

An Organometallic Approach to Gold Nanoparticles: Synthesis and X-Ray Structure of CO-Protected $\text{Au}_{21}\text{Fe}_{10}$, $\text{Au}_{22}\text{Fe}_{12}$, $\text{Au}_{28}\text{Fe}_{14}$, and $\text{Au}_{34}\text{Fe}_{14}$ Clusters**

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Ligand-stabilized quasi-molecular gold nanoparticles, consisting of chunks of cubic close-packed lattice featuring “magic numbers” of gold atoms, are being intensively investigated both experimentally and theoretically,^[1] owing to their potential use in miniaturized basic devices in electronics,^[2] models and precursors of metallic catalysts,^[3] stains of biological samples,^[4] and imaging nanoprobe for drug screening and diagnosis.^[5] The monodispersity of several of the above thiol/thiolate or phosphine monolayer protected gold nanoparticles has been disputed.^[6] Furthermore, STM experiments and DFT calculations of adsorption of methylthiolate on Au(111) showed formation of linear RS-Au-SR staple motives.^[7] This feature was later confirmed by calculations on an Au_{38} cluster of O_h symmetry,^[8] and has been experimentally demonstrated by the first structural characterizations of gold-thiolate clusters, namely $[\text{Au}_{25}(\text{SCH}_2\text{CH}_2\text{Ph})_{18}]^{-19}$ and the giant $[\text{Au}_{102}(p\text{-MBA})_{44}]$ ($p\text{-MBA} = p\text{-mercaptobenzoic acid}$) cluster.^[10] The only other structurally characterized gold particles exceeding 1 nm in at least one dimension are the ligand-protected $[\text{Au}_{16}(\text{AsPh}_3)_8\text{Cl}_6]$,^[11] $[\text{Au}_{25}(\text{PPh}_3)_{10}(\text{SET})_5\text{Cl}_2]^{2+}$,^[12] and $[\text{Au}_{39}(\text{PPh}_3)_{14}\text{Cl}_6]\text{Cl}_2$ clusters,^[13] which do not display such a feature. Herein, we report an organometallic approach to a new kind of molecular ligand-stabilized gold nanoparticle, consisting of the synthesis of Au-Fe colloidal nanoparticles, in which $\{\text{Fe}(\text{CO})_x\}$ ($x = 3, 4$) moieties take the place of thiol or thiolate ligands in protecting and stabilizing the gold kernel. These iron carbonyl groups share and may also exceed the bonding versatility of thiols/thiolates.^[14]

The synthesis of CO-protected Au-Fe clusters involves the oxidation of $[\text{Fe}_3(\text{CO})_{11}]^{2-}$ with $[\text{AuCl}_4]^-$ salts in acetone and under an inert atmosphere. After formation of the previously unknown yellow-orange $[\text{Au}_5\{\text{Fe}(\text{CO})_4\}_4]^{3-}$ cluster, the reaction affords brown solutions of colloidal Au-Fe

nanoparticles, which display broad and unresolved IR carbonyl absorptions shifting from 1960 to 2010 cm^{-1} as a function of the starting ratio of the reagents. The colloidal nature of these solutions was confirmed by dynamic light scattering (DLS) measurements in acetonitrile for two samples. The first sample (ν_{CO} at 1980 cm^{-1}) revealed the presence of two sets of nanoparticles displaying hydrodynamic diameters in the ranges 10–30 and 100–200 nm. The second sample (ν_{CO} at 1990 cm^{-1}) showed particles with nominal diameters of 35–60 and 110–300 nm. No evidence of smaller particles could be gathered. These results parallel recent measurements of solutions from which monodispersed $[\text{Au}_{25}(\text{SCH}_2\text{CH}_2\text{Ph})_x]$ was obtained in good yields.^[15]

