficient, clusters containing triple-capped trigonal prism and double-capped square antiprism units have been used in the starting pool for Sn_{25} , Sn_{30} and Sn_{40} . The search routine was stopped when the most recent 1000 structures being generated did not appear below 1 eV relative to the pool global minimum (GM). For Sn_{40} , *i.e.*, a total of 5000 geometries were created. The results of the global optimization are analyzed with the help of our GA analysis software.¹¹²

3.2 Geometry optimization and frequency analyses

All lowest-lying isomers are locally reoptimized using Gaussian16¹²¹ and NWChem v6.8.1.^{122,123} Orbital-based DFT with tight optimization criteria and a high-density numerical grid is applied employing the PBE0/cc-pVTZ-PP^{124–126} and LC- ω PBEh/def2-TZVPP^{125,127,128} xc functional/basis set combinations

taking into account singlet and triplet spin configurations. Scalar relativistic corrections are effectively treated by the use of such pseudopotential-supported basis sets. The choice of the xc functional/basis set combination is justified by an extensive benchmark with experimental data of Sn_6 and Sn_7 ⁵¹ as well as previous studies on doped tin clusters.^{129–132} CCSD(T)/cc-pVTZ-PP single-point energies on top of the PBE0/cc-pVTZ-PP optimized geometries have been computed using the domain-based local pair natural orbital (DLPNO) acceleration approach^{133–136} implemented in Orca v5.0.1.^{137–139} Furthermore, the CCSD(T) energies were computed using the TightPNO criteria and extrapolated to the complete basis set limit using the two-point extrapolation for the cc-pVTZ-PP and corresponding auxiliary basis sets.¹⁴⁰ Only those clusters were considered for the analysis of the experiment with a relative energy of $\Delta E \leq 0.20$ eV with respect to the predicted GM on either of the three levels of theory. Some final structural isomers of larger clusters have only been found from reoptimizing corresponding cationic GA candidates. An overview of all reoptimized cationic Sn_N^+ structures can be found in the ESI.† Harmonic frequency analyses are performed for all species in order to verify that the optimized geometries correspond to local minima on the potential energy surface. A summary of the vibrational frequencies of all structural isomers can be found in the ESI.†

3.3 Dielectric and optical response

The moments of inertia and the dielectric properties comprising the electric dipole moment components and the isotropic electronic polarizability were taken from PBE0/cc-pVTZ-PP results due to their already demonstrated good accuracy–cost balance for tin-based clusters.^{51,129–132} With these properties, electric deflection profiles can be predicted by simulating the rotational dynamics of an ensemble of clusters in the electric field. Here, a classical approach is considered, since even for low rotational temperatures rotational states of high angular momentum quantum number J are still significantly populated.¹⁰² To give an example, already a rotational temperature of $T_{\text{rot}} = 1$ K causes the rotational states of the smallest cluster under investigation, Sn_6 , to be significantly populated up to $J = 30$ considering a rotational constant of $B = 0.44$ m^{-1} . A canonical ensemble of 5000 clusters is set up with an individually defined rotational state and Boltzmann-distributed with a defined rotational temperature. These clusters enter the electric field adiabatically such that the given rotational state does not change when entering the inhomogeneous field. Within the field, the Euler equations for a classical rotor are solved in a trajectory simulation for every time step.^{102,141} The resulting distribution of the electric dipole moments of the ensemble is then convoluted with the Gaussian function describing the experimental beam profile without applied electric field. This procedure yields the simulated electric deflection profiles which can be compared to the experiment. Note that this approach is only valid for rigid rotors. The shift of the simulated deflection profile is connected to the computed electronic polarizability of the cluster.^{37,102} Since the rotational temperature can only be estimated for a given nozzle temperature and the electric dipole moments resulting from



quantum chemical calculations come with an uncertainty of up to 10–15%,¹⁴² it is necessary to investigate their influence on the simulated electric beam profiles of the tin clusters. For this, the electric deflection profile together with the classical trajectory simulations are shown exemplary for Sn₁₀ with varied values of the electric dipole moments and rotational temperatures in the ESI.† This cluster is well suited for such a study because there is only one structural isomer present, which possesses only one non-zero component of the electric dipole moment. The results indicate that both reducing the electric dipole moment component and increasing the rotational temperature also yield a simulated deflection profile which describes the experimental data to a good approximation, so that one has to keep in mind that there is a certain freedom in choosing these two parameters. Because the experimental data for larger clusters with $N > 18$ could only be obtained with a higher nozzle temperature of 30 K, setting the rotational temperature to $T_{\text{rot}} = 20$ K is considered appropriate.^{108,132} Furthermore, the calculation of an effective polarizability by using second-order perturbation theory (*cf.* ESI† for details) demonstrates that a rotational temperature of 20 K is the most reasonable choice for most cluster species. With these two things in mind, all trajectory simulations in this work have been carried out with the computed electric dipole moments and a rotational temperature of $T_{\text{rot}} = 20$ K for reasons of consistency.

Singlet excited states and corresponding transition dipole moments are calculated by means of time-dependent DFT (TDDFT) at the LC- ω PBEh/def2-TZVPP level of theory employing Gaussian16 and NWChem v6.8.1 to simulate the experimentally measured optical absorption spectra. Up to 1500 excited states for Sn₄₀ had to be considered in order to cover the experimental accessible photon energy range. A natural transition orbital (NTO) analysis has been carried out aiming for a more intuitive picture of which (mixed) orbitals are involved in particle-hole excitations.¹⁴³ The obtained oscillator strengths f are connected to the absorption cross sections σ by integration over the corresponding spectral bands given a certain full width at half maximum (FWHM), which is found to be 0.27 eV for photodissociation spectra recorded at 300 K and 0.20 eV at 32 K.

$$f = \frac{4\epsilon_0 m_e c}{he^2} \int_{\text{band}} d(\hbar\omega) \sigma(\hbar\omega) \quad (3)$$

0.911 Å⁻² eV⁻¹

This approach has been validated in a benchmark study of Sn₆ and Sn₇ as well as the already covered Sn₁₄ cluster.⁵¹ Here, the Sn₆ and Sn₇ clusters are shown again in order to exemplary draw a picture of the multiphoton absorption which is observed in many of the photodissociation spectra.

4 Preliminary considerations

4.1 Structural classification

In order to tackle the prolate shape transition and to conceptualize the cluster's prolateness, the cluster shapes are viewed

as ellipsoids with their three semiaxes A_i ($A_1 \geq A_2 \geq A_3$).

$$A_i = \sqrt{\frac{1}{Nm_{\text{Sn}}^2} \left(\sum_{i \neq j=1}^3 I_i - I_j \right)} \quad (4)$$

Herein, I_i gives the computed axis-dependent moments of inertia of the Sn _{N} clusters. A dimensionless relative quantity A_i^{rel} is chosen such that a perfect sphere gives a value of one and the prolateness can be measured by the deviation of the largest component A_1^{rel} from this value.^{144,145}

$$A_i^{\text{rel}} = \frac{3A_i}{\sum_{j=1}^3 A_j} \quad (5)$$

Fig. 1 depicts these semiaxes for all structural isomers; a corresponding table can be found in the ESI.† Clearly, the prolate growth pattern ($A_1^{\text{rel}} > A_2^{\text{rel}} \approx A_3^{\text{rel}}$) for clusters with more than approximately twelve atoms becomes apparent. This behavior continues for the whole size range studied, although quasispherical structures appear (here as an oblate spheroid with $A_1^{\text{rel}} \approx A_2^{\text{rel}} > A_3^{\text{rel}}$) and become competing candidates starting around Sn₄₀.

4.2 Fragmentation mechanisms

Before discussing the dielectric and optical behavior, it is worth looking first at the fragmentation mechanisms obtained for the investigated clusters. Since it is far less trivial to study this experimentally for neutral clusters compared to the charged species,^{42,63,64,111} the photodissociation experiment does not yield cluster-specific information on the photoinduced fragments. However, when varying the delay time of the spectroscopy laser, assuring that fragments will be ionized and detected by the

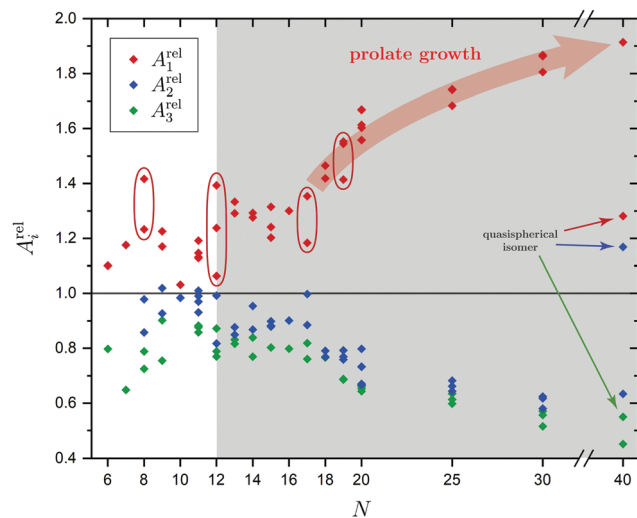


Fig. 1 Relative ellipsoid semiaxes A_i^{rel} based on the PBE0/cc-pVTZ-PP calculated moments of inertia shown for every optimized Sn _{N} structural isomer. A significant deviation from $A_1^{\text{rel}} = 1$ indicates a prolate geometry approximately starting from $N \geq 12$ as highlighted by the gray box. Those species with isomers showing a rather large difference in their prolateness are stressed by red ellipses.



TOF-MS, a considerable gain in the Sn₇ and Sn₁₀ mass signals is observed. This can be viewed as evidence for those fragmentation channels to be the most dominant ones. Moreover, dissociation energies for the energetically-lowest fragmentation paths have been calculated and compared with experimental values for the smallest clusters,¹⁴ as shown in Fig. 2 and further deepened in the ESI.† Although the correspondence of the calculated dissociation energies with the experimental ones is not quantitative, the trend with cluster size is covered reasonably well. Due to the particularly stable Sn₇ and Sn₁₀ units, three significant fragmentation channels can be observed. Sn_N clusters with N < 12 favor the loss of a tin atom. Clusters with N < 19 dissociate into a Sn₇ unit whereas clusters with N ≥ 19 lead into a Sn₁₀ fragment and another daughter cluster. It should be noted that the Sn₁₅ unit is also comparatively stable which is in agreement with recent theoretical findings,^{44,47} but does not replace the Sn₁₀ fragmentation channel for this size range. Interestingly, the computed dissociation energies resemble quite well the anti-magic abundance pattern of the clusters Sn₁₃, Sn₁₄, Sn₁₇ and Sn₂₀, which fragment more easily given the energy difference between the F₂ excimer laser energy of 7.89 eV and their ionization potentials all being in the

range 6.17–6.43 eV.⁵⁹ The energetics of these dissociation mechanisms have a far-reaching impact on the evaluation of our photodissociation spectra when interpreted in terms of optical absorption, which is discussed in detail for Sn₆, Sn₇ and Sn₁₀ in Sections 5.1 and 5.2 taking also multiphoton processes into account.

