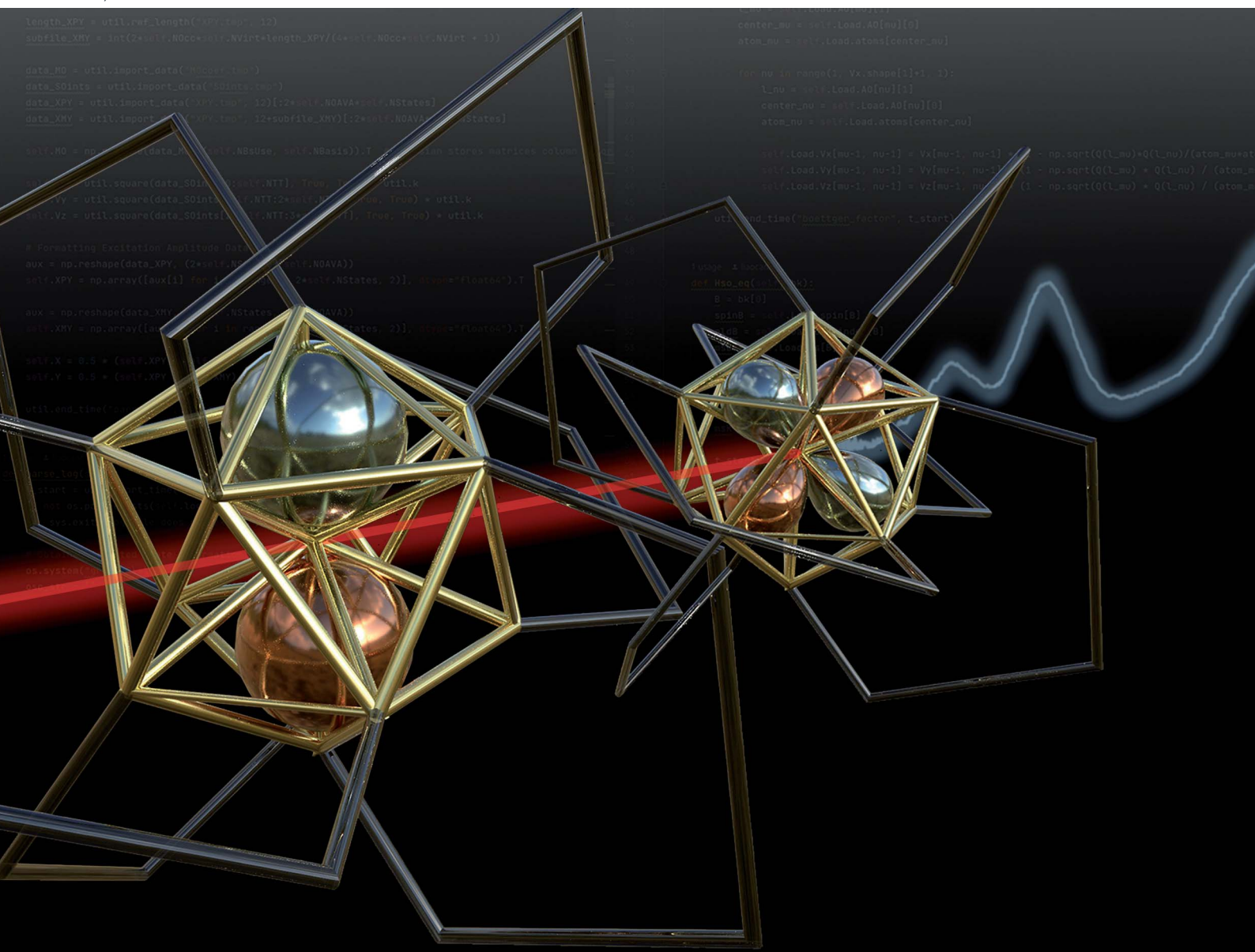


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**EDGE ARTICLE**

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# Manifestation of the interplay between spin–orbit and Jahn–Teller effects in Au<sub>25</sub> superatom UV-Vis fingerprint spectra†

