Medium-sized Au$_{40}(SR)_{24}$ and Au$_{52}(SR)_{32}$ nanoclusters with distinct gold-kernel structures and spectroscopic features

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Medium-sized Au$_{40}$(SR)$_{24}$ and Au$_{52}$(SR)$_{32}$ nanoclusters with distinct gold-kernel structures and spectroscopic features†

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We have analyzed the structures of two medium-sized thiolate-protected gold nanoparticles (RS-AuNPs) Au$_{40}$(SR)$_{24}$ and Au$_{52}$(SR)$_{32}$ and identified the distinct structural features in their Au kernels [Sci. Adv., 2015, 1, e1500425]. We find that both Au kernels of the Au$_{40}$(SR)$_{24}$ and Au$_{52}$(SR)$_{32}$ nanoclusters can be classified as interpenetrating cuboctahedra. Simulated X-ray diffraction patterns of the RS-AuNPs with the cuboctahedral kernel are collected and then compared with the X-ray diffraction patterns of the RS-AuNPs of two other prevailing Au-kernels identified from previous experiments, namely the Ino-decahedral kernel and icosahedral kernel. The distinct X-ray diffraction patterns of RS-AuNPs with the three different types of Au-kernels can be utilized as signature features for future studies of structures of RS-AuNPs. Moreover, the simulated UV/Vis absorption spectra and Kohn–Sham orbital energy-level diagrams are obtained for the Au$_{40}$(SR)$_{24}$ and Au$_{52}$(SR)$_{32}$, on the basis of time-dependent density functional theory computation. The extrapolated optical band-edges of Au$_{40}$(SR)$_{24}$ and Au$_{52}$(SR)$_{32}$ are 1.1 eV and 1.25 eV, respectively. The feature peaks in the UV/Vis absorption spectra of the two clusters can be attributed to the d → sp electronic transition. Lastly, the catalytic activities of the Au$_{40}$(SR)$_{24}$ and Au$_{52}$(SR)$_{32}$ are examined using CO oxidation as a probe. Both medium-sized thiolate-protected gold clusters can serve as effective stand-alone nanocatalysts.

Introduction

Thiolate-protected gold nanoclusters (RS-AuNPs) have received considerable attention over the past decade owing to their unique structures and physicochemical properties compared with other types of gold nanoparticles,1–7 and their potential applications in catalysis, nanotechnology, and biomedicine, among others.8–11 To date, the total atomic structures of a number of RS-AuNPs, including Au$_{110}$(SC$_2$H$_4$)$_{14}$,12 Au$_{100}$(TBHT)$_{16}$ (TBHT = SP$_3$-p-$^2$Bu)$_{13}$, Au$_{124}$(SAdm)$_{16}$,14 Au$_{23}$(PET)$_{18}$–1/0 (PET = phenylethylthiolate SC$_2$H$_2$Ph)$_{15–17}$ Au$_{28}$(TBHT)$_{20}$,18 Au$_{10}$S(S-t-Bu)$_{18}$,19 Au$_{36}$(TBHT)$_{24}$,20 Au$_{18}$(PET)$_{24}$,21 Au$_{102}$(p-MBA)$_{34}$ (p-MBA = p-mercaptopbenzoic acid, SC$_2$O$_2$H$_2$)$_{22}$ Au$_{130}$(p-MBT)$_{30}$ (p-MBT = para-methylbenzenethiol),23 Au$_{133}$(TBHT)$_{32}$,24,25 etc. have been successfully resolved via X-ray crystallography. Among these clusters, Au$_{102}$(p-MBA)$_{34}$ and Au$_{133}$(PET)$_{18}$–1/0 were the first two crystallized and characterized in 2007 and 2008, respectively. The total structure determination of Au$_{102}$(p-MBA)$_{34}$ and Au$_{133}$(PET)$_{18}$–1/0 provides profound insights into many new characteristics such as gold–sulfur bonding, atomic packing structure in RS-AuNPs, staple motifs of ligands, as well as some generic rules of Au distribution among the Au-kernels and ligands. Until now, Au$_{133}$(TBHT)$_{32}$ is the largest ligand-covered gold cluster whose crystal structure has been obtained.