4.3 Dissociation kinetics

To comprehend the fragmentation of the clusters on the experimental time scale, it is favorable to investigate the kinetics of the photodissociation process. Many small metal clusters have been found to fragment *via* statistical unimolecular dissociation¹¹⁰ such that the rate constant for dissociation *k* can be evaluated in terms of the Rice–Ramsperger–Kassel–Marcus (RRKM) theory.^{146–149}

$$k(E + E_{\text{int}}) = \frac{s n^{\ddagger} (E + E_{\text{int}}^{\ddagger} - E_{\text{D}})}{h \rho(E + E_{\text{int}})} \quad (6)$$

Here, *E* is the photon energy, *E*_{int} and *E*_{int}[‡] the internal energy comprising the “active” vibrational and rotational degrees of freedom at the initial and transition state,¹⁵⁰ *E*_D the dissociation energy, *s* the reaction path degeneracy, *n*[‡] the sum of ro-vibrational states at the transition state and *ρ* the density of ro-vibrational states. The latter two quantities are computed employing the Beyer–Swinehart direct count algorithm.¹⁵¹ The reaction path degeneracy *s* varies depending on the dissociation process as well as the cluster geometry and is chosen individually for all structural isomers. To give some examples, *s* is set to the number of atoms for an atom-loss process of the Sn₆ and Sn₇ parent cluster, whereas it is lowered for larger clusters whenever there are peripheral tin atoms leaving behind a stable daughter subunit. The same holds for other fragmentation channels of larger clusters where a Sn₇ or Sn₁₀ cluster fragment is discernible in the parent cluster structure. The clusters are well known to have highly excited angular momentum already at very low rotational temperatures, which is why the angular momentum dependence in eqn (6) has been considered in the first place. However, in estimating approximate lifetimes of the clusters under experimental conditions, the angular momentum dependence is shown to cause only insignificant changes and could be reasonably dropped. Furthermore, it has been found to be of minor importance which vibrational mode is responsible for dissociation, especially as the cluster size increases. Therefore, anharmonic corrections to the vibrational frequencies have been neglected, although they can become important in certain situations due to the high degree of vibrational excitation common in unimolecular dissociation processes. The assumption is made that no activation barrier is related to the fragmentation process, thus drawing a direct relation between the transition state and dissociation energy for reactions of the type Sn_N → Sn_{N–M} + Sn_M. It should be noted that care has to be taken, since the activation barriers in general are not obtainable in a straightforward way and thus not regarded here. By calculating the rate of dissociation, lifetimes *τ* of the clusters can be estimated as a function of their excess energy *E* – *E*_D as shown in Fig. 3 and listed fully in the ESI,† providing a useful insight into the dissociation behavior under the given experimental conditions. Due to the

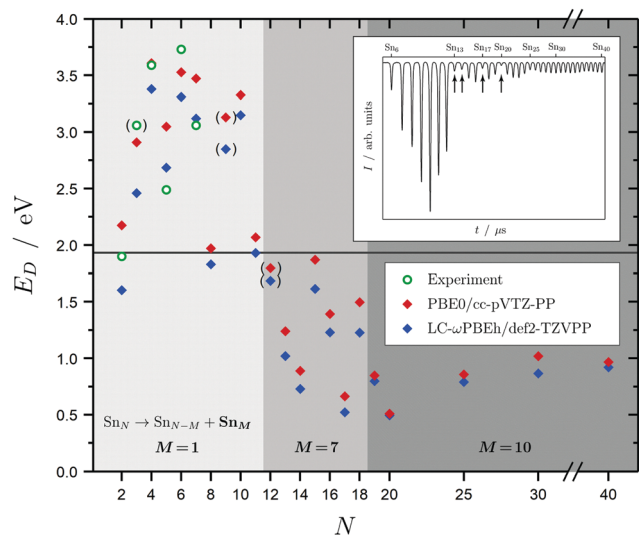


Fig. 2 Dissociation energies *E*_D in eV for the energetically most favorable fragmentation channels of the Sn_N clusters with N = 2–20, 25, 30, 40 averaged over all isomers in the given energy range. Experimental values for the incremental dissociation energies are estimated based on atomization enthalpies obtained from Knudsen cell mass spectrometry.¹⁴ The horizontal line at 1.9 eV gives the smallest accessible single-photon energy available in our photodissociation experiment. Over the size range studied, three regions can be distinguished: dissociation under the loss of an atom (light gray), a Sn₇ (gray) and a Sn₁₀ fragment (dark gray). Calculated dissociation energies in parentheses point to a differing fragmentation mechanism, *i.e.*, loss of a Sn₂ fragment for the Sn₉ parent cluster and loss of Sn₆ fragments for the Sn₁₂ parent cluster. The Sn₃ experimental dissociation energy in parentheses has been evaluated assuming a linear geometry¹² (in its lowest ¹A₁ and ³B₂ state Sn₃ is an isoscele triangle²¹) and has thus to be considered with care. The inset shows a typical time-of-flight mass spectrum after F₂ excimer laser photoionization (7.89 eV) of the Sn_N clusters given as arbitrary intensity *I* in dependence of the flight time *t*; the naturally low-abundant clusters Sn₁₃, Sn₁₄, Sn₁₇ and Sn₂₀ are highlighted by arrows.



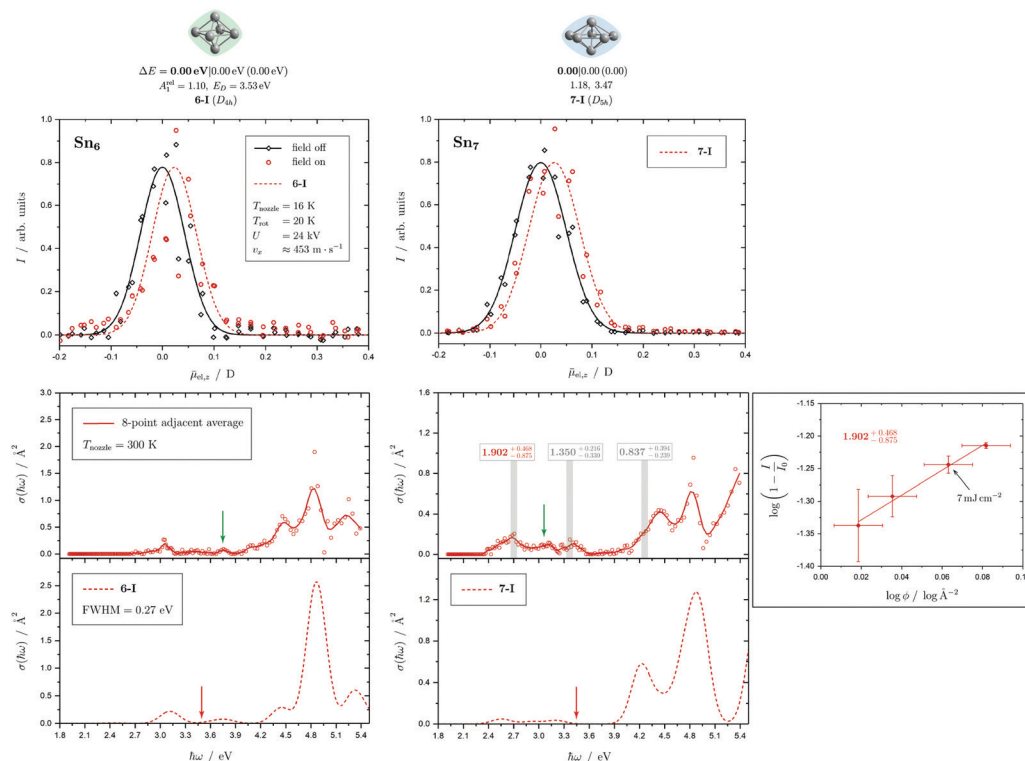


Fig. 4 Summary of all structural isomers (top), electric beam deflection measurements together with the corresponding classical trajectory simulations (center) and all photodissociation spectra together with the TDDFT simulations (bottom), here shown for the clusters Sn₆ and Sn₇. This caption serves as basis also for the following figures. Top: The geometries are shown together with their molecular point groups. The first line gives the energies ΔE relative to the GM in eV at the PBE0/cc-pVTZ-PP[CCSD(T)/cc-pVTZ-PP(LC- ω PBEh/def2-TZVP)] levels of theory. The second line gives the largest dimensionless semiaxis A_1^{el} for an ellipsoid as a measure of the cluster's prolateness, followed by the dissociation energy E_D for the lowest fragmentation channel in eV at the PBE0/cc-pVTZ-PP level of theory. Recurring building units are highlighted in color: distorted octahedron (green), pentagonal bipyramid (blue), triple-capped trigonal prism (yellow), single- (violet) and double-capped square antiprism (red). All lowest-lying isomers have been found to be spin singlets. Center: The electric beam deflection profiles, given as arbitrary normalized intensity I as a function of the mean projection of the dipole moment on the field direction $\bar{\mu}_{\text{el},z}$, are shown with (red circles) and without applied electric field (black diamonds). The latter one has been fitted by a Gaussian around the center of deflection (solid black line). The electric deflection profiles were recorded at lowest possible nozzle temperatures of 16 K and a deflection voltage of 24 kV for clusters $N \leq 19$ as well as 30 K and 20 kV for $N > 19$ assuming uniform rotational temperatures of 20 K. The velocities were measured to be $453 \pm 3 \text{ ms}^{-1}$. Trajectory simulations have been performed employing moments of inertia, dipole moments and polarizabilities at the PBE0/cc-pVTZ-PP level of theory (dashed lines). Bottom: The photodissociation spectra, given as absorption cross section σ as a function of the photon energy $\hbar\omega$, are shown by discrete points (red circles) traced by an eight-point adjacent average (red solid line) to guide the eye. The photodissociation spectra were recorded at nozzle temperatures of 300 K (shown here) and 32 K (not shown here). For Sn₇, at 2.70 eV (shown to the right), 3.35 eV and 4.25 eV, indicated by the gray vertical bars, the depletion $1 - I/I_0$ has been plotted against the photon fluence ϕ double-logarithmically, such that the number of absorbed photons can be extracted from the slope (given in gray boxes).^{110,152,153} The rather large uncertainties are due to the nature of the depletion experiment and are estimated from the deviation determined by the depletion and fluence error bars. Significantly lower energy densities than the indicated 7 mJ cm^{-2} in order to avoid such multiphoton absorption processes cannot be easily applied due to vanishingly small differences in mass signal ratios. TDDFT simulations were obtained at the LC- ω PBEh/def2-TZVP level of theory by convoluting the discrete electronic transitions with Gaussians of an experimental FWHM of 0.27 eV at 300 K (dashed lines, shown here) and 0.20 eV at 32 K (dotted line, not shown here). Experimentally estimated dissociation energies are indicated by green arrows¹⁴ and computed dissociation energies at the PBE0/cc-pVTZ-PP level of theory for the energetically-lowest fragmentation channels averaged over all isomers are shown by red arrows when being above 1.8 eV.

3.7 and 4.1 eV result from the polar structure, the broad asymmetric band between 4.5–5.0 eV is mainly caused by the non-polar isomer. If the nozzle is cooled down to 32 K, the bands at 3.7 and 4.1 eV are enhanced, while the rising edge of the broad asymmetric band decreases. Hence, as the clusters are cooled, the fraction of the polar isomer increases significantly, just as observed in the electric deflection experiment at 16 K. Considering not only the intensity ratios directly obtained from electric deflection and spectral data, but also incorporating the impact of the temperature-dependent Boltzmann distribution (*cf.* ESI† for details), an upper bound for the energy difference ΔE

of the two isomers can be determined. Neglecting differences in vibrational frequencies and rotational constants, an estimated value of $\Delta E < 0.01 \text{ eV}$ is obtained, taking into account the different molecular point groups. Thus, the energy difference of almost 0.2 eV predicted theoretically is clearly too large. The fact that the CCSD(T) calculation only insignificantly improves on the relative energy of this isomer points towards the applied level of theory to be insufficient to accurately predict ΔE , possibly due to strongly contributing relativistic and static correlation effects. Due to the D_{2h} symmetry of isomer 8-II, a maximum isomer mixture I:II of 68:32 should be observed at large temperatures.



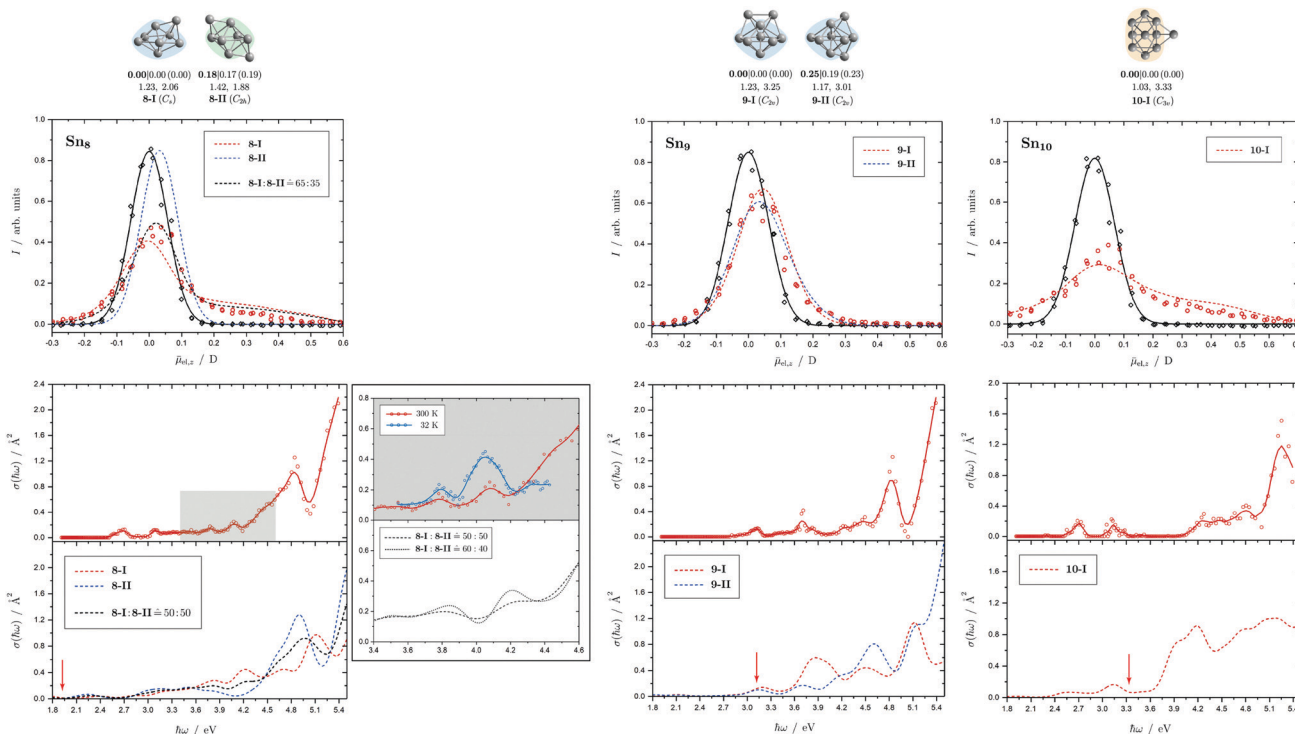


Fig. 5 Summary of all structural isomers (top), electric beam deflection measurements together with the corresponding trajectory simulations (center) and all photodissociation spectra together with the TDDFT simulations (bottom) for the clusters Sn_8 , Sn_9 and Sn_{10} . For details of this figure it is referred to the caption of Fig. 4. Bottom: For Sn_8 at 300 K an isomer ratio 8-I:8-II $\hat{=}$ 50:50 is obtained which is shifted to 8-I:8-II $\hat{=}$ 60:40 at 32 K.