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Atomically precise nanoclusters play an important role in nanoscale catalysis, photonics, and quantum information science. Their nanochemical properties arise from their unique superatomic electronic structures. As the flagship of atomically precise nanochemistry, the Au<sub>25</sub>(SR)<sub>18</sub> nanocluster exhibits tunable spectroscopic signatures that are sensitive to the oxidation state. This work aims to unravel the physical underpinnings of the spectral progression of Au<sub>25</sub>(SR)<sub>18</sub> nanocluster using variational relativistic time-dependent density functional theory. The investigation will focus on the effects of superatomic spin–orbit coupling, its interplay with Jahn–Teller distortion, and their manifestations in the absorption spectra of Au<sub>25</sub>(SR)<sub>18</sub> nanoclusters of different oxidation states.

The Au<sub>25</sub>(SR)<sub>18</sub> nanocluster is the flagship of atomically precise nanochemistry.<sup>1</sup> Over one thousand papers have been published about it and its derivatives since its identification in 2005 (ref. 2) and structure determination in 2008.<sup>1,3–7</sup> However, the influence of spin–orbit coupling (SOC) over different oxidation states of the Au<sub>25</sub>(SR)<sub>18</sub> nanocluster has not been fully understood, despite it being the most studied cluster in the field and recent attempts.<sup>8–11</sup> SOC plays an important role in the electronic structure and properties of molecules and materials, especially for heavier elements such as Au. The inclusion of SOC in *ab initio* calculations of ground and excited state nanomaterials has not been routinely pursued due to the high computational cost. This challenge has prevented a full understanding of some of the most studied nanosystems.

Fig. 1 shows the experimental UV-Vis absorption spectra of Au<sub>25</sub>(SR)<sub>18</sub> nanoclusters (abbreviated as Au<sub>25</sub>) at three different oxidation states. The differences among them can clearly be seen. The low energy region from 1.2 eV to 2.0 eV, known as the fingerprint band, corresponds mainly to the transitions between frontier orbitals: the double peak in Au<sub>25</sub><sup>−</sup> changes to an asymmetric band in neutral Au<sub>25</sub> with further blue-shifts in Au<sub>25</sub><sup>+</sup>.<sup>12–15</sup> The frontier orbitals of Au<sub>25</sub> can be well-understood in the absence of SOC by the superatomic complex model:<sup>16</sup> Au<sub>25</sub><sup>−</sup> being an 8-electron system with a superatomic electron

configuration of (1S)<sup>2</sup>(1P)<sup>6</sup>, with 1D orbitals being the LUMO. Although the double peaks of Au<sub>25</sub><sup>−</sup> were explained by the SOC splitting of the 1P orbitals based on two-component time-dependent density functional theory (2C-TDDFT),<sup>8</sup> the role of SOC in the neutral and cationic spectra remains unexplored.

Another consideration is the interplay between geometry and electronic structure as a result of SOC. Jahn–Teller distortion plays a crucial role in the electronic structure of metal nanoclusters including ligand-protected metal clusters.<sup>15,17,18</sup> It is well-known that filling degenerate or near-degenerate orbitals with unpaired electrons is met by geometric distortions that lower the nanocluster symmetry and break orbital degeneracies. Metal nanoclusters tend to not follow Hund's rule since the energy lowered by Jahn–Teller distortion can be greater than that by Hund's spin–spin coupling. This perspective is

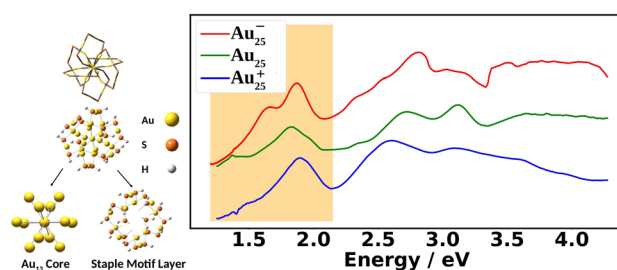


Fig. 1 On the left, ball-and-stick and wire frame models of Au<sub>25</sub>(SR)<sub>18</sub> are shown. The nanocluster can be separated into two regions: the Au<sub>13</sub> core and the staple motif layer. In our computational model, we use R = H to reduce computational cost. On the right, the experimental UV-Vis absorption spectra of Au<sub>25</sub>(SPET)<sub>18</sub> (SPET = 2-phenylethanethiolate: SCH<sub>2</sub>CH<sub>2</sub>Ph) at the −1, 0, and +1 charge states recorded at 78 K is shown. The shaded region is the “fingerprint” band, which is the focus of this work.