Although tremendous progress has been made in the structural determination of RS-AuNPs from small to large-size via X-ray crystallography, the crystal structures in the medium-size range from Au$_{38}$(PET)$_{24}$ to Au$_{102}$(p-MBA)$_{34}$ are largely unknown. The lack of crystal structures of medium-sized thiolate-protected gold clusters hinders the understanding of the structure–property relationship for many known stable clusters, such as Au$_{44}$(TBHT)$_{28}$,26 Au$_{55}$(PET)$_{31}$,27 Au$_{64}$(SC$_6$H$_{11}$)$_{32}$,28 Au$_{42}$(PET)$_{35}$,29 Au$_{72}$(SC$_6$H$_{13}$)$_{40}$,30 Au$_{76}$(4-MEBA)$_{44}$ (4-MEBA = 4-[2-mercaptopethyl]benzoic acid)$_{31}$ and Au$_{64}$(SPh)$_{42}$,32,33 all synthesized or isolated in the laboratory. Recently, Azubel et al. applied single-particle transmission electron microscopy (SP-TEM) to determine the positions of 68 Au atoms without a single crystal of Au$_{64}$(3-MBA)$_{32}$–34 However, this SP-TEM
measurement still requires theoretical input on the protection ligands to achieve the total structure of the cluster.

Very recently, two medium-sized clusters, Au\(_{40}(o\text{-MBT})_{24}\) and Au\(_{52}(TBTT)_{32}\), have been successfully crystallized by Zeng et al.\(^{35}\) It was found that both nanoclusters are composed of 4-atom tetrahedral units which can coil up into a Kekulé-like ring in the Au\(_{40}(o\text{-MBT})_{24}\) (o-MBT = ortho-methylbenzenethiol) cluster or a DNA-like double helix in Au\(_{52}(TBTT)_{32}\). In particular, the Au\(_{52}(TBTT)_{32}\) cluster is a new species to the RS-AuNP family, while the Au\(_{40}(o\text{-MBT})_{24}\) cluster is akin to the known phenylethylthiolate-capped Au\(_{40}(PET)_{24}\) cluster which is an intermediate in the synthesis of the Au\(_{54}(PET)_{34}\) cluster.\(^{36,37}\) However, the marked differences in the absorption spectrum indicate that Au\(_{40}(o\text{-MBT})_{24}\) and Au\(_{40}(PET)_{24}\) have different structures.\(^{18}\) The Kekulé-like ring in Au\(_{40}(o\text{-MBT})_{24}\) is completely different from the Au\(_{54}\) bi-icosahedral core in the Au\(_{40}(PET)_{24}\) originally proposed by Malola et al.\(^{39}\) The experimental work by Zeng et al. sheds new light on the total structures of RS-AuNPs within the range of Au\(_{38}(PET)_{24}\) and Au\(_{102}(P\text{-MBA})_{44}\).\(^{35}\)

In this communication, we perform an in-depth kernel structure analysis of Au\(_{40}(o\text{-MBT})_{24}\) and Au\(_{52}(TBTT)_{32}\); We classify all known RS-AuNPs into three groups: (1) cubocahedra, (2) Ino-decahedra, and (3) icosahedra. The calculated powdered X-ray diffraction (XRD) results reveal different characteristic peaks for the Au-kernel structures in these groups, which can be useful in identifying the structures of gold nanoparticles when the crystal structure is lacking. In addition, the computed optical absorption spectra of Au\(_{40}(o\text{-MBT})_{24}\) and Au\(_{52}(TBTT)_{32}\) are found to be consistent with experimental measurements. Lastly, the catalytic properties of both clusters are examined by using CO oxidation as a probe.

**Computational methods**

The theoretical powder X-ray diffraction (XRD) curve is calculated using the Debye formula:

\[
I(s) = \sum_i \sum_{j \neq i} \frac{\cos \theta}{(1 + \alpha \cos(2\theta))} \exp \left( - \frac{Bs^2}{2} \right) f_i f_j \sin(2\pi d_{ij}) \frac{2\pi d_{ij}}{2\pi d_{ij}},
\]

where \(s\) is the diffraction vector length and \(\theta\) is the scattering angle, satisfying \(s = 2\sin \theta / \lambda\), \(\lambda\) and \(\alpha\) are determined by the experimental setup and are set to be 0.1051967 nm and 1.01, respectively. \(B\) is the damping factor, which reflects thermal vibrations, and is set to be 0.03 nm\(^2\). The corresponding atomic numbers are used for the scattering factors \(f_i\) and \(d_{ij}\) is the distance between atoms \(i\) and \(j\). The atomic distance \(d_{ij}\) is taken from the optimized structure of clusters based on the density functional theory (DFT) code DMol\(^3\) (version 7.0).\(^{40,41}\)