Since the absorption spectrum at 300 K demonstrates isomer 8-II to be present in slightly higher proportion, the D_{2h} symmetry must be slightly broken which is also predicted at the LC- ω PBEh/def2-TZVPP level of theory (distortion towards C_s). Then a maximum isomer mixture of 50:50 should be possible which is also approximately confirmed in the optical experiment.

For Sn_9 , the previous quantum chemical calculations found a double-capped pentagonal bipyramid (9-II), which could explain the electric deflection behavior very well.³⁷ Global optimization now indicates a different structure as GM, which is also a double-capped pentagonal bipyramid (9-I). The electric dipole moments of the two structural isomers differ only slightly, such that the electric deflection experiment can also be described very well with the structure now predicted as GM. The absorption spectra are also similar, although the agreement with respect to the band signature in the region between 3.0 and 5.0 eV fits better with the new GM in comparison with the experiment. Therefore, it can be assumed that this structural isomer is experimentally present.

A quadruple-capped trigonal prism (C_{3v}) is found for Sn_{10} as the only low-energy structure in the global optimization. A distorted quadruple-capped trigonal prism with C_s symmetry, which has been identified as an energetically low-lying isomer in the cationic Sn_{10}^+ species, appears to be a transition state in the neutral system as discussed previously.³⁷ With the neutral structural isomer the electric deflection experiment can be described sufficiently well. The corresponding absorption spectrum shows a good agreement with the experimental photodissociation

spectrum. Both the two weak bands at 2.7 and 3.2 eV as well as the diffuse signature starting at 4.0 eV are correctly predicted theoretically, although the latter one is estimated to begin already at 3.7 eV by the TDDFT. With a comparatively large computed dissociation energy of 3.33 eV, this cluster is forecasted by the RRKM theory to fragment on the experimental time scale with one of the largest excess energies for its size range (*cf.* Fig. 3). This could be a reason why the shoulder of the absorption band at 3.7 eV in the theoretical absorption spectrum is suppressed in the photodissociation spectrum. Since the longest wavelength bands appear before and around this dissociation energy and, in comparison, have a high intensity, they are probably also due to multiphoton absorption processes.

5.3 Sn_{11} and Sn_{12}

The electric deflection experiment indicates that already at nozzle temperatures above 50 K, the clusters with more than ten tin atoms are thermally excited to such an extent that the structures are no longer rigid but partially floppy. Accordingly, structure discrimination with the electric deflection experiment is only possible if the clusters are cooled down accordingly. In addition, the probability of finding several quasidegenerate structural isomers tends to increase. This is already the case for Sn_{11} . Of the total of four predicted structural isomers, two are energetically nearly degenerate. On the one hand a triple-capped square antiprism (11-I) and on the other hand a quintuple-capped trigonal prism (11-II) is predicted. The isomers 11-III and 11-IV become even less plausible when



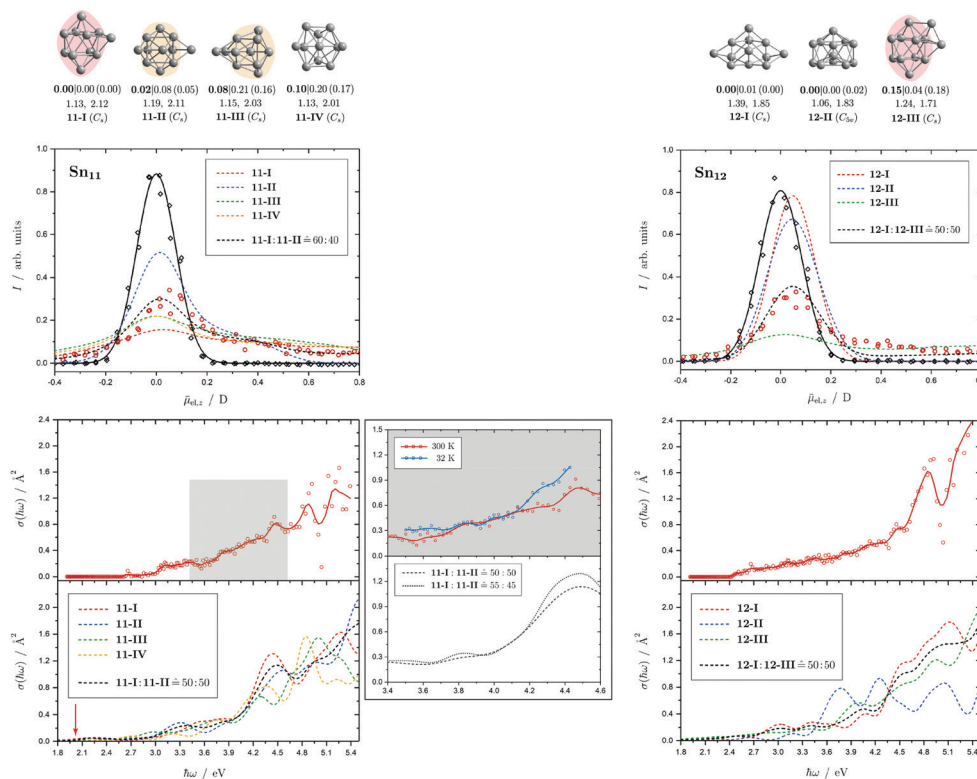


Fig. 6 Summary of all structural isomers (top), electric beam deflection measurements together with the corresponding trajectory simulations (center) and all photodissociation spectra together with the TDDFT simulations (bottom) for the clusters Sn_{11} and Sn_{12} . For details of this figure it is referred to the caption of Fig. 4. Bottom: For Sn_{11} at 300 K an isomer ratio 11-I : 11-II \approx 50 : 50 is obtained which is shifted to 11-I : 11-II \approx 55 : 45 at 32 K. For Sn_{12} both at 300 K and 32 K (not shown) a fitted isomer mixture of approximately 12-I : 12-III \approx 50 : 50 is found.

considering the CCSD(T) relative energies. The electric deflection experiment indicates that isomers 11-I and 11-II are present in an approximately similar ratio in the experiment. The calculated absorption spectra of both isomers can reproduce the measured spectrum quite well, especially the increase in light absorption from 3.1 eV. However, the calculated spectra possess a stronger patterning in the region between 3.5 and 5.4 eV than is observed experimentally. This could indicate that both structural isomers are present in the experiment, thus washing out the theoretically predicted signatures. Cooling the cluster nozzle down leads to a slight increase in the triple-capped square antiprism fraction, so that this structural isomer probably represents the GM.

For Sn_{12} , the structural isomers are predicted to be two quadruple-capped square antiprisms of which one is substantially distorted (12-I) and one being very polar (12-III) as well as an icosahedron with a missing corner (12-II). The electric deflection experiment indicates the presence of the very polar isomer 12-III and the less polar isomer 12-I. CCSD(T) calculations support this finding by lowering the relative energy of 12-III compared to the DFT results. The time-averaged electric dipole moment is already partially quenched at nozzle temperatures of 50 K, so that structural discrimination requires low nozzle temperatures here as well. The observed absorption spectrum can also be interpreted as a superposition of the two antiprisms, such that a slowly increasing light absorption starting at 2.5 eV is observed without

distinct absorption bands. Since the theoretically predicted absorption spectrum derived from the icosahedron has pronounced bands between 3.5 and 4.5 eV, this structural isomer can be excluded. It is interesting to note that the absorption spectrum of Sn_{12} is nearly independent of the nozzle temperature indicating quasidegenerate structural isomers.

5.4 Sn_{13} , Sn_{14} , Sn_{15} and Sn_{16}

The two structural isomers of Sn_{13} are derived on the one hand from a triple-capped trigonal prism on which four more atoms are attached peripherally (13-I), and on the other hand from a pentagonal bipyramid, which is connected to a triple-capped trigonal prism *via* a triangular plane (13-II). This results in a slightly prolate shape in each case. An icosahedral isomer (I_h) predicted for cationic Sn_{13}^+ , which has already been observed experimentally,⁴² is also found for the neutral species ($\Delta E = 1.42$ eV) since the a_g HOMO is not subject to Jahn-Teller (and pseudo Jahn-Teller) stabilization, but is not expected to be particularly stable due to a lowered HOMO-LUMO gap. The electric deflection measurements can best be explained by the presence of 13-II, although 13-I cannot be completely excluded. Isomer 13-II also explains the observed absorption spectrum very well. The overall result is a poorly structured broad light absorption that starts at 2.5 eV and then increases continuously to 5.4 eV. The pronounced absorption band of 13-I at 4.7 eV cannot be observed experimentally. However, a small amount



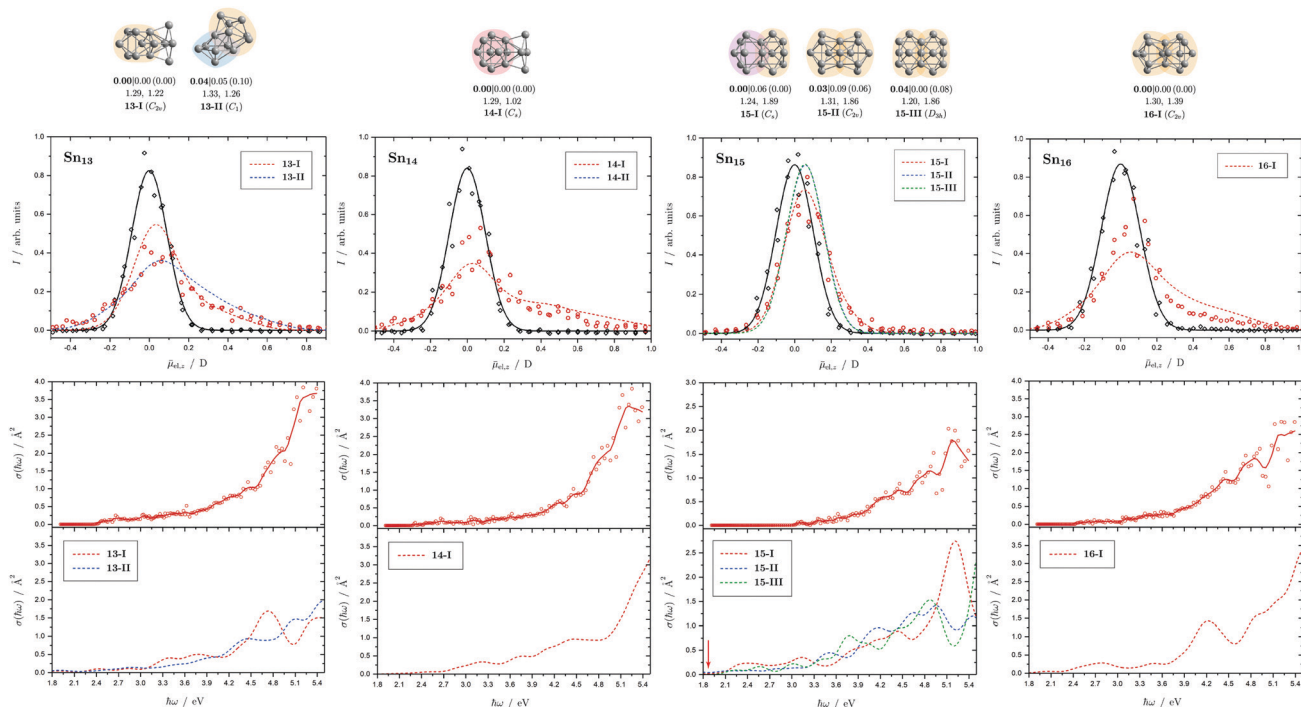


Fig. 7 Summary of all structural isomers (top), electric beam deflection measurements together with the corresponding trajectory simulations (center) and all photodissociation spectra together with the TDDFT simulations (bottom) for the clusters Sn₁₃, Sn₁₄, Sn₁₅ and Sn₁₆. For details of this figure it is referred to the caption of Fig. 4.