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paramount to understanding the evolution of the UV-Vis fingerprint band with respect to charge. As the nanocluster oxidizes from anionic ( $-1$ ) to neutral ( $0$ ) and cationic ( $+1$ ) oxidation states, the superatomic orbital electron configuration changes from  $1P^6$  to  $1P^5$  and  $1P^4$ , respectively, presenting a clear opportunity for Jahn–Teller distortion to act. Indeed, the Jahn–Teller effects have been shown to be present in  $Au_{25}$  which can be further manipulated to tune its redox properties.<sup>15,19</sup>

The goal of the present work is, therefore, to fully understand the role of SOC in the fingerprint absorption band at 1.2 eV to 2.0 eV for the  $-1$ ,  $0$ , and  $+1$  oxidation states of the  $Au_{25}(SR)_{18}$  nanocluster and to correlate the SOC-induced electron structure modulation with Jahn–Teller distortion.  $Au_{25}(SH)_{18}$  was used as a model in the study to reduce computational cost since it has been shown that optical properties in this energy region are insensitive to ligand choice.<sup>12,20,21</sup> All calculations were performed using a developmental version of the Gaussian electronic structure package.<sup>22</sup> Two-component time-dependent density functional theory calculations were performed using the PBE0,<sup>23,24</sup> functional with the relativistic CRENL<sup>25–29</sup> effective core potential (ECP) including SOC and its complementary basis set for Au and S atoms. See ESI† for more details.

Fig. 2 compares the simulated spectra with the experiment for the three different oxidation states of  $Au_{25}$ . The low energy satellite peak at 1.64 eV was labeled as  $\alpha_1$  and the main peak at 1.91 eV was labeled as  $\alpha_2$  in Fig. 2a for the anionic cluster. Upon oxidation of the anion to the neutral nanocluster, the satellite peak lowered in intensity and was red-shifted to 1.43 eV, labeled as  $\beta_1$  in Fig. 2b. Broadening of the main peak was observed along with a slight red-shift to 1.82 eV, labeled as  $\beta_3$  in Fig. 2b.  $\beta_3$  exhibited a low energy tail with significant intensity at 1.58 eV that connects  $\beta_1$  and  $\beta_3$ , labeled as  $\beta_2$  in Fig. 2b.  $\beta_3$  also exhibited a high energy shoulder, labeled  $\beta_4$ . Further oxidation to the cationic state resulted in the disappearance of the low energy satellite peak along with a blue-shift of the main peak to 1.89 eV, labeled  $\gamma_2$ . The low-energy tail remains at 1.70 eV and is labeled  $\gamma_1$ . The high energy shoulder of the main peak is no longer present in the experimental spectrum, but remains in the simulated spectrum. Overall, one can see an excellent agreement between the simulations and the experimental spectra for the three different charge states.

We next correlate the change in the optical absorption spectra to the change in the geometry of the  $Au_{25}$  nanocluster as the charge is varied, with a focus on SOC electronic structure. We note that Jahn–Teller distortion in  $Au_{25}$  nanoclusters has been correlated to experimental optical absorption and voltammograms.<sup>15,18,31</sup> Fig. 3 shows the change in the nanocluster  $Au_{13}$  core geometry and diagonal lengths for each oxidation state. The anion core was nearly a perfect icosahedron. The slight imperfections left the core with a  $T_h$  symmetry rather than the  $I_h$  symmetry expected of a perfect icosahedron. Oxidation of the anion to the neutral nanocluster resulted in a distortion of the core lowering its symmetry to  $C_i$ . The distortion appeared as a tilting of the 10 equatorial triangular faces of the icosahedron caused by the increase in bond angles between the purple, blue, and orange diagonals. In addition, every diagonal decreased in length leading to an overall shrinking of the core. Much greater