The generalized gradient approximation with the Perdew–Burke–Ernzerhof (PBE)\(^{42}\) functional and the double numeric polarized (DNP) basis set coupled with semi-core pseudo-potential are employed. Self-consistent calculations are done with a convergence criterion of \(10^{-5}\) hartree for the total energy. The linear and quadratic synchronous transit (LST/QST) method is used to locate the transition state of CO oxidation on the Au\(_{40}(o\text{-MBT})_{24}\) and Au\(_{52}(TBTT)_{32}\).\(^{43–48}\) In all calculations, the \(o\text{-MBT}\) in Au\(_{40}(o\text{-MBT})_{24}\), TBTT in Au\(_{52}(TBTT)_{32}\), and the ligands of other RS-AuNPs are simplified by methyl groups to lower computation cost.

**Results and discussion**

To understand the structure formation of Au\(_{40}(SR)_{24}\) and Au\(_{52}(SR)_{32}\), their kernel structures are analyzed first. The formation of the Au\(_{34}\) kernel of Au\(_{40}(SR)_{24}\) can be divided into two steps, with the Au\(_{13}\) cuboctahedra as building blocks, as shown in Fig. 1. In step 1, the two Au\(_{13}\) cuboctahedra interpenetrate each other to form a Au\(_{26}\) structure by sharing six Au atoms. This rod-like Au\(_{20}\) geometry has been found as a kernel in the structures of Au\(_{30}(SR)_{20}\) and Au\(_{30}(S(R)_{18}\).\(^{18,19}\) Next, one of the Au\(_{13}\) cuboctahedra in the Au\(_{20}\) interpenetrates with the other two Au\(_{13}\) cuboctahedra in the same way as in step 1 to form the kernel of Au\(_{34}\) with a fcc-like structure. Likewise, as shown in Fig. 2, the formation of the Au\(_{44}\) kernel of Au\(_{52}(SR)_{32}\) can be divided into three steps in the same way as Au\(_{40}\) in Au\(_{40}(SR)_{24}\). In step 1 three Au\(_{28}\) geometries interpenetrate each other to form an Au\(_{26}\) structure. Next, the Au\(_{35}\) structure interpenetrates with another Au\(_{13}\) cuboctahedra to form a Au\(_{28}\) geometry, a kernel found in the structure of Au\(_{30}(SR)_{20}\).\(^{17}\) In step 3, the Au\(_{44}\) kernel with fcc-like symmetry can be formed through two interpenetrating Au\(_{28}\) geometries. The above structure analysis on the kernels of Au\(_{40}(SR)_{24}\) and Au\(_{52}(SR)_{32}\), as well as Au\(_{30}(SR)_{20}\), Au\(_{30}(S(R)_{18}\), and Au\(_{36}(SR)_{24}\) (ESI Fig. S1†), shows that the Au kernels of all these clusters with
fcc-like symmetry are comprised of several interpenetrating cuboctahedra.

Besides Au_{28}(SR)_{20}, Au_{30}S(SR)_{18}, Au_{36}(SR)_{24}, Au_{40}(SR)_{24}, and Au_{52}(SR)_{32}, structure analysis for the kernels of other RS-AuNPs such as Au_{25}(SR)_{18}, Au_{38}(SR)_{24}, Au_{102}(SR)_{44}, Au_{130}(SR)_{50}, and Au_{133}(SR)_{52} are presented in ESI Fig. S2 and S3.† Interestingly, according to different types of kernel structures, those crystallized RS-AuNPs larger than Au_{25}(SR)_{18} can be classified into three categories, as shown in Fig. 3: the first category includes Au_{28}(SR)_{20}, Au_{30}S(SR)_{18}, Au_{36}(SR)_{24}, Au_{40}(SR)_{24}, and Au_{52}(SR)_{32}, whose Au kernels possess fcc-like symmetry, are comprised of several interpenetrating cuboctahedra. The Au_{102}(SR)_{44} and Au_{130}(SR)_{50} nanoclusters in which the Au kernels possess Ino-decahedral (D_{5h}) symmetry can be grouped into the second category. The last category includes the remaining RS-AuNPs such as Au_{25}(SR)_{18} and Au_{133}(SR)_{52} with icosahedral Au kernels, and Au_{38}(SR)_{24} with a fused bicosahedral kernel.