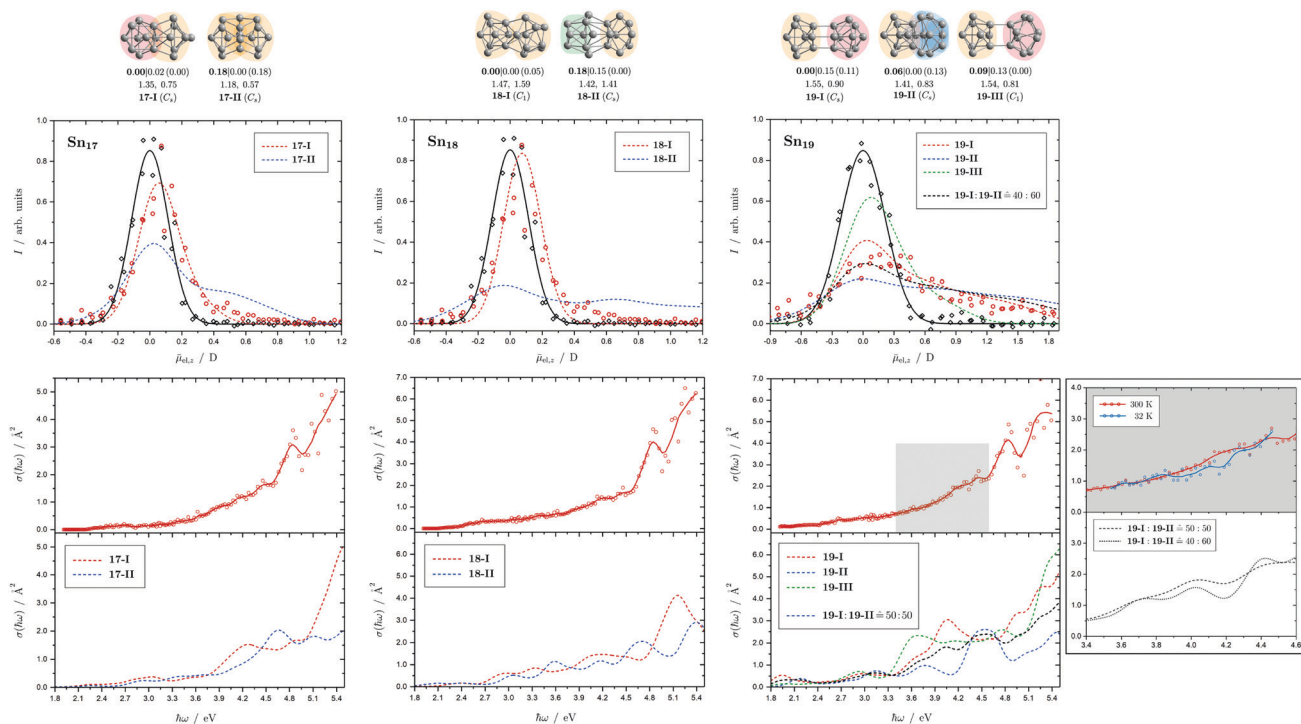


Fig. 8 Summary of all structural isomers (top), electric beam deflection measurements together with the corresponding trajectory simulations (center) and all photodissociation spectra together with the TDDFT simulations (bottom) for the clusters Sn₁₇, Sn₁₈ and Sn₁₉. For details of this figure it is referred to the caption of Fig. 4. Bottom: For Sn₁₉ at 300 K an isomer ratio 19-I : 19-II ≈ 50 : 50 is obtained which is shifted to 19-I : 19-II ≈ 40 : 60 at 32 K.



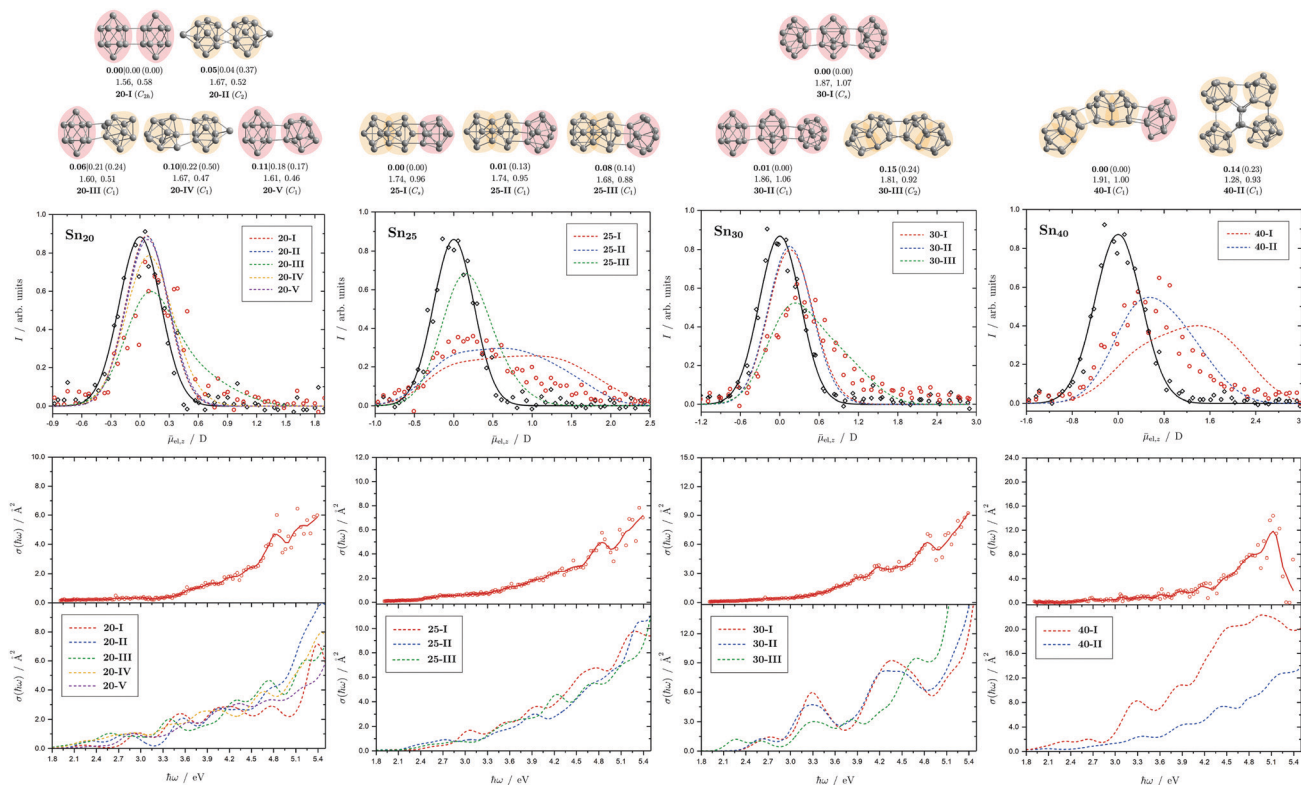


Fig. 9 Summary of all structural isomers (top), electric beam deflection measurements together with the corresponding trajectory simulations (center) and all photodissociation spectra together with the TDDFT simulations (bottom) for the clusters Sn_{20} , Sn_{25} , Sn_{30} and Sn_{40} . For details of this figure it is referred to the caption of Fig. 4.

of 13-I also cannot be excluded based on the absorption spectrum.

One structural isomer has been identified for Sn_{14} in the given energy range, which is derived from a double-capped square antiprism to which additional four atoms are attached on one side. Another isomer much higher in energy has already been discussed and can be excluded.⁵¹ The electric deflection data are matched reasonably well. The observed absorption spectrum of Sn_{14} is rather featureless, with light absorption starting at 2.6 eV, then remaining approximately constant until 3.5 eV, increasing and then raising again more strongly from 4.7 eV onward. The calculated absorption spectrum shows a similar course.

Three isomers of Sn_{15} were found here, which all share a triple-capped trigonal prism building unit. In 15-I it is connected *via* a triangular face with a capped square antiprism, whereas in 15-II and 15-III two trigonal prisms, each triple-capped, are linked by a triangular face. The electric deflection data are very well described by 15-I. The light absorption increases steadily starting from 3.0 eV and shows a distinct band at 5.0 eV, which is also theoretically obtained for 15-I. However, the predicted absorption bands between 2.0 and 3.0 eV are not observed experimentally, which may be related to the fact that the single-photon absorption taking place is not sufficient for the photodissociation of the clusters. The comparatively large dissociation energy in the RRKM picture supports this observation.

The predicted structure of Sn_{16} consists of two triple-capped trigonal prisms connected by a square plane. A series of related structures were found for the cationic Sn_{16}^+ , all corresponding to transition states of the neutral species. The neutral GM agrees moderately well with electric deflection measurements. The observed absorption spectrum can be well explained. On the one hand, this includes the onset of light absorption at 2.7 eV, which then remains approximately constant until 3.6 eV, followed by a stronger increase of the absorption cross sections until 5.4 eV. Only the theoretically predicted local maximum in the light absorption at 4.2 eV cannot be detected in such a pronounced way experimentally.

5.5 Sn_{17} , Sn_{18} and Sn_{19}

Two structural isomers are predicted for Sn_{17} , a prolate quadruple-capped trigonal prism attached *via* two atoms to a double-capped square antiprism (17-I) and a more compact isomer constructed by superimposition of three triple-capped trigonal prisms (17-II). The prolate structural isomer can describe the electric deflection data very well. The calculated absorption spectrum of isomer 17-I can also explain well the measured one, although the abrupt increase in light absorption at 4.0 eV is not particularly pronounced experimentally. The predicted rise at 4.5 eV to a plateau predicted for the more compact isomer is definitely not observed in experiment.

The structure of the predicted Sn_{18} GM (18-I) consists of two triple-capped trigonal prisms fused together. The second



structural isomer (18-II) contains a triple-capped trigonal prism connected to nine further atoms containing a distorted octahedron. While 18-II is very polar, 18-I has a low electric dipole moment and the electric deflection measurements are thus well described by isomer 18-I. The agreement between measured and calculated absorption spectra for 18-I is satisfactory, and in particular the strong increase in light absorption starting at 4.6 eV is well reproduced.

Three isomers are obtained for Sn_{19} , all of which are quite polar. The very prolate structures of 19-I and 19-III consist of two subunits linked together, each consisting of a triple-capped trigonal prism and a double-capped square antiprism. In contrast, 19-II is much more compact and also consists of a triple-capped trigonal prism connected to ten more atoms forming a superimposition of three pentagonal bipyramids. The electric deflection data indicate the presence of 19-I or 19-II, possibly both isomers are present together. All predicted optical spectra forecast a sudden increase in light absorption at 4.0 eV (19-I), 4.5 eV (19-II) and 3.5 eV (19-III). The fact that this is not observed experimentally points also to the presence of two structural isomers, presumably 19-I and 19-II, resulting in a rather uniform increase in light absorption. Lowering the nozzle temperature from 300 K to 32 K changes the course of the optical absorption slightly in favor of 19-II.

5.6 Sn_{20} , Sn_{25} , Sn_{30} and Sn_{40}

The structural elucidation of Sn_{20} using electric deflection data is particularly challenging. In fact, a total of five structural isomers were found in an energy range of 0.11 eV relative to the GM and all of them have considerably low vibrational frequencies (*cf.* corresponding frequency plot in ESI†). They are all strongly prolate structures built from two interconnected subunits, namely different combinations of quadruple-capped trigonal prisms and double-capped square antiprisms. Isomers 20-I, 20-II, and 20-V have electric dipole moments that are clearly too small to explain the electric deflection data. In contrast, the dipole moment of 20-III appears to be slightly too large and that of 20-IV again too small; possibly both isomers are also present together, with the proportion of the more polar isomer 20-III clearly predominating. The strong increase in light absorption predicted for 20-I, 20-II and 20-IV starting at 5.0 eV cannot be confirmed experimentally. 20-III also seems best suited to describe the course of the absorption spectrum.

For Sn_{25} , a total of three isomers were found with the GA. All isomers are combinations of two triple-capped trigonal prisms fused together *via* a triangular plane (two of them resembling isomer 15-II), each interconnected with a double-capped square antiprism. The electric deflection data can best be described by the GM or isomer 25-II, although the simulated beam profiles are further broadened and suggest a larger electric dipole moment. This can be explained by taking into account the vibrational excitation. Since the larger clusters exhibit very soft vibrational modes, it is reasonable to assume a significant vibrational excitation and, hence, a partial quenching of the observed electric dipole moment even at a nozzle temperature of 30 K. This is more pronounced for clusters with a larger electric dipole moment, which is why this effect is not as

prominent for the Sn_{20} cluster with even softer vibrational modes. The calculated absorption spectrum of isomer 25-I also fits very well to the rather structureless experimentally observed light absorption with a prominent peak at 4.8 eV, which, in contrast, is not predicted for 25-II and 25-III.

