Toward the Solution Synthesis of the Tetrahedral Au$_{20}$ Cluster

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We report the observation in solution of the recently discovered tetrahedral Au$_{20}$ cluster coordinated with eight PPh$_3$ (Ph = phenyl) ligands. The composition and molecular weight of the Au$_{20}$(PPh$_3$)$_8$ complex were confirmed by the isotopic pattern and accurate mass measurement of its doubly charged cation using high-resolution mass spectrometry. A collision-induced dissociation experiment showed that four PPh$_3$ ligands can be easily removed from Au$_{20}$(PPh$_3$)$_8^{2+}$, resulting in a highly stable Au$_{20}$(PPh$_3$)$_4^{2+}$ ion. This observation is consistent with the tetrahedral Au$_{20}$, in which the four apex sites are expected to bond to the PPh$_3$ ligands strongly, and is confirmed by theoretical calculations that predict a highly stable Au$_{20}$(PH$_3$)$_4$ complex with Au–PH$_3$ bond energies of ~1 eV. The current experimental and theoretical observations suggest that large quantities of ligand-stabilized tetrahedral Au$_{20}$ can be obtained, opening the door for exploring its anticipated novel chemical, optical, and catalytic properties.

One of the longstanding objectives of cluster science is to discover highly stable clusters and use them as building blocks for bulk cluster-assembled materials. The discovery of C$_{60}$ in the gas phase$^1$ and its subsequent bulk synthesis$^2$ provide classical inspiration and a prototypical example. However, it is generally believed that few other stable gaseous clusters may be assembled into bulk materials because of cluster–cluster interactions leading to agglomeration. Indeed, no cluster-assembled materials have been synthesized on the basis of stable gaseous clusters other than the fullerenes, despite intensive experimental and theoretical efforts.$^3$–$^9$ Recently, we reported that a 20-atom gold cluster possesses a tetrahedral (T$_d$) structure with a remarkably large HOMO–LUMO energy gap, suggesting that it would be highly chemically inert and may have novel optical and catalytic properties.$^{10}$ Here we report the observation of the tetrahedral Au$_{20}$ cluster in solution, ligated with triphenyl phosphine (PPh$_3$) ligands, and its confirmation by theoretical calculations. The current work represents a successful synthetic effort directly guided and motivated by a gas-phase observation, validating the gas-phase-to-condensed-phase approach for the discovery of cluster-assembled nanomaterials.

There has been a long history of investigations on gold nanoparticles.$^{11}$ Recent interest has been stimulated by the discovery of catalytic effects in supported gold nanoparticles$^{12}$–$^{15}$ and the potential applications of gold nanoparticles in nanoelectronics and nanobiology.$^{16}$–$^{22}$ Gold nanoparticles are generally synthesized in solution with closely packed thiol ligands on their surfaces as a protection layer.$^{11}$,$^{13}$ The smallest thiol-protected gold nanoparticles are ~0.9 nm in diameter and are proposed to contain 28 Au atoms.$^{24}$ Many small multinuclear gold clusters with PPh$_3$ ligands have been synthesized and characterized crystallographically.$^{25}$ The largest structurally
characterized gold clusters with PPh₃ ligands contain 13 and 39 atoms, although a “greengold” compound has been identified and proposed to contain 75 Au atoms. The tetrahedral Au₂₀ occupies a unique position in the hierarchy of gold clusters and nanoparticles. There are no known PPh₃-coordinated gold clusters, nor are there known thiol-protected gold nanoparticles with a T₄Au₂₀ core. The T₄ structure of Au₂₀ is highly unique; it can be viewed as a fragment of the bulk fcc lattice, but unlike the bulk crystal, it has a large energy gap. It is also the largest metal cluster with close packing and yet with every atom on the surface. The only other known transition-metal cluster compound that has been structurally characterized to have a 20-atom T₄ core is [Os₂₀(CO)₄₀]²⁻.