Simulated powder X-ray diffraction (XRD) curves of these crystallized RS-AuNPs exhibit different characteristic peaks for nanoclusters with different Au kernel structures. First, we confirm the reliability of the theoretical formula for computing the XRD curves by comparing the simulated XRD curves and the measured XRD curves of Au_{25}(SR)_{18} and Au_{38}(SR)_{24}, as shown in ESI Fig. S4.† Next, the XRD spectra of the RS-AuNPs in all three categories are presented in Fig. 4. From Fig. 4(a), one can see that the Au_{28}(SR)_{20}, Au_{30}S(SR)_{18}, Au_{36}(SR)_{24}, Au_{40}(SR)_{24}, and Au_{52}(SR)_{32} nanoclusters exhibit similar diffraction patterns, where a main peak is located at about 4.0 nm\(^{-1}\), and the two weaker peaks are located at 6.5 nm\(^{-1}\) and 7.5 nm\(^{-1}\), respectively. In the second category (Fig. 4(b)), the consistent XRD curves of Au_{102}(SR)_{44} and Au_{130}(SR)_{50} nanoclusters, where only one less pronounced peak is located at 7.7 nm\(^{-1}\), are significantly different from those in the first category. For the last category, as shown in Fig. 4(c), the diffraction pattern of the Au_{133}(SR)_{52} nanocluster, where four weaker peaks are located in the range 5–9 nm\(^{-1}\), differ from those of Au_{25}(SR)_{18} and Au_{38}(SR)_{24}. The multi-peak diffraction...
pattern can be attributed to the multi-shell structure of the \( \text{Au}_{133}(\text{SR})_{52} \) nanocluster. The analysis above shows that the diffraction pattern is closely related to the structure of the Au kernel in RS-AuNP, which can offer clues to theoretical prediction of the structure of RS-AuNP even when the single crystal is lacking.

Fig. 5 presents the simulated UV/Vis optical absorption spectra of \( \text{Au}_{40}(\text{SR})_{24} \) and \( \text{Au}_{52}(\text{SR})_{32} \) based on the time-dependent DFT (TD-DFT) computation. In addition, the atomic orbital (AO) component of Kohn–Sham molecular orbitals is displayed. As shown in Fig. 5(a), the extrapolated optical band-edge of \( \text{Au}_{40}(\text{SR})_{24} \) is 1.1 eV, consistent with the measured optical gap.\(^{39} \) The extrapolated optical band-edge of \( \text{Au}_{52}(\text{SR})_{32} \) is 1.25 eV (Fig. 5(c)). The population analysis of Kohn–Sham (KS) molecular orbitals, as shown in Fig. 5(b) and (d), demonstrates that the occupied frontier molecular orbitals are mainly contributed from the Au(5d), denoted as the d band, while the Au(6sp) atomic orbitals show dominant contributions to the unoccupied orbitals (sp band). The feature absorption peaks can therefore be assigned to the d → sp interband transition.

Finally, the catalytic properties of \( \text{Au}_{40}(\text{SR})_{24} \) and \( \text{Au}_{52}(\text{SR})_{32} \) are examined by using CO oxidation as a probe. To this end, a few surface staple motifs on the clusters are removed to allow for the catalytic reaction.\(^{49,50} \) The computed catalytic reaction pathway for CO oxidation on the \( \text{Au}_{40}(\text{SR})_{24} \) cluster is shown in Fig. 6(a). CO and O\(_2\) are coadsorbed initially with a binding energy of −1.76 eV. Then the O\(_2\) molecule moves toward the adsorbed CO molecule to form a bridge-like metastable intermediate state characterized by the O–C–O–O species with the

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**Fig. 4** Theoretical powder X-ray diffraction (XRD) curves of the crystallized RS-AuNPs.