All structural isomers found for Sn_{30} resemble the prolate growth pattern either *via* interconnecting three double-capped square antiprisms (30-I and 30-II) or two 15-II units (30-III) indicating a similarity to the Sn_{25} cluster. The electric beam deflection profile is covered best by isomer 30-III (predicted as GM for the Sn_{30}^+ cluster), since 30-I and 30-II have dipole moment components being significantly too low. However, the additional presence of a very polar isomer being responsible for the large deflection is suggested. The absorption spectrum is again given by a monotonous increase with a characteristic signature between 3.9 and 5.1 eV. Since for the Sn_{30} cluster a significantly large excess energy is needed for photodissociation, the low-energy part of the spectrum is likely governed by multiphoton processes. Isomers 30-I and 30-II have strong peaks at 3.3 and 4.3 eV in common which are not observed experimentally and can thus be ruled out. Isomer 30-III yields a better correspondence although the features starting at 3.9 eV experimentally are predicted to appear approximately 0.8 eV below. Starting from Sn_{30} , the experimental absorption cross sections are substantially lower than forecasted by the TDDFT implying that non-dissociative relaxation pathways might play a growing role in clusters of this size regime.

Interestingly, two structurally distinct isomers for Sn_{40} were found. Isomer 40-I continues the prolate and thread-like shape evolution by adding a double-capped square antiprism to a 30-III unit and isomer 40-II fuses four triple-capped trigonal prisms together peripherally *via* four central atoms forming an oblate geometry and opening up the appearance of quasispherical structures. The electric deflection data disagree with the more polar isomer 40-I due to its overestimated electric dipole moment. Isomer 40-II gives a much better fit with deviations in the simulated deflection profile within the expected error range. The onset of light absorption in the photodissociation spectrum starts around 2.4 eV, which, compared to the particularly large excess energy needed for photodissociation, hints again towards involved multiphoton phenomena. The measured data points exhibit more scattering being due to a lower abundance in the mass spectra. Therefore, the decline in absorption cross section > 5.1 eV, where also the photon fluence is lowest and most fluctuating, has to be regarded with care. Whereas the strongly prolate isomer 40-I continues the trend of increasing light absorption with cluster size, the oblate isomer 40-II provides an absorption spectrum of significantly lower absorption cross section much better fitting the experimental course.

6 Discussion

6.1 Structural motifs and shape evolution

There are a few theoretical studies which deal with the structures of neutral Sn_N clusters up to $N = 20$,^{29–36,38,40,43,44,48–50} but



only one joint computational and experimental investigation³⁷ our results will be compared to. The structure elucidation is strongly improved by carrying out electric deflection measurements at 16 K together with additional photodissociation spectroscopy at 300 and 32 K as well as more sophisticated quantum chemical calculations. The pentagonal bipyramid found in the clusters with $N = 6-9$ is demonstrated not to be a dominant building unit for the assembly of larger clusters, but rather are the triple-capped trigonal prism and double-capped square antiprism which emerge at $N = 10$ and 11. New isomers could be confirmed to be (pro rata) present in the molecular beam for the clusters Sn_9 , Sn_{11} , Sn_{12} , Sn_{15} , Sn_{17} , Sn_{18} , Sn_{19} and Sn_{20} which were either not found before by the GA or did not show enough differences in their electric deflection profiles alone for an unambiguous assignment. The experimental results of the clusters Sn_8 , Sn_{11} , Sn_{12} and Sn_{19} suggest that more than one isomer, often differing in their polarities, are present in the molecular beam simultaneously, thus giving rise to a temperature-dependent study of their mixture and the energetics involved. While for the clusters Sn_8 , Sn_{11} and Sn_{19} a lowering of the nozzle temperature could adjust the isomer mixture in favor of a thermodynamically more stable structure, the cluster Sn_{12} shows a uniform distribution of the present isomers over the investigated temperature range.

The prolate growth pattern and the gradual transition to more compact or quasispherical structures of the semiconductor clusters Si, Ge and Sn is well-known by now and was first probed experimentally by Jarrold *et al.* who performed ion mobility measurements on Sn_N^+ clusters.⁶⁰ Up to at least $N = 65$ elongated geometries were found with quasispherical isomers starting to appear from $N = 35$, thus indicating a delayed structural rearrangement compared to cationic Si and Ge clusters.^{61,62} For Pb clusters quasispherical structures were discovered for all size regimes. However, non-vanishing electric dipole moments and anomalies in the electronic polarizabilities support a covalent rather than metallic bonding up to at least $N = 36$.¹⁵⁴ Cui *et al.* additionally found indirect evidence for the shape transition by recording photoelectron spectra of Sn_N^- clusters which pointed towards a semiconductor-to-metal transition at $N = 42$.⁸⁰ Many of the theoretically predicted structures were identified both in the neutral and cationic species, although often significantly distorted in the latter, but experimental evidence for the shape transition in neutral clusters is still sparse. Bachelors and Schäfer estimated the crossover region from elongated to quasispherical geometries calorimetrically to occur at $N = 45-85$.^{99,100} From our selected studies on the clusters Sn_{25} , Sn_{30} and Sn_{40} we can qualitatively confirm this observation. It is reasonable to believe that at least up to $N = 30$ prolate clusters are dominantly present in the molecular beam studies. The thread-like growth is characterized by connecting triple-capped trigonal prisms and double-capped square antiprisms as building blocks whose appearance already dates back to the smallest clusters under investigation. With Sn_{40} adopting an oblate plate-like motif, the gradual crossover in structure is limited to happen at $30 < N \leq 40$. This is also in agreement with DFT studies of Zang *et al.* who predict not only plate-like isomers for $40 \leq N \leq 44$, but also the existence of Y-shaped isomers for $36 \leq N \leq 39$ as emerging

quasispherical structures.⁴⁶ Li *et al.* further give indications for the presence of thread-like isomers as low-energy competitors up to $N = 50$ ⁴⁴ and above.⁴⁷

The evolution of elongated structures is also resembled by the cluster's binding energy, *i.e.* for an atomization or arbitrary dissociation process. As can be seen from Fig. 2, the dissociation energy for the lowest-energy fragmentation channel drops successively as the clusters start to grow in a prolate manner. Once a certain size is reached, it stays approximately constant due to the nearly invariable surface-to-volume ratio of the clusters, *i.e.* due to the approximately size-independent binding energy per atom.¹⁰⁰

6.2 Microscopic impact of shape on optical absorption

The electronic transitions in the absorption spectra of the Sn_N clusters below 7 eV are given by a complicated manifold of $5p \rightarrow 5p$ excitations with significant contributions of $5s$ orbitals both in the initial and final states. A d orbital polarization is present to a larger extent compared to Si and Ge clusters, but still pretty small. Already in the smallest clusters absorption bands correspond to many single-electron excitations with non-negligible oscillator strengths, which itself are most often described by more than two significantly contributing NTO pairs, impeding a clear and simple characterization of the nature of certain regions of the absorption spectra. It has been shown for Sn_6 that several reference configurations are required in order to adequately treat the low-lying excited states.²⁷ However, the oftentimes satisfactory up to quantitative agreement of the TDDFT spectra with the experimental course suggests that at least for the general features the disregard of multireference (and spin-orbit) effects is justified in several cases. Absorption spectra of the smallest clusters are dominated by distinct features which can be traced back to their molecular shape and symmetry as well as well separated (discrete) energy levels. From Sn_{12} onward, together with the approximate start of the prolate growth, the spectra become less inimitable, showing now a monotonous increase in light absorption. Interestingly, this observation is on par with findings from the absorption spectra of silicon clusters.¹⁵⁵⁻¹⁶⁰ This can partly be understood by the growing density of electronic states with cluster size, but does not explain fully the overall lack of features compared to the TDDFT absorption spectra. Although the vibrational finestructure has not been resolved in the photodissociation spectra, the large extent of vibrational excitation present at 300 K is causing a complicated hot band signature which is expected to, at least to some degree, alter the overall course of the measured spectrum. As already discussed in detail, the low-energy regime of the photodissociation spectra are frequently governed by multiphoton absorption as well as hidden absorption characteristics due to the cluster's excess energies needed for dissociation on the experimental time scale. Therefore, no reliable information can be extracted from the experimental absorption spectra and excited states computed by TDDFT have to be considered. In this regard no band gap closure has been observed for the size range studied and the optical band gap with an oscillator strength $f \geq 0.005$ is still above 1.0 eV for the Sn_{30} and Sn_{40} cluster, *inter alia*, because of their relative



stability (*cf.* band gap plot in ESI[†]) as predicted previously.⁴⁴ In the high-energy regime of the absorption spectra of larger clusters charge-transfer excitations become quite frequent, although they never appear as distinct and individual features.

Based on the analysis of the calculated electronic excitations, it would be of great interest to develop a microscopic picture which links the structural transition of the clusters to their optical absorption. This relation is well-documented for classical spheroids by the Mie–Gans theory utilizing dielectric functions of the corresponding bulk material.^{105,161,162} Whether this behavior also holds for the smallest clusters and if a plausible picture can be drawn quantum chemically is discussed in the following. Therefore, two cases have been examined in detail, the Sn₁₉ cluster which exhibits a more and less prolate isomer as well as the Sn₄₀ cluster which is shown to be strongly oblate, but has a very prolate isomer in theory too. The TDDFT spectra of the isomers 19-I and 19-II shown again in Fig. 10 are comparable in a sense that their low-energy region shares the same characteristics followed by an intense band at approximately 4.0 eV and 4.5 eV, respectively, before the absorption cross section continues to increase further. Interestingly, when the bonds connecting the two subunits in the more prolate isomer 19-I are compressed slightly, it is the

intense band which is shifted towards higher photon energies by roughly 0.6 eV causing a subsequent increase of the absorption cross section now appearing at > 5.0 eV. Due to the large amount of electronic transitions involved in this band, an unambiguous orbital picture could not be derived for this geometric alteration. However, it can be understood qualitatively in classical terms. The absorption spectrum of the undistorted isomer 19-I is quite well resembled by the Mie–Gans theory for a prolate spheroid with geometric dimensions taken from quantum chemistry. Upon distortion the predicted spectral course approaches the result of the Mie–Gans theory of a perfect sphere. The TDDFT spectrum of isomer 19-II being less prolate is already close to the result of a sphere. Regarding the prolate isomer 40-I and the experimentally suspected oblate isomer 40-II a similar agreement with the Mie–Gans theory is observed as depicted in Fig. 11.

One quantum chemical conspicuity is, however, worth noting. When comparing the distorted isomer 19-I to its undistorted structure (and also isomer 19-II to 19-I and isomer 40-II to 40-I) an increase in kinetic energy by about 1.5 eV is observed which can be traced back to those molecular orbitals involved in the transitions > 3.5 eV. This situation is remotely reminiscent of the particle-in-a-cylinder model which has been applied to explain longitudinal plasmon resonances in the absorption spectra of metal nanorods.¹⁶³ At first glance this problem does not seem to be transferable to prolate tin clusters with their intricate manifold of excitations leading to broad

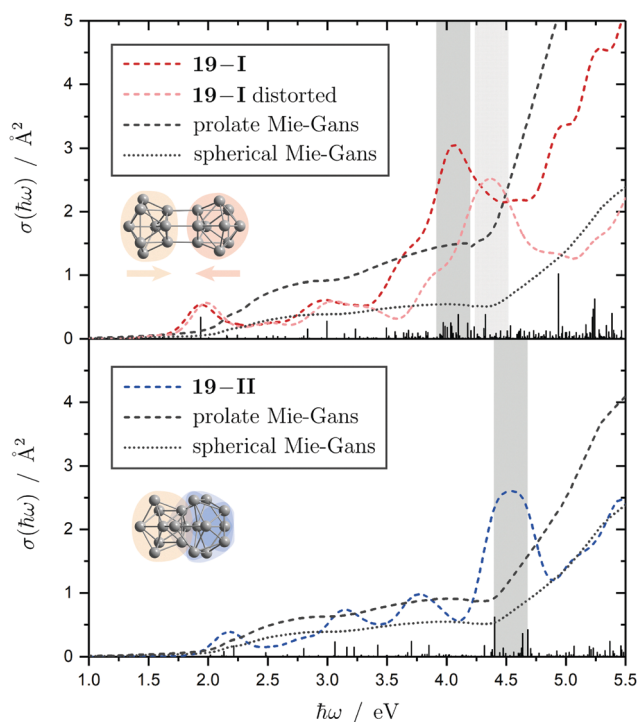


Fig. 10 Calculated Sn₁₉ absorption spectra for isomers 19-I (red) and 19-II (blue) at the LC- ω PBEh/def2-TZVPP level of theory. The discrete electronic transitions are convoluted by Gaussians with a FWHM of 0.27 eV. Characteristic absorption peaks are highlighted by gray bars. Additionally, an absorption spectrum is shown where the more prolate isomer 19-I is compressed about 0.15 Å along its C_s mirror plane (light red). The results of the Mie–Gans theory for a spherical and prolate particle (with an aspect ratio¹⁰⁵ of $A/B = 2.4$ for isomer 19-I and $A/B = 1.6$ resembling isomer 19-II) are depicted for comparison.