The addition of Au^{3+} salts in excess gives rise to separation of gold powder and the formation of the dark-green $[\text{Au}\{\text{Fe}_2(\text{CO})_8\}_2]^-$ cluster, two isomers of which have previously been isolated and characterized by other routes.^[16] In our investigations of the above two samples we have so far isolated five molecular species, namely, $[\text{NET}_4]_3[\text{Au}_5\{\text{Fe}(\text{CO})_4\}_4]$ (ν_{CO} in CH_3CN at 1945s, 1861s cm^{-1}), $[\text{NET}_4]_6[\text{Au}_{21}\{\text{Fe}(\text{CO})_4\}_{10}]\cdot\text{Cl}$ (ν_{CO} in CH_3CN at 1982s, 1937sh, 1889sh cm^{-1}), $[\text{NET}_4]_6[\text{Au}_{22}\{\text{Fe}(\text{CO})_4\}_{12}]\cdot(\text{CH}_3)_2\text{CO}\cdot 0.5\text{C}_6\text{H}_{14}$ (ν_{CO} in CH_3CN at 1980s, 1925sh, 1880sh cm^{-1}), $[\text{NET}_4]_8[\text{Au}_{28}\{\text{Fe}(\text{CO})_3\}_4\{\text{Fe}(\text{CO})_4\}_{10}]\cdot 6\text{CH}_3\text{CN}$ (ν_{CO} in CH_3CN at 1985s, 1927sh, 1887sh cm^{-1}) and $[\text{NET}_4]_{10}[\text{Au}_{34}\{\text{Fe}(\text{CO})_3\}_6\{\text{Fe}(\text{CO})_4\}_8]\cdot 2\text{Cl}\cdot 7.6\text{CH}_3\text{CN}$ (ν_{CO} in CH_3CN at 1990s, 1932sh, 1900sh cm^{-1}). Their structures have been determined by single-crystal X-ray diffraction studies.^[17]

The $[\text{Au}_5\{\text{Fe}(\text{CO})_4\}_4]^{3-}$ cluster (**1**) is isostructural with the corresponding $[\text{Cu}_5\{\text{Fe}(\text{CO})_4\}_4]^{3-}$ ^[18] and $[\text{Ag}_5\{\text{Fe}(\text{CO})_4\}_4]^{3-}$ ^[14] species (see Figure S1 in the Supporting Information). As shown in Figure 1, $[\text{Au}_{22}\{\text{Fe}(\text{CO})_4\}_{12}]^{6-}$ (**2**) may be formally envisioned to derive by sandwiching two $[\text{Au}_5]^{3+}$ fragments between three $[\text{Au}_4\{\text{Fe}(\text{CO})_4\}_4]^{4-}$ ^[19] moieties in a triple-decker fashion. The outer $\{\text{Fe}(\text{CO})_4\}$ groups adopt C_{3v} local symmetry of the carbonyl groups and behave as triply bridging (μ_3) ligands, whereas the central $\{\text{Fe}(\text{CO})_4\}$ groups adopt C_{2v} local symmetry of the carbonyl groups and behave as μ_4 ligands.

The $[\text{Au}_{21}\{\text{Fe}(\text{CO})_4\}_{10}]^{5-}$ structure (**3**) may be envisioned as a molecular model of Licurgo's cup^[1] (Figure 2). The metal frame consists of an inner Au-centered pentagonal antiprism, at the top and bottom of which two pentagonal Au_5Fe_5 rings are condensed. As a result the whole metal frame may be described as deriving from two fused concave cups generated by a $\text{Au}_5\text{Fe}_5\text{-Au-Au-Au-Au}_5\text{Fe}_5$ sequence of layers, sharing the unique Au atom and with opposite orientations. In spite of

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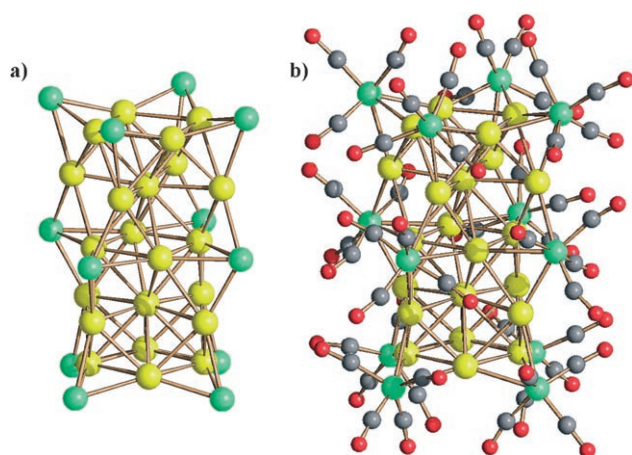


Figure 1. The metal frame (a) and the complete structure (b) of **2** (Au yellow, Fe green, C gray, O red).