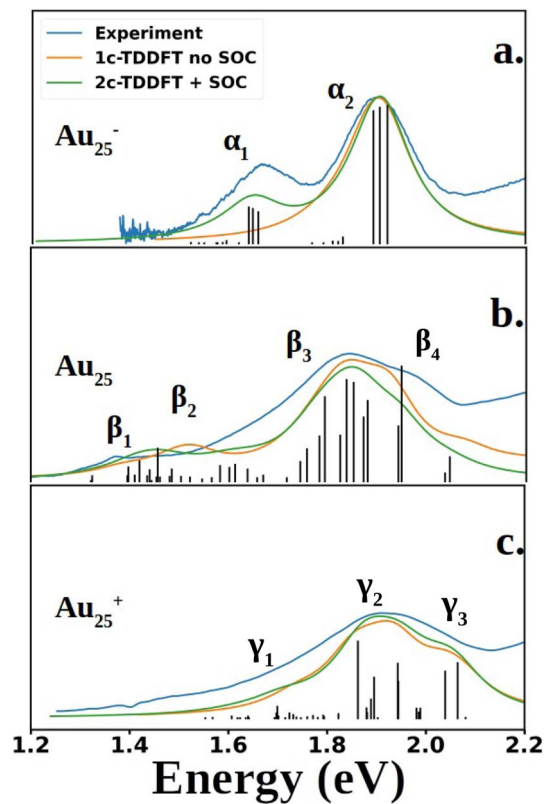


Fig. 2 Experimental and computed spectra of  $Au_{25}(SR)_{18}$ . The excited states obtained from 2C-TDDFT are plotted as black lines. Spectra were generated by applying Lorentzian broadening to the excited states with a half-width at half max of 0.07 eV. The anion spectrum (a) is compared to the experimental spectrum of  $Au_{25}(C_6H_{13})_{18}^-$  ( $SCH_6H_{13}$ : 1-hexanethiolate), reproduced from data obtained from Ramakrishna *et al.*<sup>30</sup> The neutral (b) and cation (c) spectra are compared to experimental spectra of  $Au_{25}(SPET)_{18}$  and  $Au_{25}(SPET)_{18}^+$  respectively. All spectra were recorded at 78 K. The computed neutral spectrum was red-shifted by 0.11 eV to align with the experimental spectrum.

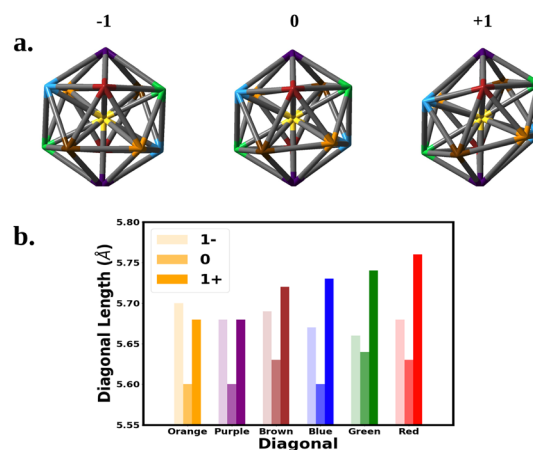


Fig. 3 The structure of the  $Au_{13}$  core at the  $-1$ ,  $0$ , and  $+1$  oxidation states are shown above. The vertices represent Au atoms. Atoms of the same color share a diagonal that runs through the center. The diagonal lengths are plotted for each oxidation state in the bottom chart.



distortion was observed when oxidized to the cation. The symmetry of the cation core remained  $C_i$  but the symmetry deviated further from icosahedron. The equatorial tilting of the icosahedron became much more pronounced as the bond angles between the purple, blue, and orange diagonals further increased. The length of each diagonal also increased with most diagonals surpassing their anionic lengths.