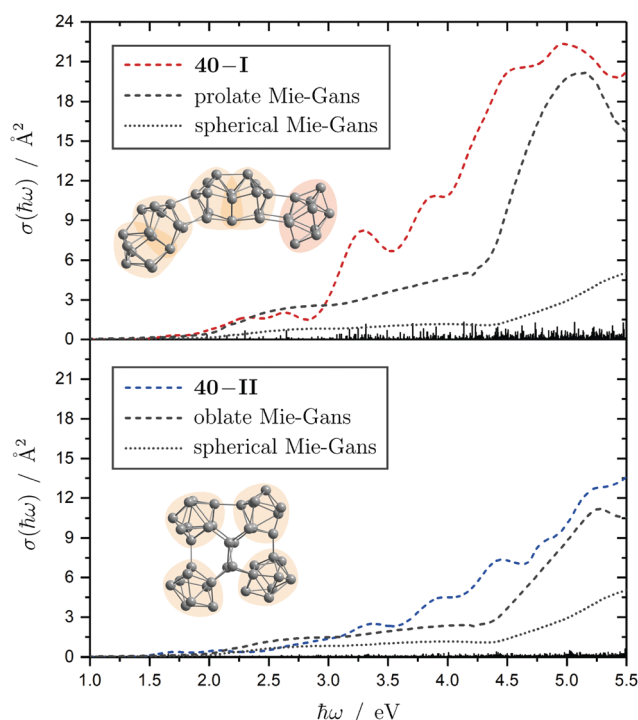


Fig. 11 Calculated Sn₄₀ absorption spectra for isomers 40-I (red) and 40-II (blue) at the LC- ω PBEh/def2-TZVPP level of theory. The discrete electronic transitions are convoluted by Gaussians with a FWHM of 0.27 eV. The results of the Mie–Gans theory for a spherical and prolate/oblate particle (with an aspect ratio¹⁰⁵ of $A/B = 3.7$ for isomer 40-I and $A/B = 0.4$ resembling isomer 40-II) are depicted for comparison.



absorption bands, but given that band gaps of semiconductor nanowires were also shown to depend on the wire diameter,^{164,165} the applicability of this approach should be further tested.

6.3 Limitations of the experimental methodology

In this study it became clear that both the electric deflection and the photodissociation spectroscopy experiment, though oftentimes being interpreted successfully, suffer from certain limitations which will be summarized in the following. The key assumption to interpret the electric deflection profiles is based on the clusters being rigid rotors. Especially larger Sn_N clusters, as discussed for $N \geq 20$, exhibit low vibrational modes and hence become floppy as a result of thermal excitation which partially quenches the electric dipole moment even at nozzle temperatures below 30 K. The electric dipole moments required for the classical trajectory simulations stem from DFT calculations and are often attributed with an error of up to 20%. Moreover, the rotational temperature, being responsible for populating the rotational states of the clusters in the trajectory simulation, is not known *a priori* and has to be estimated. For reasons of consistency the same rotational temperature of $T_{\text{rot}} = 20$ K is applied for all clusters, although larger clusters with $N \geq 19$ have been measured at higher nozzle temperatures of 30 K and thus likely have slightly higher rotational temperatures, too. In the ESI[†] electric dipole moments and polarizabilities have been deduced from the experimental deflection profiles by fitting a Gaussian function containing information about the broadening and deflection based on a first-order perturbation theory treatment. Calculated effective polarizabilities based on second-order perturbation theory¹⁰² as a function of the experimental electric dipole moments agree well with experimentally extracted polarizabilities giving confidence in the chosen rotational temperature of $T_{\text{rot}} = 20$ K. This holds for many clusters even though they are treated as spherically-symmetric rotors, a clear contradiction to most of the clusters showing strongly prolate geometries. However, larger clusters exhibit effective polarizabilities which are too small compared to the experimentally extracted ones stressing again the thermal excitation of weak vibrational modes and the consequent quenching of the electric dipole moment. In some cases a Gaussian function is not an appropriate choice for the fit as is discussed for, *i.e.*, Sn_{11} in the ESI.[†] Whenever two isomers seem to be present in the molecular beam, two Gaussian functions need to be fitted hampering this task further and adding a certain degree of freedom to those deduced parameters.

The photodissociation spectrum corresponds to the true absorption spectrum only when there are no non-dissociative relaxation pathways present. The probability for this is expectedly altered over the studied size range of the clusters due to an increasing density of excited states with cluster size. In the spectra of the largest clusters the measured absorption cross sections are quite a bit lower than predicted by TDDFT which could owe to the mentioned phenomenon. Larger clusters with many vibrational modes and low dissociation energies are shown to exhibit great excess energies which impede a fragmentation on the experimental timescale rendering the

light absorption invisible to this action spectroscopy. The excess energy takes values up to >4 eV (for the Sn_{40} cluster) scrutinizing a large region of the spectrum as to whether it can be simply compared to TDDFT results. The low-energy regime is further complicated due to the tin clusters being prone to multiphoton absorption processes as demonstrated for the Sn_7 cluster and suspected to occur in several other species. Because of this no band gaps have been extracted here. Although the vibronic finestructure is not resolved, the significant excitation of virtually all vibrational modes at 300 K causes a complicated hot band structure which potentially alters the general course of the photodissociation spectrum slightly. Together with the electric deflection experiment, information on the energetics of the clusters have been obtained whenever more than one isomer were found to be present in the molecular beam simultaneously. This was possible due to the global assumption that the clusters are equilibrated within the nozzle. Although there is certain evidence for this assumption, it is still somewhat debatable whether the clusters are indeed perfectly equilibrated. It is also not fully clarified if a Boltzmann distribution of the thermally excited vibrational modes is the most appropriate assumption under the given conditions.

7 Conclusions

Electric beam deflection measurements and photodissociation spectroscopy together with quantum chemical calculations are applied to shine light on the geometric evolution as well as to unravel the optical absorption of neutral Sn_N clusters with $N = 6-20, 25, 30, 40$. In comparison to a previous study³⁷ the size range of the clusters is extended and the dielectric response is probed at a lower nozzle temperature of 16 K. An improved GA routine and more sophisticated levels of theory, including the hybrid functionals PBE0 and LC- ω PBEH plus CCSD(T) calculations, are employed for the structural reoptimization as well as the calculation of properties such as the electric dipole moments and polarizabilities. In this regard, new energetically low-lying isomers for clusters with $N = 9, 11, 12, 15, 17-20$ are confirmed to be (partially) present in the molecular beam. Triple-capped trigonal prisms and double-capped square antiprisms are identified as dominant building blocks over almost the entire size range with the former one building Sn_{15} sub-units which are particularly conspicuous for clusters with $N > 20$. The clusters are shown to adopt prolate shapes up to at most the oblate Sn_{40} cluster from where quasispherical structures are expected to gradually take over. Although this structural transition is well known from studies on charged tin clusters, it can be regarded the first size-specific experimental evidence of this transition for neutral tin clusters. Fragmentation likely occurs *via* the loss of a tin atom for clusters with $N < 12$, a Sn_7 cluster for $N < 19$ and a Sn_{10} cluster for $N \geq 19$ based on computation and evidence given by photodissociation. The impact of the rotational temperature, the electric dipole moment and thermal excitation of vibrational modes on simulating electric deflection profiles is discussed diligently.



The latter manifests in the clusters becoming floppy and, hence, quenching the electric dipole moment to some extent. This was shown to be especially important for larger clusters with $N \geq 19$.

Photodissociation spectra of neutral tin clusters have been recorded for the first time illustrating similarities to absorption spectra of small silicon clusters. The spectra show distinct absorption features for the smallest clusters under study and become less structured with increasing cluster size starting approximately at $N = 12$ coinciding with the cluster's prolate growth. In many cases a comparison between the photodissociation spectra and absorption spectra predicted by TDDFT complement the results from the electric beam deflection leading to a more explicit structural elucidation. The electronic transitions are found to consist of a manifold of $5p \rightarrow 5p$ excitations with significant contribution of $5s$ orbitals hampering an unambiguous classification of the observed absorption bands and their relation to structure. However, the general course of the absorption spectra can be understood in terms of the classical Mie-Gans theory for prolate/oblate spheroids and spheres. By varying the laser fluence multiphoton absorption is detected to be present in the low-energy regime of the spectra of many clusters. Furthermore, the dissociation kinetics on the experimental time scale between photoexcitation and detection play a tremendous role. RRKM theory is applied to estimate excess energies required for excitation energies being sufficient to cause dissociation. These can be as low as 0.6 eV for the smallest cluster, Sn_6 , and can raise up to >4.0 eV for Sn_{40} rendering large parts of the absorption spectrum invisible for photodissociation. Thermal excitation of vibrational modes and coupling of vibrational and electronic degrees of freedom is expected to also contribute to the spectra recorded at 300 K, further washing out the spectral signature possibly due to hot bands.

In both conducted experiments the simultaneous presence of two isomers for the clusters with $N = 8, 11, 12$ and 19 is observed. By comparing the isomer ratios in the electric deflection experiment at 16 K and photodissociation spectroscopy experiment at 32 K and 300 K, information about the energetics and dynamics of the structural rearrangement of the cluster isomers can be obtained. The findings scrutinize the isomer's relative energies which seem to be overestimated by quantum chemistry. Here, it would be valuable if more sophisticated results incorporating the multireference and relativistic nature become available, but also to tackle vibronic excitations. For the future it is highly desirable to merge dielectric deflection and absorption spectroscopy in a combined experiment in order to perform isomer-selective (photodissociation) spectroscopy on these species.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Financial support for this project was provided by the Deutsche Forschungsgemeinschaft (Grant SCHA 885/15-2). A. L. thanks

the Verband der Chemischen Industrie for a scholarship. Extensive calculations were conducted for this work on the Lichtenberg high-performance computer of the Technical University of Darmstadt. The authors thank the Hessian Competence Center for High Performance Computing funded by the Hessen State Ministry of Higher Education, Research, and the Arts for helpful advice.

References

- J.-X. Zhang, F. K. Sheong and Z. Lin, *50th Anniversary of Electron Counting Paradigms for Polyhedral Molecules. Structure and Bonding*, Springer International Publishing, 2021, ch. 5, pp. 197–257.
- L. D. Anh, K. Takase, T. Chiba, Y. Kota, K. Takiguchi and M. Tanaka, *Adv. Mater.*, 2021, **33**, 2104645.
- C. C. Striemer, R. Krishnan and P. M. Fauchet, *JOM*, 2004, **56**, 20–25.
- L. Pavesi, L. Dal Negro, C. Mazzoleni, G. Franzò and F. Priolo, *J. Appl. Phys.*, 2004, **96**, 3164–3171.
- L. Pavesi, *Silicon Nanocrystals: Fundamentals, Synthesis and Applications*, Wiley-VCH, Weinheim, 2010.
- X. Wen, P. Zhang, T. A. Smith, R. J. Anthony, U. R. Kortshagen, P. Yu, Y. Feng, S. Shrestha, G. Coniber and S. Huang, *Sci. Rep.*, 2015, **5**, 12469.
- W. M. Zhang, J. S. Hu, Y. G. Guo, S. F. Zheng, L. S. Zhong, W. G. Song and L. J. Wan, *Adv. Mater.*, 2008, **20**, 1160–1165.
- K. Kravchyk, L. Protesescu, M. I. Bodnarchuk, F. Krumeich, M. Yarema, M. Walter, C. Guntlin and M. V. Kovalenko, *J. Am. Chem. Soc.*, 2013, **135**, 4199–4202.
- Z. Zhu, S. Wang, J. Du, Q. Jin, T. Zhang, F. Cheng and J. Chen, *Nano Lett.*, 2014, **14**, 153–157.
- L. R. Parent, Y. Cheng, P. V. Sushko, Y. Shao, J. Liu, C. M. Wang and N. D. Browning, *Nano Lett.*, 2015, **15**, 1177–1182.
- R. E. Honig, *J. Chem. Phys.*, 1953, **21**, 573–574.
- K. A. Gingerich, D. L. Cocke and U. V. Choudary, *Inorg. Chim. Acta*, 1975, **14**, 10–12.
- K. A. Gingerich, E. A. Ramakrishnan and J. E. Kingcade, *High Temp. Sci.*, 1986, **21**, 1–15.
- G. Meloni, R. W. Schmude, J. E. Kingcade and K. A. Gingerich, *J. Chem. Phys.*, 2000, **113**, 1852–1856.
- V. E. Bondybey, M. Heaven and T. A. Miller, *J. Chem. Phys.*, 1983, **78**, 3593–3598.
- K. Pak, M. F. Cai, T. P. Dzugan and V. E. Bondybey, *Faraday Discuss. Chem. Soc.*, 1988, **86**, 153–162.
- A. B. Anderson, *J. Chem. Phys.*, 1975, **63**, 4430–4456.
- L. L. Lohr, *Inorg. Chem.*, 1981, **20**, 4229–4235.
- K. Balasubramanian and K. S. Pitzer, *J. Chem. Phys.*, 1983, **78**, 321–327.
- G. Pacchioni, *Mol. Phys.*, 1985, **55**, 211–223.
- D. Dai and K. Balasubramanian, *J. Chem. Phys.*, 1986, **85**, 3401–3406.
- D. Dai and K. Balasubramanian, *J. Phys. Chem.*, 1992, **96**, 9236–9243.