**Fig. 5** Simulated UV/Vis absorption spectra and Kohn–Sham orbital energy level diagrams of \( \text{Au}_{40}(\text{SR})_{24} \) (a and b) and \( \text{Au}_{52}(\text{SR})_{32} \) (c and d). The energies are in eV. Each Kohn–Sham orbital is drawn to show the relative contributions (line length with color labels) of the atomic orbitals of Au(6sp) olive, Au(5d) blue, S(3p) magenta, and other orbital contributions from C and H atoms are in black.
O–O bond length being 1.45 Å, and the CO oxidation requires overcoming a reaction barrier of 0.51 eV (TS1). The O–O bond length is further elongated to 1.81 Å, while CO fully grasps an O atom of O2 to form a CO2 molecule. The CO2 can eventually desorb by overcoming a relatively low-energy barrier of 0.35 eV (TS2), leaving the other O atom adsorbed on the gold cluster.

For CO oxidation on the Au52(SR)32 cluster, as shown in Fig. 6(b), the CO and O2 molecules can be favorably co-adsorbed on two neighboring low-coordinated Au atoms, with the coadsorption energy of CO and O2 being about −1.66 eV. The reaction pathway is similar to that for the Au40(SR)24, in which the two molecular species require to overcome reaction barriers of 0.35 eV (TS1) and 0.57 eV (TS2), respectively, to arrive at the final product state. Both reaction barriers are comparable to those of typical nanogold catalysts, indicating that the Au40(SR)24 and Au52(SR)32 clusters can be a stand-alone nanoscale catalyst for CO oxidation.

Conclusions

We have performed a systematic structure analysis of two medium-sized Au40(SR)24 and Au52(SR)32 nanoclusters. We find that the kernel structures of both nanoclusters can be viewed as several interpenetrating cuboctahedra. Based on this observation, we suggest classification of the crystallized RS-AuNPs into three groups according to their kernel structures. In the first group, Au kernels of RS-AuNPs such as Au28(SR)20, Au30(SR)18, Au36(SR)24, Au40(SR)24, and Au52(SR)32 exhibit fcc-like symmetry, comprising several interpenetrating cuboctahedra. In the second group, Au kernels of RS-AuNPs such as Au102(SR)44 and Au130(SR)50 possess icosahedral (D5h) symmetry. The third group includes Au25(SR)18, Au133(SR)52 with icosahedral Au kernels, or Au38(SR)24 with a fused icosahedral Au kernel. Clusters in each group give rise to distinct diffraction patterns, thereby providing important clues for the theoretical prediction of the structure of RS-AuNPs as long as the XRD data are known. We have also computed the UV/Vis absorption spectra and Kohn–Sham orbital energy level diagrams for Au40(SR)24 and Au52(SR)32. Our calculation indicates that the first absorption peak can be assigned to the d → sp interband transition. Examination of the catalytic properties of Au40(SR)24 and Au52(SR)32 suggests that both clusters can be stand-alone nanoscale catalysts for CO oxidation.

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References

Electronic Supplementary Information

Medium-Sized $\text{Au}_{40}(\text{SR})_{24}$ and $\text{Au}_{52}(\text{SR})_{32}$ Nanoclusters with Distinct Gold-Kernel Structure and Spectroscopy Features

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Figure S1. Formation of Au$_{20}$ kernel of Au$_{28}$(SR)$_{20}$ and Au$_{30}$S(SR)$_{18}$ (a), and Au$_{28}$ kernel of Au$_{36}$(SR)$_{24}$ (b). The Au atoms marked with the same number and color are fused together.
Figure S2. Two orthogonal views of multi-shell Au kernels of $\text{Au}_{102}(\text{SR})_{44}$ (a) and $\text{Au}_{130}(\text{SR})_{50}$ (b). All the structures displayed possess Ino-decahedral ($D_{5h}$) symmetry. The shells are shown in different colors.
Figure S3. (a) Formation of $Au_{23}$ kernel of $Au_{38}(SR)_{24}$ by fusing two icosahedral $Au_{13}$. The Au atoms marked with the same color are fused together. (b) Two orthogonal views of multi-shell Au kernel of $Au_{133}(SR)_{52}$. The icosahedral $Au_{13}$ (icosahedral $Au_{13}$ is also the kernel of $Au_{25}(SR)_{18}$) and $Au_{55}$ structures are displayed. The shells are shown in different colors.
Figure S4. The theoretical (black) and experimental (red) powder X-ray diffraction (XRD) curves of Au$_{25}$(SR)$_{18}$ (a) and Au$_{38}$(SR)$_{24}$ (b). The experimental XRD curves of Au$_{25}$(SR)$_{18}$ and Au$_{38}$(SR)$_{24}$ are adapted from refs. 1 and 2, respectively.

Supplemental References