Because of potential cluster–cluster agglomeration, Au₂₀ must also be protected by ligands in order to use it as a building block for cluster-assembled materials. To maintain the unique structural and electronic properties of the T₄Au₂₀, the ligands have to be carefully chosen. The thiophenol ligands have strong chemical interactions with Au and would alter the electronic structure of the T₄Au₂₀. In addition, thiol ligands tend to form a close-packed monolayer on gold nanoparticles, leaving no atomic sites for catalysis. Thus, we chose the zero-valent PPh₃ ligands, which form only dative bonding to Au and offer the potential that both the T₄Au₂₀ core and its unique electronic structure would not be destroyed. Inasmuch as the four apex sites of Au₂₀ are more reactive, we reasoned that the bulky PPh₃ ligands may coordinate only to these sites, leaving 16 uncoordinated surface sites for chemisorption and catalysis. Preliminary theoretical calculations (Figure 1A–C) revealed that the Au₂₀(PR₃)₄ (R = H, Ph) complexes indeed possess high stability. Figure 1A shows the optimized structure of Au₂₀ coordinated with four PPh₃ ligands; its van der Waals surface is shown in Figure 1B. We also calculated Au₂₀(PH₃)₄ for computational simplicity (Figure 1C), expecting that the effect of PH₃ on the Au₂₀ core would be similar to that of PPh₃. Indeed, the only effect that the phosphine ligands have on the Au₂₀ core is that the four face-centered Au atoms tend to be pushed outward from the four Au₁₀ faces: the calculated distance between the face-centered atoms is increased from 3.1 Å in Au₂₀ to 3.4 Å (4.0 Å) in Au₂₀(PPH₃)₄ [Au₂₀(PH₃)₄]. But the ligated complexes still maintain a large HOMO–LUMO energy gap: 1.44 eV in Au₂₀(PPh₃)₄ and 1.82 eV in Au₂₀(PH₃)₄, compared to 1.77 eV in the parent Au₂₀.

We started our synthesis of PPh₃-protected Au₂₀ using a two-phase system similar to that used in the literature for the synthesis of small phosphine-stabilized gold nanoparticles. AuCl(PPh₃) (1.482 g, 3 mmol) and tetrabutylammonium bromide (1.803 g, 3.3 mmol) were dissolved in a toluene/water mixture (200 mL/150 mL). After ca. 15 min, 1.134 g (30 mmol) of NaBH₄ dissolved in 25 mL of water was added dropwise via cannula under vigorous stirring. The reaction mixture turned dark brown instantly under bubbling. The reaction was stirred overnight. The organic phase was separated from the aqueous layer and washed with water (2 × 200 mL). The toluene was removed in vacuo under gentle heating (60 °C). The resulting black solid was dissolved in dichloromethane and precipitated by the slow addition of pentane. The solid was filtered and washed several times with hexanes (3 × 100 mL), a 1:1

**Figure 1.** Computed structures of Au₂₀ and its phosphine complexes. (A) Au₂₀(PPh₃)₄. (B) van der Waals surface of Au₂₀(PPh₃)₄. (C) Au₂₀(PH₃)₄. (D) Au₂₀(PH₃)₈.
methanol/water mixture (3 \times 100 \text{ mL}), diethyl ether (50 \text{ mL}), and pentane (2 \times 50 \text{ mL}). The final product was dried in vacuo with a yield of 507 mg.

A small portion of the solid sample was dissolved in an ethanol/CH\textsubscript{2}Cl\textsubscript{2} mixed solvent (10:1 volume ratio), which was centrifuged to remove the large particles. The solution turned from dark to brownish after the centrifugation. A drop of the brownish solution was spread onto an amorphous carbon-coated copper grid and dried in preparation for high-resolution transmission electron microscopy (HRTEM). The HRTEM image (Figure 2) showed that the soluble samples contained gold nanoparticles with diameters as large as 3 nm, but the majority of the particles have diameters of less than 1 nm. Each edge of the bare Au\textsubscript{20} was calculated to be 0.81 nm and was not changed in the PH\textsubscript{3}-capped complex. Thus, the \textit{T}_{\text{L}} Au\textsubscript{20} core is expected to give a triangular image with edges slightly less than 1 nm. Indeed, numerous triangular particles with dimensions of less than 1 nm can be found in the TEM image, consistent with the anticipated PPH\textsubscript{3}-capped Au\textsubscript{20}.