Molecular orbital analysis is necessary in the characterization of the absorbance spectra. Fig. 4 shows the anionic MO diagram calculated with and without SOC along with visualization of the superatomic orbitals. The highest occupied and unoccupied orbitals are the 1P and 1D, respectively. The 1P orbitals generally have the same appearance but lie on different axes. The 1D superatomic orbitals were more diverse in their appearance including having a  $d_{z^2}$  or “four-leaf clover” center. Because superatomic orbitals can be identified with molecular orbitals, molecular symmetry dictates how atomic spin-orbit coupling collectively manifests in superatomic orbitals.

The anion superatomic orbitals split in the same manner as atomic orbitals in a  $T_h$  field as shown in Fig. 4. Double group symmetry is used in the discussion of molecular orbitals under SOC. Irreducible representations (irrep) of the double symmetry is denoted with a tilde. In the discussion of orbital degeneracies, we take the perspective of spin-orbitals (e.g., a non-relativistic hydrogen 2p manifold is a six-fold degeneracy). Without SOC, the 1D superatomic orbitals split into  $t_g$  and  $e_g$  while the 1P superatomic orbitals remain degenerate transforming as  $t_u$ . With SOC, the  $t_g$  manifold splits into a doubly degenerate  $\tilde{e}_{1/2,g}$

and a four-fold degeneracy  $\tilde{f}_{3/2,g}$  while the  $e_g$  manifold remains as a four-fold degeneracy (re-designated as  $\tilde{f}_{3/2,g}$  under double group theory). The  $t_u$  manifold splits into a four-fold degeneracy  $\tilde{f}_{3/2,u}$  and a doubly degenerate  $\tilde{e}_{1/2,u}$ . The relativistic field splitting was consistent with the predictions of double group theory. Below the 1P superatomic orbitals was a dense region with orbitals delocalized onto the staple motif layer. It was observed that these orbitals do play a key role in the neutral and cationic spectra within the fingerprint region.

The MO diagram of the neutral and cationic nanoclusters with SOC are shown in Fig. 5 (see ESI† for MO diagrams without SOC). The driver of Jahn–Teller distortion was the removal of electrons from the former 1P  $\tilde{f}_{3/2,u}$  manifold that breaks the spherical symmetry of the 1P shell. The nanocluster symmetry for both oxidation states lowered to  $C_i$  to break the partially filled  $\tilde{f}_{3/2,u}$  degeneracy. The neutral cluster experienced minor distortion since the destabilized 1P was still singly occupied. This was reflected in the reminiscence of the large energy gap between the former 1D  $\tilde{f}_{3/2,g}$  and  $\tilde{e}_{1/2,g}$  manifolds, resembling  $T_h$  orbital splitting. The distortion was much greater for the cation since the depopulated 1P can further destabilize without raising overall energy. The 1D superatomic orbitals further separate closing the large energy gap. In both oxidation states, the 1P degeneracy was broken.

Shown in Fig. 2 are the experimental and simulated spectra for the three oxidation states of  $Au_{25}$ . The most straightforward spectrum is the anionic spectrum shown in Fig. 2a. The anion