- 23 D. Dai and K. Balasubramanian, *J. Chem. Phys.*, 1992, **96**, 8345–8353.
- 24 W. Plass, H. Stoll, H. Preuss and A. Savin, *J. Mol. Struct.*, 1995, **339**, 67–81.
- 25 D. Dai and K. Balasubramanian, *J. Chem. Phys.*, 1996, **100**, 19321–19325.
- 26 D. Dai and K. Balasubramanian, *J. Chem. Phys.*, 1998, **108**, 4379–4385.
- 27 C. Zhao and K. Balasubramanian, *J. Chem. Phys.*, 2001, **115**, 3121–3133.
- 28 L. Si-Dian, R. Johnston and J. Murrell, *Acta Phys.-Chim. Sin.*, 1993, **9**, 642–649.
- 29 P. Jackson, I. G. Dance, K. J. Fisher, G. D. Willett and G. E. Gadd, *Int. J. Mass Spectrom. Ion Process.*, 1996, **157/158**, 329–343.
- 30 B. Wang, L. M. Molina, M. J. López, A. Rubio, J. A. Alonso and M. J. Stott, *Ann. Phys.*, 1998, **7**, 107–119.
- 31 Z. Y. Lu, C. Z. Wang and K. M. Ho, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2000, **61**, 2329–2334.
- 32 C. Jo and K. Lee, *J. Chem. Phys.*, 2000, **113**, 7268–7272.
- 33 C. Majumder, V. Kumar, H. Mizuseki and Y. Kawazoe, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2001, **64**, 233405.
- 34 R. Pushpa, S. Narasimhan and U. Waghmare, *J. Chem. Phys.*, 2004, **121**, 5211–5220.
- 35 C. Majumder, V. Kumar, H. Mizuseki and Y. Kawazoe, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2005, **71**, 035401.
- 36 S. Krishnamurty, K. Joshi, D. G. Kanhere and S. A. Blundell, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2006, **73**, 045419.
- 37 S. Schäfer, B. Assadollahzadeh, M. Mehring, P. Schwerdtfeger and R. Schäfer, *J. Phys. Chem. A*, 2008, **112**, 12312–12319.
- 38 C. C. Yang and S. Li, *J. Phys. Chem. C*, 2009, **113**, 14207–14212.
- 39 E. Oger, R. Kelting, P. Weis, A. Lechtken, D. Schooss, N. R. Crawford, R. Ahlrichs and M. M. Kappes, *J. Chem. Phys.*, 2009, **130**, 124305.
- 40 B. Assadollahzadeh, S. Schäfer and P. Schwerdtfeger, *J. Comput. Chem.*, 2010, **31**, 929–937.
- 41 A. Lechtken, N. Drebov, R. Ahlrichs, M. M. Kappes and D. Schooss, *J. Chem. Phys.*, 2010, **132**, 211102.
- 42 N. Drebov, E. Oger, T. Rapps, R. Kelting, D. Schooss, P. Weis, M. M. Kappes and R. Ahlrichs, *J. Chem. Phys.*, 2010, **133**, 224302.
- 43 N. Drebov and R. Ahlrichs, *J. Chem. Phys.*, 2011, **134**, 124308.
- 44 H. Li, H. Du, W. Chen, Q. Q. Shan, Q. Sun, Z. X. Guo and Y. Jia, *J. Phys. Chem. C*, 2012, **116**, 231–236.
- 45 A. Wiesel, N. Drebov, T. Rapps, R. Ahlrichs, U. Schwarz, R. Kelting, P. Weis, M. M. Kappes and D. Schooss, *Phys. Chem. Chem. Phys.*, 2012, **14**, 234–245.
- 46 Q. Zang, G. Chen and W. Lu, *Chem. Phys. Lett.*, 2012, **552**, 69–72.
- 47 H. Li, W. Chen, F. Wang, Q. Sun, Z. X. Guo and Y. Jia, *Phys. Chem. Chem. Phys.*, 2013, **15**, 1831–1836.
- 48 Q. J. Zang, G. J. Chen, W. Qin, L. Z. Zhao and W. C. Lü, *Chem. Res. Chin. Univ.*, 2013, **29**, 579–583.
- 49 P. G. A. Leyva, E. M. Sosa-Hernández, J. M. Montejano-Carrizales and F. Aguilera-Granja, *Eur. Phys. J. D*, 2015, **69**, 1–7.
- 50 D. Wu, Q. Du, X. Wu, R. Shi, L. Sai, X. Liang, X. Huang and J. Zhao, *J. Chem. Phys.*, 2019, **150**, 174304.
- 51 A. Lehr, M. Jäger, M. Gleditzsch, F. Rivic and R. Schäfer, *J. Phys. Chem. Lett.*, 2020, **11**, 7827–7831.
- 52 J. Drowart and P. Goldfinger, *Angew. Chem.*, 1967, **13**, 589–628.
- 53 T. P. Martin and H. Schaber, *J. Chem. Phys.*, 1985, **83**, 855–858.
- 54 J. R. Heath, Y. Liu, S. C. O'Brien, Q.-L. Zhang, R. F. Curl, F. K. Tittel and R. E. Smalley, *J. Chem. Phys.*, 1985, **83**, 5520–5526.
- 55 J. C. Phillips, *J. Chem. Phys.*, 1987, **87**, 1712–1716.
- 56 K. LaiHing, R. G. Wheeler, W. L. Wilson and M. A. Duncan, *J. Chem. Phys.*, 1987, **87**, 3401–3409.
- 57 M. Watanabe, Y. Saito, S. Nishigaki and T. Noda, *Jpn. J. Appl. Phys.*, 1988, **27**, 344–347.
- 58 D. S. Cornett, M. Peschke, K. LaiHing, P. Y. Cheng, K. F. Willey and M. A. Duncan, *Rev. Sci. Instrum.*, 1992, **63**, 2177–2186.
- 59 S. Yoshida, K. Fuke and I. Introduction, *Chem. Phys.*, 1999, **111**, 3880–3890.
- 60 A. A. Shvartsburg and M. F. Jarrold, *Phys. Rev. A: At., Mol., Opt. Phys.*, 1999, **60**, 1235–1239.
- 61 A. A. Shvartsburg and M. F. Jarrold, *Chem. Phys. Lett.*, 2000, **317**, 615–618.
- 62 A. A. Shvartsburg, R. R. Hudgins, P. Dugourd and M. F. Jarrold, *Chem. Soc. Rev.*, 2001, **30**, 26–35.
- 63 Y. Tai, J. Murakami, C. Majumder, V. Kumar, H. Mizuseki and Y. Kawazoe, *J. Chem. Phys.*, 2002, **117**, 4317–4322.
- 64 Y. Tai, J. Murakami, C. Majumder, V. Kumar, H. Alizuseki and Y. Kawazoe, *Eur. Phys. J. D*, 2003, **24**, 295–298.
- 65 G. Ganteför, M. Gausa, K. H. Meiwes-Broer and H. O. Lutz, *Faraday Discuss.*, 1988, **86**, 197–208.
- 66 G. Ganteför, M. Gausa, K. H. Meiwes-Broer and H. O. Lutz, *Z. Phys. D*, 1989, **12**, 405–409.
- 67 J. Ho, M. L. Polak and W. C. Lineberger, *J. Chem. Phys.*, 1992, **96**, 144–154.
- 68 J. Thøgersen, L. D. Steele, M. Scheer, C. A. Brodie and H. K. Haugen, *J. Phys. B*, 1996, **29**, 1323–1330.
- 69 V. D. Moravec, S. A. Klopčič and C. C. Jarrold, *J. Chem. Phys.*, 1999, **110**, 5079–5088.
- 70 Y. Negishi, H. Kawamata, A. Nakajima and K. Kaya, *J. Electron Spectros. Relat. Phenomena*, 2000, **106**, 117–125.
- 71 S. Osmekhin, M. Tchapyguine, M. H. Mikkilä, M. Huttula, T. Andersson, O. Björneholm and S. Aksela, *Phys. Rev. A: At., Mol., Opt. Phys.*, 2010, **81**, 023203.
- 72 A. A. Shvartsburg and M. F. Jarrold, *Phys. Rev. Lett.*, 2000, **85**, 2530–2532.
- 73 H. Haberland, *Phys. World*, 2000, **13**, 27–28.
- 74 K. Joshi, D. G. Kanhere and S. A. Blundell, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2002, **66**, 155329.
- 75 K. Joshi, D. G. Kanhere and S. A. Blundell, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2003, **67**, 235413.