We further characterized the sample using a high-resolution Fourier transform ion cyclotron resonance (FTICR) mass spectrometer, which was accurately calibrated and equipped with an electrospray ionization (ESI) source.\textsuperscript{a} The solution sample was introduced into the ESI ion source through direct infusion using a syringe pump. The mass analysis was carried out in an electrospray ionization (ESI) source.\textsuperscript{34} The solution sample was first isolated from a broadband mass-to-charge (\textit{m}/\textit{z}) range corresponding to Au\textsubscript{20}(PPH\textsubscript{3})\textsubscript{8}\textsuperscript{2+} (\textit{m}/\textit{z} = 2493.848), as shown in Figure 3A. Prominent singly charged ions were observed, corresponding to five- and six-atom gold clusters, but no doubly charged ions at \textit{m}/\textit{z} = 2493.848 were observed. However, a careful examination of the mass spectrum revealed that doubly charged ions corresponding to Au\textsubscript{20} clusters with eight and seven PPH\textsubscript{3} ligands were observed at \textit{m}/\textit{z} = 3018.030 and 2886.984, respectively. We also observed mass peaks corresponding to Au\textsubscript{20}, but familiar clusters such as the undecanuclear Au\textsubscript{11} species were not observed under our experimental conditions.

Figure 3B shows the observed isotopic pattern of the Au\textsubscript{20}(PPH\textsubscript{3})\textsubscript{8}\textsuperscript{2+} cation primarily due to the \textsuperscript{13}C isotope, compared with the simulated isotopic pattern (Figure 3C). The perfect agreement between the experimental and the simulated isotopic patterns, as well as the accurate \textit{m}/\textit{z} measurement, unequivocally confirmed the correct identification of the composition of the Au\textsubscript{20}(PPH\textsubscript{3})\textsubscript{8}\textsuperscript{2+} cluster. Such good agreement was also obtained for the observed Au\textsubscript{20}(PPH\textsubscript{3})\textsubscript{7}\textsuperscript{2+} ions. To obtain structural information for the Au\textsubscript{20}(PPH\textsubscript{3})\textsubscript{7}\textsuperscript{2+} cluster, we conducted collision-induced dissociation (CID) experiments in the FTICR cell. The Au\textsubscript{20}(PPH\textsubscript{3})\textsubscript{8}\textsuperscript{2+} ions were first isolated from a broadband detection (Figure 4A) and then excited to collide with background gas molecules (10\textsuperscript{-7} Torr of benzene in this case). The excitation method used was the so-called sustained off-resonance

**Figure 2.** High-resolution transmission electron microscopy image of the synthesized Au–phosphine nanoparticles. The arrow points to one possible Au\textsubscript{20} cluster. HRTEM analysis was carried out on a JEOL JEM 2010 microscope fitted with a LaB\textsubscript{6} filament and a specified point-to-point resolution of 0.194 nm. The operating voltage of the microscope was 200 kV. The image was digitally recorded with a slow-scan CCD camera (image size 1024 \times 1024 pixels, Digital Micrograph software from Gatan, Pleasant, CA).

**Figure 3.** High-resolution mass spectroscopic characterization of Au–PPH\textsubscript{3} complexes. (A) Mass spectrum containing the Au\textsubscript{20}(PPH\textsubscript{3})\textsubscript{8}\textsuperscript{2+} and Au\textsubscript{20}(PPH\textsubscript{3})\textsubscript{7}\textsuperscript{2+} doubly charged clusters. (B) Isotopic pattern of the Au\textsubscript{20}(PPH\textsubscript{3})\textsubscript{8}\textsuperscript{2+} mass peak. (C) Simulated isotopic pattern using the natural isotopic abundances of Au, P, C, and H.
is intact in the PPh$_3$-coordinated clusters, which is consistent with our initial expectation and calculation that the four apex PPh$_3$ ligands can be readily dissociated from Au$_{20}$(PPh$_3$)$_8^{2+}$, were terminated at x = 4, and no more PPh$_3$ could be removed under the current CID condition.