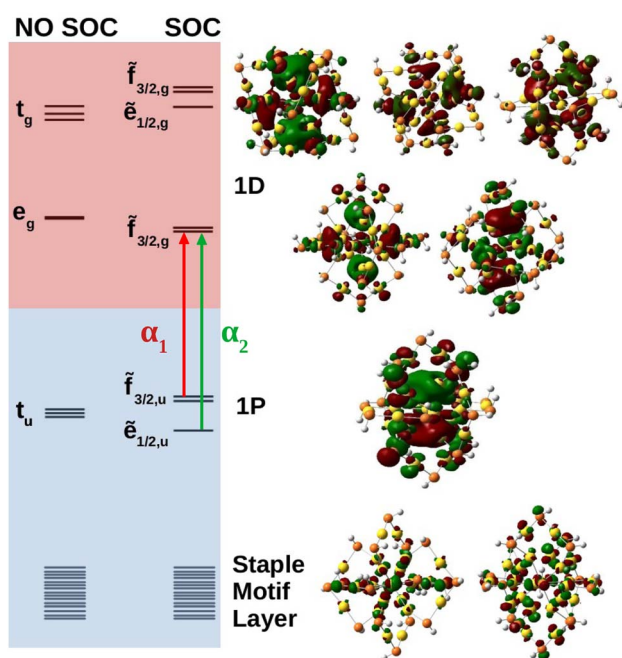


Fig. 4 MO diagram of  $Au_{25}(SH)_{18}^-$  with and without SOC, where each line is a Kramers pair. Blue and red shaded orbitals denote occupied and unoccupied orbitals, respectively. Below the superatomic orbitals exists a high density of states region where electron density is delocalized onto the staple motif layer. Images of the orbitals are shown on the right.

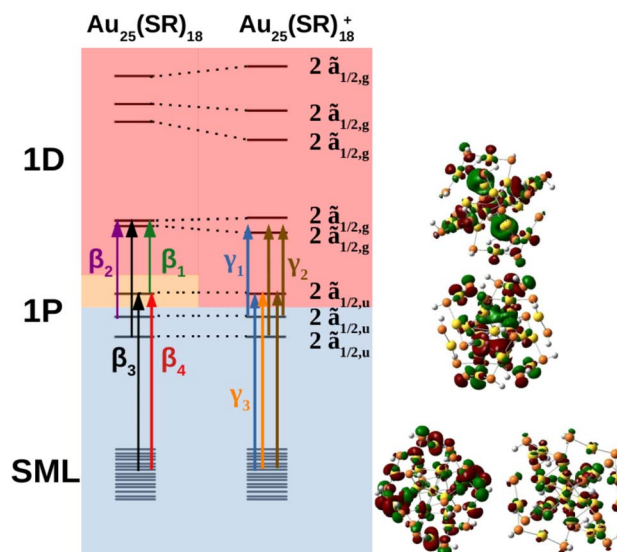


Fig. 5 MO diagram depicting the Jahn–Teller distortion from the neutral  $Au_{25}(SR)_{18}$  to  $Au_{25}(SR)_{18}^+$  where each line is a Kramers pair. Orbital transitions responsible for features  $\beta_1$ ,  $\beta_2$ ,  $\beta_3$ , and  $\beta_4$  on the neutral spectrum; and  $\gamma_1$ ,  $\gamma_2$ , and  $\gamma_3$  on the cationic spectrum, are shown. Blue, orange, and red shaded orbitals are doubly-occupied, singly-occupied, and unoccupied respectively. The double group irrep of the orbitals are shown on the right and are the same for both oxidation states. The “2” in front of the irrep indicates that the Kramers pair is a pair of spin-orbitals that transform as the given irrep (e.g.,  $2\tilde{a}_{1/2,u}$  indicates that two spin-orbitals that transform as  $\tilde{a}_{1/2,u}$  make up the Kramers pair).



**Table 1** Percentage of triplet and singlet eigenstate contribution to excited states responsible for  $\alpha_1$  and  $\alpha_2$  in the anion spectrum. Excited states are labeled by increasing energy within a given peak

Peak	Excited state	% triplet	% singlet
$\alpha_1$	1	74.91%	25.09%
	2	67.22%	32.78%
	3	76.65%	23.35%
$\alpha_2$	1	34.03%	65.97%
	2	26.37%	73.63%
	3	37.74%	72.26%

spectrum consisted of two peaks,  $\alpha_1$  and  $\alpha_2$ . The orbital excitations responsible for  $\alpha_1$  and  $\alpha_2$  were  $1P \tilde{e}_{1/2,u} \rightarrow 1D \tilde{f}_{3/2,g}$  and  $1P \tilde{f}_{3/2,u} \rightarrow 1D \tilde{f}_{3/2,g}$ , respectively. From the density of state point of view,  $\alpha_2$ , arising from four-fold degenerate  $\tilde{f}_{3/2,u}$  orbitals, should have a higher intensity than  $\alpha_1$ , which arises from two-fold degenerate  $\tilde{e}_{1/2,u}$  orbitals. However, this is not the case in experiment nor our calculations. Spin projection of 2C-TDDFT results shows that the excited states responsible for  $\alpha_2$  are comprised mostly of singlet character whereas those responsible for  $\alpha_1$  are comprised mostly of triplet character, as shown in Table 1. Given that the ground state is a closed-shell singlet, the discrepancy in peak intensities is clearly due to the spin selection rule ( $\Delta S = 0$ ).