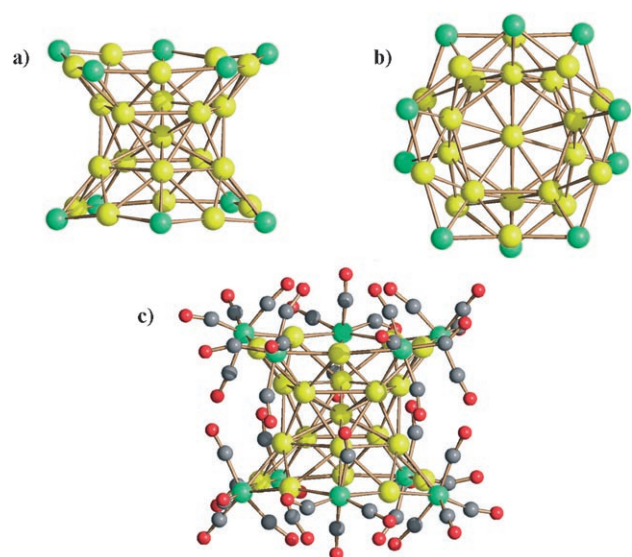


Figure 2. Two views of the metal frame (a, b) and the complete structure (c) of **3** (color legend as in Figure 1).

the width of the Au_5Fe_5 rims, the two cups do not host any molecule or ion since one CO ligand of each $\mu_3\text{-}\{\text{Fe}(\text{CO})_4\}$ fragment is directed toward the center of the rim and gives rise to a “lid” of the cup. Although such a structure does not have any precedent, certain structural motives are reminiscent of some ligand-protected Au and Au–Ag nanoparticles. Thus, the Au atoms describe two fused noncentered and incomplete icosahedrons as the bis(icosahedral) $[\text{Ag}_{12}\text{Au}_{13}(\text{p-Tol}_3\text{P})_{10}\text{Br}_8]^+$ rotamer (Tol = tolyl).^[20] Secondly, the Au_5Fe_5 rims of the two cups represent a cyclic variation of the staple motives present in both $[\text{Au}_{25}(\text{SCH}_2\text{CH}_2\text{Ph})_{18}]^{-[9]}$ and $[\text{Au}_{102}(\text{p-MBA})_{44}]^{-[10]}$.

In contrast to **2** and **3**, $[\text{Au}_{28}\{\text{Fe}(\text{CO})_3\}_4\{\text{Fe}(\text{CO})_4\}_{10}]^{8-}$ (**4**) contains both $\{\text{Fe}(\text{CO})_4\}$ and $\{\text{Fe}(\text{CO})_3\}$ moieties and displays a rather complex metal frame (Figure 3 and Figure S2 in the Supporting Information). The metal framework of **4** consists of an inner tetra-capped, centered pentagonal prismatic Au_{15} core. Two fused and centered pentagonal prisms are the

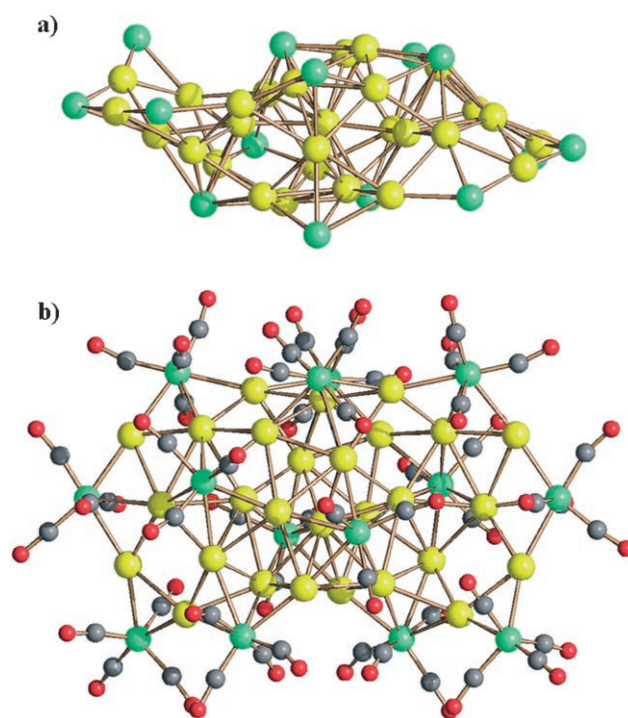


Figure 3. The metal frame (a) and the complete structure (b) of **4** (color legend as in Figure 1).