Irradiation (SORI), which has proven to be more efficient for the dissociation of large molecules such as large peptides and proteins. The CID experiment (Figure 4B) revealed that four PPh$_3$ ligands can be readily dissociated from Au$_{20}$(PPh$_3$)$_8^{2+}$, resulting in a highly stable Au$_{20}$(PPh$_3$)$_4^{2+}$ cluster ion, which could not be dissociated further under our current experimental conditions.

The CID results suggested that the tetrahedral core of Au$_{20}$ is intact in the PPh$_3$-coordinated clusters, which is consistent with our initial expectation and calculation that the four apex sites of Au$_{20}$ are the most reactive sites that bind strongly to the four PPh$_3$ ligands. We further computed a Au$_{20}$ cluster coordinated by eight PPh$_3$ (Figure 1D), with the four additional PPh$_3$ ligands coordinated to the face-centered sites. Our calculations showed that the T$_d$ Au$_{20}$ core is intact in Au$_{20}$(PPh$_3$)$_8^{2+}$, which also has a large energy gap (1.55 eV) that is only slightly smaller than that of Au$_{20}$. The edge length of the T$_d$ Au$_{20}$ was not influenced by the PPh$_3$ coordination; the only structural change is the outward move of the face-centered atoms. The four face-centered atoms in Au$_{20}$ form a smaller tetrahedron, with a Au–Au distance of 3.1 Å that is increased to 4.0 Å in Au$_{20}$(PPh$_3$)$_4$ and 4.7 Å in Au$_{20}$(PPh$_3$)$_8$. The average binding energy of PPh$_3$ to the face-centered sites was calculated to be 16 kcal/mol, which is still sizeable but smaller than that of the apex sites. We should point out that the Au–PPh$_3$ binding energy is expected to be larger for the apex sites and smaller for the face-centered sites because of steric repulsion between the bulkier PPh$_3$ ligands compared to that of the PPh$_3$ ligands used in the calculations. Thus, our theoretical results were completely consistent with the CID experiment and confirmed that the T$_d$ Au$_{20}$ core is a robust cluster building block.

The phosphine ligands are electron donors, causing the increase in the HOMO energies of Au$_{20}$ from -5.60 eV in the bare cluster to -4.51 eV in Au$_{20}$(PPh$_3$)$_4$ and -3.89 eV in Au$_{20}$(PPh$_3$)$_8$. This result suggests that the latter has a relatively low ionization energy and may exist as doubly charged closed-shell cations in solution. This explains why we were not able to observe singly charged Au$_{20}$(PPh$_3$)$_6^{+}$ from the ESI source. It also explains why we did not observe negatively charged Au$_{20}$(PPh$_3$)$_6^{-}$ ions when the negative mode was used in the ESI. The weaker Au$_{20}$(PPh$_3$)$_8^{2+}$ ion signals may be produced from the dissociation of the parent Au$_{20}$(PPh$_3$)$_8^{2+}$ in the ESI source. The current experimental and theoretical results suggest that the T$_d$ Au$_{20}$ cluster coordinated with phosphine ligands may be obtained in bulk quantity. It is expected that by increasing the size of the ligands one can synthesize the T$_d$ Au$_{20}$ clusters with only the four apex sites coordinated. These clusters may be promising catalysts with the highest surface area and well-defined surface sites.