The neutral nanocluster spectrum is shown in Fig. 2b. Many more excited states were responsible for the neutral spectrum than for the anionic spectrum. The features  $\beta_1$  and  $\beta_2$  were direct consequences of Jahn–Teller distortion. Recall that Jahn–Teller distortion breaks the  $\tilde{f}_{3/2,u}$  degeneracy into a stabilized doubly-occupied 1P and a destabilized singly-occupied 1P. Orbital excitations from the destabilized singly-occupied 1P to the lowest two 1D gave rise to  $\beta_1$ . Orbital excitations from the stabilized doubly-occupied 1P to the lowest two 1D gave rise to  $\beta_2$ . Along with excitations between superatomic orbitals, the vacancy in 1P opened opportunities for the low energy orbitals delocalized onto the staple motif layer (SML) to excite into the core-localized 1P, suggesting possible ligand–metal charge transfer (LMCT) in the fingerprint region. Mixing of these orbital excitations were pervasive in the excited states throughout the spectrum. Along with orbital excitations from the lowest 1P to the lowest two 1D, SML  $\rightarrow$  1P LMCT excitations also contributed to electronic transitions that form  $\beta_3$ . The rise of SML  $\rightarrow$  1P LMCT excitations with considerable oscillator strength also led to the broadening of  $\beta_3$  and the rise of  $\beta_4$ . Because ligands in the theoretical model differ from experiment ( $R = SH$  as opposed to SPET), it is reasonable for the theoretical spectrum to disagree with experiment at  $\beta_4$ .

The simulated cation spectrum is shown in Fig. 2c. With the depletion of the highest 1P,  $\beta_1$  was not present in the cationic spectrum, resulting in a single broad peak, labeled  $\gamma_2$ , and a low energy tail, labeled  $\gamma_1$ . Excitations between the middle 1P and the two lowest 1D along with SML  $\rightarrow$  1P LMCT excitations were observed in  $\gamma_1$ . The high intensity excited state at 1.82 eV that constitutes  $\gamma_2$  also consists mainly of excitations between the middle 1P and the two lowest 1D. The rest of the excited states that constitute  $\gamma_2$  were from orbital excitations between the lowest 1P

and the two lowest 1D along with SML  $\rightarrow$  1P LMCT. The blue-shift of the main peak compared to the neutral spectrum arose from further destabilization of the depleted 1P. Such destabilization increased the energy of excited states involving SML  $\rightarrow$  1P LMCT, leading to a blue-shift of the main peak. The lack of high intensity excited states at  $\gamma_3$  due to a minimal ligand caused the theoretical spectrum to exhibit a high energy shoulder instead of a single broad peak encompassing both  $\gamma_2$  and  $\gamma_3$ , as seen in experiment.

## Conclusion

Herein, relativistic 2C-TDDFT with a variational treatment of SOC was employed to investigate the physical underpinnings behind the spectral progression of the Au<sub>25</sub> nanocluster UV-Vis absorption spectrum as it oxidizes from  $-1$  to  $0$  and  $+1$  oxidation states. In addition to reproducing absorption spectra with high accuracy, the effect of the interplay between spin–orbit and Jahn–Teller distortion was carefully analyzed. To our knowledge, this was the first study to provide a thorough picture of the Au<sub>25</sub> nanocluster relating SOC, Jahn–Teller distortion, and spectroscopic characteristics.

## Data availability

Data are available from the corresponding authors upon request.

## Author contributions

X. Li, D. Jiang, and M. Zhu conceived the project; C. Liao performed the computational study and data analysis; M. Zhu conducted experiments; X. Li was responsible for acquisition of funding; all authors contributed to the development of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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