innermost core of the giant $\text{Au}_{102}(\text{p-MBA})_{44}$ cluster.^[10] Capping of the two pentagonal faces of the Au_{15} moiety with $\{\text{Fe}(\text{CO})_3\}$ groups and addition of two $\{\text{Au}_2\text{Fe}(\text{CO})_3\}$ moieties (to give rise to two additional $\{\text{Au}_5\text{Fe}(\text{CO})_3\}$ pentagonal bipyramids) generates a compact $\text{Au}_{19}\text{Fe}_4$ central core of the cluster. Assembly on both sides of two quasi-planar $\{\text{Au}_5\{\text{Fe}(\text{CO})_4\}_3\}$ and $\{\text{Au}_4\{\text{Fe}(\text{CO})_4\}_3\}$ staples (recalling the $\{\text{Au}_5(\text{SR})_2\}$ staples of $[\text{Au}_{25}(\text{SCH}_2\text{CH}_2\text{Ph})_{18}]^{-[9]}$ and $[\text{Au}_{102}(\text{p-MBA})_{44}]^{-[10]}$) generates two dangling “baskets” with opposite orientations (up and down). The structure is then completed by capping the resulting four concave Au_4 butterfly faces with four additional $\mu_4\text{-}\{\text{Fe}(\text{CO})_4\}$ groups.

The structure of $[\text{Au}_{34}\{\text{Fe}(\text{CO})_3\}_6\{\text{Fe}(\text{CO})_4\}_8]^{8-}$ (**5**) is shown in Figure 4, and a step-by-step construction of the metal framework is given in Figure S3 in the Supporting Information. The Au_{34} core consists of an inner Au_6 octahedron interstitially lodged in an Au_{24} polyhedron having 6 pentagonal and 26 triangular faces (or 6 rhombic and 14 triangular faces). The above Au_{24} polyhedron is formally generated by the condensation of a pentagonal ring of gold atoms on top of each of the six vertices of the interstitial Au_6 octahedron. Capping of these six pentagonal faces with $\mu_6\text{-}\{\text{Fe}(\text{CO})_3\}$ groups gives rise to an inner $\text{Au}_{30}\text{Fe}_6$ core. Condensation of two $\text{Au}_2\{\text{Fe}(\text{CO})_4\}_3$ staples on two opposite rhombic faces of the $\text{Au}_{30}\text{Fe}_6$ core generates two outer distorted trigonal prisms with an edge bridged by a $\mu\text{-}\{\text{Fe}(\text{CO})_4\}$ group and the two triangular faces capped by $\mu_4\text{-}\{\text{Fe}(\text{CO})_4\}$ groups. Finally, two $\mu_3\text{-}\{\text{Fe}(\text{CO})_4\}$ groups cap two opposite rhombic faces and complete the shielding of the gold particle. Once again, a staple motif is present as for $[\text{Au}_{25}(\text{SCH}_2\text{CH}_2\text{Ph})_{18}]^{-[9]}$, $[\text{Au}_{102}(\text{p-MBA})_{44}]^{-[10]}$, **3**, and **4**.

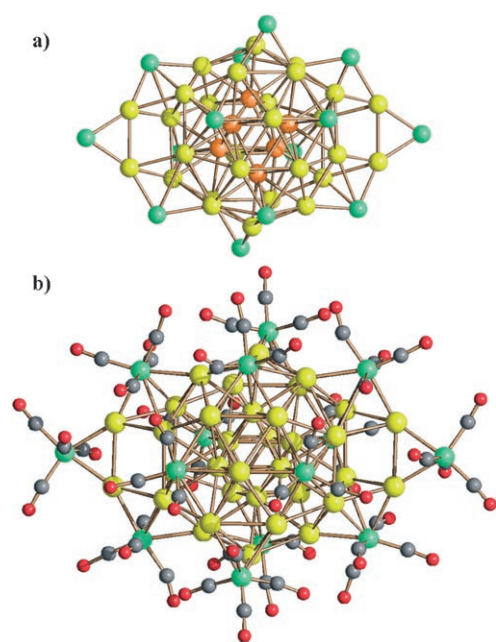


Figure 4. The metal frame (a) and the complete structure (b) of **5** (color legend as in Figure 1; the interstitial Au₆ octahedral moiety in (a) is marked in orange).

In conclusion, framework fragments of C_5 local symmetry seem to dominate all structures, with the notable exception of **2**. Even relatively small variations in composition of otherwise similarly sized CO-protected Au–Fe clusters result in large changes of the structure of the gold kernel. These changes are probably favored by the bonding versatility of the {Fe(CO)₄} and {Fe(CO)₃} moieties. However, the correspondence of some structural motives between the so far structurally characterized thiolate-protected gold clusters and the title compounds confirms the need of some wariness in determining supposedly molecular features by measurements carried out on collections of quasi-monodispersed ligand-protected gold nanoparticles, rather than a single nanoparticle.^[21]

Moreover, staple motives, the oxidation state of surface gold atoms, and the energy of Au atomic orbitals are likely to concur in delaying the insulator-to-metal transition as their nuclearity increases, relative to the more compact transition-metal carbonyl clusters.^[22] As suggested by EHMO calculations, the HOMO–LUMO gaps of **2–5** are in the range 0.8–1.3 eV (see the Supporting Information), in fair agreement with calculated^[8] and experimental values of monodispersed Au₃₈ molecule-like nanoparticles.^[23]