- 76 F. C. Chuang, C. Z. Wang, S. Ögüt, J. R. Chelikowsky and K. M. Ho, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2004, **69**, 165408.
- 77 G. A. Breaux, D. A. Hillman, C. M. Neal and M. F. Jarrold, *J. Phys. Chem. A*, 2005, **109**, 8755–8759.
- 78 L. F. Cui, X. Huang, L. M. Wang, D. Y. Zubarev, A. I. Boldyrev, J. Li and L. S. Wang, *J. Am. Chem. Soc.*, 2006, **128**, 8390–8391.
- 79 S. Cantrill, *Nat. Nanotechnol.*, 2006, DOI: [10.1038/nnano.2006.11](https://doi.org/10.1038/nnano.2006.11).
- 80 L. F. Cui, L. M. Wang and L. S. Wang, *J. Chem. Phys.*, 2007, **126**, 064505.
- 81 L.-F. Cui, X. Huang, L.-M. Wang, J. Li and L.-S. Wang, *Angew. Chem.*, 2007, **119**, 756–759.
- 82 V. Kumar and Y. Kawazoe, *Appl. Phys. Lett.*, 2003, **83**, 2677–2679.
- 83 U. Rohrmann and R. Schäfer, *Phys. Rev. Lett.*, 2013, **111**, 133401.
- 84 T. M. Fuchs and R. Schäfer, *Phys. Rev. A*, 2018, **98**, 063411.
- 85 S. Bobev and S. C. Sevov, *Inorg. Chem.*, 2001, **40**, 5361–5364.
- 86 B. Kesanli, J. Fettinger and B. Eichhorn, *Chem. – Eur. J.*, 2001, **7**, 5277–5285.
- 87 M. Brynda, R. Herber, P. B. Hitchcock, M. F. Lappert, I. Nowik, P. P. Power, A. V. Protchenko, A. Růžička and J. Steiner, *Angew. Chem., Int. Ed.*, 2006, **45**, 4333–4337.
- 88 S. J. Kim, S. D. Hoffman and T. F. Fässler, *Angew. Chem., Int. Ed.*, 2007, **46**, 3144–3148.
- 89 G. Prabusankar, A. Kempter, C. Gemel, M. K. Schöter and R. A. Fischer, *Angew. Chem., Int. Ed.*, 2008, **47**, 7234–7237.
- 90 C. Schrenk, I. Schellenberg, R. Pöttgen and A. Schnepf, *Dalton Trans.*, 2010, **39**, 1872–1876.
- 91 J. Q. Wang, S. Stegmaier, B. Wahl and T. F. Fässler, *Chem. – Eur. J.*, 2010, **16**, 1793–1798.
- 92 S. Stegmaier and T. F. Fässler, *J. Am. Chem. Soc.*, 2011, **133**, 19758–19768.
- 93 C. Schrenk, F. Winter, R. Pöttgen and A. Schnepf, *Chem. – Eur. J.*, 2015, **21**, 2992–2997.
- 94 J. Wiederkehr, C. Wölper and S. Schulz, *Chem. Commun.*, 2016, **52**, 12282–12285.
- 95 A. P. Purdy, R. J. Butcher, J. P. Yesinowski, S. A. Fischer, D. Gunlycke and B. L. Chaloux, *Inorg. Chem.*, 2018, **57**, 4921.
- 96 M. Binder, C. Schrenk and A. Schnepf, *Chem. Commun.*, 2019, **55**, 12148–12151.
- 97 L. Qiao, C. Zhang, C. C. Shu, H. W. Morgan, J. E. McGrady and Z. M. Sun, *J. Am. Chem. Soc.*, 2020, **142**, 13288–13293.
- 98 J. E. McGrady, F. Weigend and S. Dehnen, *Chem. Soc. Rev.*, 2022, **51**, 628–649.
- 99 T. Bachelors and R. Schäfer, *Chem. Phys. Lett.*, 1999, **300**, 177–182.
- 100 T. Bachelors, R. Schäfer and H. J. Güntherodt, *Phys. Rev. Lett.*, 2000, **84**, 4890–4893.
- 101 S. Schäfer, M. Mehring, R. Schäfer and P. Schwerdtfeger, *Phys. Rev. A: At., Mol., Opt. Phys.*, 2007, **76**, 052515.
- 102 S. Heiles and R. Schäfer, *Dielectric Properties of Isolated Clusters: Beam Deflection Studies*, Springer, Heidelberg, 2014.
- 103 A. Shayeghi, R. L. Johnston and R. Schäfer, *Phys. Chem. Chem. Phys.*, 2013, **15**, 19715–19723.
- 104 M. Jäger, A. Shayeghi, V. Klippenstein, R. L. Johnston and R. Schäfer, *J. Chem. Phys.*, 2018, **149**, 244308.
- 105 A. Lehr, M. Jäger and R. Schäfer, *J. Phys. Chem. C*, 2020, **124**, 1070–1076.
- 106 L. P. Maguire, S. Szilagyí and R. E. Scholten, *Rev. Sci. Instrum.*, 2004, **75**, 3077–3079.
- 107 L. P. Maguire, S. Szilagyí and R. E. Scholten, *Rev. Sci. Instrum.*, 2007, **78**, 019903.
- 108 T. M. Fuchs, F. Rivic and R. Schäfer, *Phys. Rev. A*, 2021, **104**, 012820.
- 109 J. M. Kellogg, I. I. Rabi, N. F. Ramsey and J. R. Zacharias, *Phys. Rev.*, 1939, **56**, 728–743.
- 110 U. Ray, M. F. Jarrold, J. E. Bower and J. S. Kraus, *J. Chem. Phys.*, 1989, **91**, 2912–2921.
- 111 M. F. Jarrold and E. C. Honea, *J. Phys. Chem.*, 1991, **95**, 9181–9185.
- 112 M. Jäger, R. Schäfer and R. L. Johnston, *Nanoscale*, 2019, **11**, 9042–9052.
- 113 P. Giannozzi, S. Baroni, N. Bonini, M. Calandra, R. Car, C. Cavazzoni, D. Ceresoli, G. L. Chiarotti, M. Cococcioni, I. Dabo, A. Dal Corso, S. De Gironcoli, S. Fabris, G. Fratesi, R. Gebauer, U. Gerstmann, C. Gougoussis, A. Kokalj, M. Lazzeri, L. Martin-Samos, N. Marzari, F. Mauri, R. Mazzarello, S. Paolini, A. Pasquarello, L. Paulatto, C. Sbraccia, S. Scandolo, G. Sclauzero, A. P. Seitsonen, A. Smogunov, P. Umari and R. M. Wentzcovitch, *J. Phys.: Condens. Matter*, 2009, **21**, 395502.
- 114 P. Giannozzi, O. Andreussi, T. Brumme, O. Bunau, M. B. Nardelli, M. Calandra, R. Car, C. Cavazzoni, D. Ceresoli, M. Cococcioni, N. Colonna, I. Carnimeo, A. Dal Corso, S. de Gironcoli, P. Delugas, R. A. Distasio Jr., A. Ferretti, A. Floris, G. Fratesi, G. Fugallo, R. Gebauer, U. Gerstmann, F. Giustino, T. Gorni, J. Jia, M. Kawamura, H.-Y. Ko, A. Kokalj, E. Küçükbenli, M. Lazzeri, M. Marsili, N. Marzari, F. Mauri, N. L. Nguyen, H.-V. Nguyen, A. Otero-de-la Roza, L. Paulatto, S. Poncé, D. Rocca, R. Sabatini, B. Santra, M. Schlipf, A. P. Seitsonen, A. Smogunov, I. Timrov, T. Thonhauser, P. Umari, N. Vast, X. Wu and S. Baroni, *J. Phys.: Condens. Matter*, 2017, **29**, 465901.
- 115 J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**, 3865–3868.
- 116 J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1997, **78**, 1396.
- 117 D. Vanderbilt, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1985, **32**, 8412–8415.
- 118 A. M. Rappe, K. M. Rabe, E. Kaxiras and J. D. Joannopoulos, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1990, **41**, 1227–1230.
- 119 M. Methfessel and A. T. Paxton, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1989, **40**, 3616–3621.
- 120 S. Louie, S. Froyen and L. Cohen, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1982, **26**, 1738–1742.
- 121 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato,



- A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. J. A. Montgomery, J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman and D. J. Fox, *Gaussian 16, Revision C.01*, 2016.
- 122 M. Valiev, E. J. Bylaska, N. Govind, K. Kowalski, T. P. Straatsma, H. J. Van Dam, D. Wang, J. Nieplocha, E. Apra, T. L. Windus and W. A. De Jong, *Comp. Phys. Commun.*, 2010, **181**, 1477–1489.
- 123 E. Aprà, E. J. Bylaska, W. A. De Jong, N. Govind, K. Kowalski, T. P. Straatsma, M. Valiev, H. J. Van Dam, Y. Alexeev, J. Anchell, V. Anisimov, F. W. Aquino, R. Attafynn, J. Autschbach, N. P. Bauman, J. C. Becca, D. E. Bernholdt, K. Bhaskaran-Nair, S. Bogatko, P. Borowski, J. Boschen, J. Brabec, A. Bruner, E. Cauët, Y. Chen, G. N. Chuev, C. J. Cramer, J. Daily, M. J. Deegan, T. H. Dunning, M. Dupuis, K. G. Dyall, G. I. Fann, S. A. Fischer, A. Fonari, H. Früchtel, L. Gagliardi, J. Garza, N. Gawande, S. Ghosh, K. Glaesemann, A. W. Götz, J. Hammond, V. Helms, E. D. Hermes, K. Hirao, S. Hirata, M. Jacquelin, L. Jensen, B. G. Johnson, H. Jónsson, R. A. Kendall, M. Klemm, R. Kobayashi, V. Konkov, S. Krishnamoorthy, M. Krishnan, Z. Lin, R. D. Lins, R. J. Littlefield, A. J. Logsdail, K. Lopata, W. Ma, A. V. Marenich, J. Martin Del Campo, D. Mejia-Rodriguez, J. E. Moore, J. M. Mullin, T. Nakajima, D. R. Nascimento, J. A. Nichols, P. J. Nichols, J. Nieplocha, A. Otero-De-La-Roza, B. Palmer, A. Panyala, T. Pirojsirikul, B. Peng, R. Peverati, J. Pittner, L. Pollack, R. M. Richard, P. Sadayappan, G. C. Schatz, W. A. Shelton, D. W. Silverstein, D. M. Smith, T. A. Soares, D. Song, M. Swart, H. L. Taylor, G. S. Thomas, V. Tipparaju, D. G. Truhlar, K. Tsemekhman, T. Van Voorhis, A. Vázquez-Mayagoitia, P. Verma, O. Villa, A. Vishnu, K. D. Vogiatzis, D. Wang, J. H. Weare, M. J. Williamson, T. L. Windus, K. Woliński, A. T. Wong, Q. Wu, C. Yang, Q. Yu, M. Zacharias, Z. Zhang, Y. Zhao and R. J. Harrison, *J. Chem. Phys.*, 2020, **152**, 184102.
- 124 J. P. Perdew, M. Ernzerhof and K. Burke, *J. Chem. Phys.*, 1996, **105**, 9982–9985.
- 125 B. Metz, H. Stoll and M. Dolg, *J. Chem. Phys.*, 2000, **113**, 2563–2569.
- 126 K. A. Peterson, *J. Chem. Phys.*, 2003, **119**, 11099–11112.
- 127 M. A. Rohrdanz, K. M. Martins and J. M. Herbert, *J. Chem. Phys.*, 2009, **130**, 054112.
- 128 F. Weigend and R. Ahlrichs, *Phys. Chem. Chem. Phys.*, 2005, **7**, 3297–3305.
- 129 S. Heiles, R. L. Johnston and R. Schäfer, *J. Phys. Chem. A*, 2012, **116**, 7756–7764.
- 130 M. Gleditzsch, M. Jäger, L. F. Pašteka, A. Shayeghi and R. Schäfer, *Phys. Chem. Chem. Phys.*, 2019, **21**, 24478–24488.
- 131 M. Gleditzsch, T. M. Fuchs and R. Schäfer, *J. Phys. Chem. A*, 2019, **123**, 1434–1444.
- 132 M. Gleditzsch, L. F. Pašteka, D. A. Götz, A. Shayeghi, R. L. Johnston and R. Schäfer, *Nanoscale*, 2019, **11**, 12878–12888.
- 133 C. Riplinger, B. Sandhoefer, A. Hansen and F. Neese, *J. Chem. Phys.*, 2013, **139**, 134101.
- 134 D. G. Liakos, M. Sparta, M. K. Kesharwani, J. M. Martin and F. Neese, *J. Chem. Theory Comput.*, 2015, **11**, 1525–1539.
- 135 D. G. Liakos and F. Neese, *J. Chem. Theory Comput.*, 2015, **11**, 4054–4063.
- 136 C. Riplinger, P. Pinski, U. Becker, E. F. Valeev and F. Neese, *J. Chem. Phys.*, 2016, **144**, 024109.
- 137 F. Neese, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2012, **2**, 73–78.
- 138 F. Neese, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2018, **8**, 4–9.
- 139 F. Neese, F. Wennmohs, U. Becker and C. Riplinger, *J. Chem. Phys.*, 2020, **152**, 224108.
- 140 D. G. Truhlar, *Chem. Phys. Lett.*, 1998, **294**, 45–48.
- 141 P. Dugourd, R. Antoine, M. Abd El Rahim, D. Rayane, M. Broyer and F. Calvo, *Chem. Phys. Lett.*, 2006, **423**, 13–16.
- 142 S. Heiles, S. Schäfer and R. Schäfer, *J. Chem. Phys.*, 2011, **135**, 034303.
- 143 R. L. Martin, *J. Chem. Phys.*, 2003, **118**, 4775–4777.
- 144 A. Heilmann, M. Quinten and J. Werner, *Eur. Phys. J. B*, 1998, **3**, 455–461.
- 145 V. G. Grigoryan, M. Springborg, H. Minassian and A. Melikyan, *Comput. Theor. Chem.*, 2013, **1021**, 197–205.
- 146 O. K. Rice and H. C. Ramsperger, *J. Am. Chem. Soc.*, 1927, **49**, 1617–1629.
- 147 O. K. Rice and H. C. Ramsperger, *J. Am. Chem. Soc.*, 1928, **50**, 617–620.
- 148 R. A. Marcus and O. K. Rice, *J. Phys. Colloid Chem.*, 1951, **55**, 894–908.
- 149 R. A. Marcus, *J. Chem. Phys.*, 1952, **20**, 359–364.
- 150 R. G. Gilbert and S. C. Smith, *Theory of Unimolecular and Recombination Reactions*, Blackwell Scientific Publications, Oxford, 1990.
- 151 T. Beyer and D. F. Swinehart, *Commun. ACM*, 1973, **16**, 379.
- 152 M. F. Jarrold and K. M. Creegan, *Int. J. Mass Spectrom. Ion Process.*, 1990, **102**, 161–181.
- 153 Y. Shi, V. A. Spasov and K. M. Ervin, *J. Chem. Phys.*, 1999, **111**, 938–949.
- 154 D. A. Götz, A. Shayeghi, R. L. Johnston, P. Schwerdtfeger and R. Schäfer, *Nanoscale*, 2016, **8**, 11153–11160.
- 155 K. D. Rinnen and M. L. Mandich, *Phys. Rev. Lett.*, 1992, **69**, 1823–1826.
- 156 K. D. Rinnen and M. L. Mandich, *Z. Phys. D*, 1993, **26**, 147–149.



- 157 E. C. Honea, J. S. Kraus, J. E. Bower and M. F. Jarrold, *Z. Phys. D*, 1993, **26**, 141–143.
- 158 A. Rubio, J. Alonso, X. Blase, L. Balbás and S. Louie, *Phys. Rev. Lett.*, 1996, **77**, 247–250.
- 159 Y. Z. Lan, Y. L. Feng, Y. H. Wen and B. T. Teng, *J. Mol. Struct.*, 2008, **854**, 63–69.
- 160 L.-Z. Zhao, W.-C. Lu, W. Qin, Q.-J. Zang, K. M. Ho and C. Z. Wang, *J. Phys. Chem. A*, 2017, **121**, 6388–6397.
- 161 G. Mie, *Ann. Phys.*, 1908, **330**, 377–445.
- 162 R. Gans, *Ann. Phys.*, 1912, **342**, 881–900.
- 163 H. E. Johnson and C. M. Aikens, *J. Phys. Chem. A*, 2009, **113**, 4445–4450.
- 164 H. Yu, J. Li, R. A. Loomis, L. W. Wang and W. E. Buhro, *Nat. Mater.*, 2003, **2**, 517–520.
- 165 H. Yu, J. Li, R. A. Loomis, P. C. Gibbons, L. W. Wang and W. E. Buhro, *J. Am. Chem. Soc.*, 2003, **125**, 16168–16169.