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References and Notes

(31) The Au$_{20}$–phosphine clusters were calculated via relativistic density functional theory using the ADF 2003 code [ADF 2003.01, SCM, Theoretical Chemistry, Vrije Universiteit: Amsterdam, The Netherlands (http://www.scm.com)]. All geometries were fully optimized by using an
analytical energy gradient technique without imposing any symmetry restrictions. The gradient-corrected PW91 exchange-correlation functional was used in the electronic structure calculations and geometry optimizations (Perdew, J. P.; Wang, Y. Phys. Rev. B 1992, 45, 13244). The scalar relativistic effects were taken into account via the zero-order-regular approximation (ZORA) (Lenthe, E. Van; Baerends, E. J.; Snijders, J. G. J. Chem. Phys. 1993, 99, 4597). Spin–orbit coupling effects were evaluated on the basis of single-point calculations performed at the optimized scalar relativistic structures. The Slater-type-orbit (STO) basis sets with triple-ζ plus two polarization functions (TZ2P) quality were used for Au and P atoms, with basis sets of double-ζ plus one polarization function (DZP) for C and H atoms (Lenthe, E. Van; Baerends, E. J. J. Chem. Phys. 1993, 99, 4597). The frozen-core approximation was applied to the [1s²] cores of C, [1s²−2p¹] core of P, and [1s²−4f⁰] core of Au (Baerends, E. J.; Ellis, D. E.; Ros, P. Chem. Phys. 1973, 2, 42).


(33) All reagents used in the synthesis were commercially available. Chlorotriphenylphosphine gold(I) was purchased from Strem Chemicals; tetraoctylammonium bromide, from Fluka; and sodium borohydride, from Aldrich. Toluene was purchased from Aldrich and was packed under nitrogen in a Sure/Seal bottle. Deionized water was used in the reaction and in the subsequent washes. The water was sparged with argon. All reagents were used as received without further purification. All manipulations were carried out under an atmosphere of argon using standard Schlenk and cannula techniques.

(34) The mass spectrometry analysis of the Au₂₀–PPh₃ clusters was carried out on an FTICR mass spectrometer equipped with a 9.4-T superconducting magnet. This instrument was manufactured by Bruker Daltonics (Billerica, MA), and it is the latest model in the BioAPEX series (APEX-Q), which has an Apollo ESI source and a quadrupole/hexapole interface to allow external ion generation, selection, and fragmentation. The instrument was mass calibrated prior to sample analysis using a Hewlett-Packard calibration mixture (Flanagan, J. M. U.S. Patent 5,872,357, 1999). The average mass measurement error in the mass range up to \( m/z \) 4000 was <2 ppm. The mass spectral data were acquired using a 1-MB data set, and the peak resolution (ΔM/M) at \( m/z \) 1000 exceeded 100,000. At such high resolution, the isotopic information of the analyte ions was totally resolved.


(37) The average collision energy for SORI-CID could be calculated using an equation introduced by Marshall’s group. See Huang, Y. L.; Pasa-Tolic, L.; Guan, S. H.; Marshall, A. G. Anal. Chem. 1994, 66, 4385. For the CID experiment performed with Au₂₀(PPh₃)²⁺, the estimated average energy was ~0.64 eV.

(38) The HOMO of \( Tₐ \) Au₂₀ is doubly degenerate. However, in Au₂₀(PPh₃)₈ the total molecular symmetry is lowered, even though the Au₂₀ core still maintains its local \( Tₐ \) symmetry. Thus, the HOMO degeneracy is lifted in Au₂₀(PPh₃)₈, leading to a closed-shell dication when two electrons are removed from Au₂₀(PPh₃)₈. It should be noted that in the dication the Au₂₀ core possesses only 18 valence electrons, which is also a major shell closing in the electron shell model, supporting the viability of the dication. We also optimized the structure of the Au₂₀(PPh₃)²⁺ dication theoretically and confirmed its stability. Our preliminary calculations showed that the structure of the dication is very similar to that of the neutral.

(39) In the negative-ion mode, only a few low-intensity singly charged anions below \( m/z \) 2500 were observed. The \( m/z \) values and the isotope distributions of these ions indicated that they were very small Au-containing clusters with fragmented ligands most likely due to extensive dissociations from larger and positively charged parent clusters during electrospray.