The measured diameters of CO-protected Au–Fe nanoparticles deserve some specific comments. Indeed, as shown in Table 1, these are at least one and two orders of magnitude smaller than those suggested by DLS experiments. A plausible explanation is that the above highly charged cluster anions are present in solution as cation–anion aggregates. Partially in keeping with this suggestion, the hydrodynamic diameters determined for the same sample in acetonitrile and DMSO are different. In the latter, there is clear evidence of nanoparticles of approximately 4 nm, in addition to other

Table 1: Maximum diameters (nm) of the metal frames.^[a]

	2	3	4	5
A	1.53	1.353	1.85	1.92
B	2.14	1.90	2.40	2.43

[a] Measured (X-ray structure) from the outermost Fe nuclei including twice the Fe metal radius (A), and the outermost oxygen atoms including twice the van der Waals radius of oxygen (B).

spread in the ranges 8–40 and 100–300 nm. The lowest value is in line with the dimensions of **2–5** within the first coordination sphere of the solvent cage.

It is, however, possible that some decomposition under laser radiation could also contribute to the large diameters detected. Finally, it can be anticipated that the above CO-protected Au–Fe clusters may turn out to be valuable precursors of miscellaneous inorganic oxide–metal composites of potential interest in the rapidly expanding field of gold catalysis.^[3] Thermal treatment of their [NEt₄]⁺ salts gives rise to Au, Au–Fe, or Au_nFe_xO_y powders, as a function of experimental conditions. The study of their catalytic performances is underway.^[24]

Experimental Section

A solution of [NEt₄][AuCl₄] (0.60 g, 0.62 mmol) in acetone (10 mL) was slowly added to a solution of [NEt₄]₂[Fe₃(CO)₁₁] (0.36 g, 0.62 mmol) in acetone (30 mL) under N₂. The final molar ratio was finely adjusted with small additions of [NEt₄][AuCl₄] to tune the main carbonyl absorption to 1980 cm^{−1}. The resulting dark brown suspension was filtered, and the filtrate was dried in vacuum and washed with THF (3 × 10 mL) to eliminate [NEt₄][AuFe₄(CO)₁₆]. A first extraction with acetone (20 mL) and precipitation by layering *n*-hexane gave orange crystals of [NEt₄]₃[Au₅Fe₄(CO)₁₆] (minor product) and black crystals of [NEt₄]₆[Au₂₂Fe₁₂(CO)₄₈]·(CH₃)₂CO·0.5C₆H₁₄ (yield 20% based on Au. Anal. results (%) found: Au 59.6, Fe 9.51; calcd: Au 59.94, Fe 9.29). Subsequent extraction of the residue with acetonitrile (20 mL) and precipitation by layering diisopropyl ether gave black crystals of [NEt₄]₆[Au₂₁Fe₁₀(CO)₄₀]·Cl (yield 5% based on Au. Elemental analysis (%): found: Au 62.1, Fe 8.32; calcd: Au 62.38, Fe 8.44).

The same procedure was applied for the synthesis of the second sample, the only variation being a fine tuning of the main IR carbonyl absorption to 1990 cm^{−1}. The resulting dark brown suspension was filtered, and the filtrate was dried in vacuum and washed with THF (3 × 10 mL) to eliminate [NEt₄][AuFe₄(CO)₁₆] and with acetone (2 × 10 mL) to eliminate [NEt₄]₃[Au₅Fe₄(CO)₁₆] and [NEt₄]₆[Au₂₂Fe₁₂(CO)₄₈]. A first extraction of the residue with acetonitrile (10 mL) and precipitation by layering diisopropyl ether gave black crystals of [NEt₄]₈[Au₂₈Fe₁₄(CO)₅₂]·6CH₃CN (yield 15% based on Au. Anal. results (%) found: Au 61.3, Fe 8.42; calcd: Au 61.00, Fe 8.67). A second extraction with acetonitrile (20 mL) gave black crystals of [NEt₄]₁₀[Au₃₄Fe₁₄(CO)₅₀]·2Cl·7.6CH₃CN (yield 20% based on Au. Elemental analysis (%): found: Au 63.20, Fe 7.50; calcd: Au 63.31, Fe 7.41). The overall yield of Au–Fe carbonyl species (including the yet uncharacterized ones) measured by Au analysis of the filtered reaction solutions was greater than 70%.

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