2013

Effect of polymer ligand structures on fluorescence of gold clusters prepared by photoreduction

Luo Li
Huazhong University of Science and Technology

Zhen Li
University of Wollongong, zhenl@uow.edu.au

Hui Zhang
Huazhong University of Science and Technology

Shoucun Zhang
Huazhong University of Science and Technology

Irfan Majeed
Huazhong University of Science and Technology

See next page for additional authors

Publication Details
Effect of polymer ligand structures on fluorescence of gold clusters prepared by photoreduction

Abstract
Blue emission fluorescent Au$_5$ clusters with maximum quantum yield of 20.1% were synthesized by a simple photoreduction method using three specially designed tridentate polymer ligands. The evolution of fluorescent Au nanoclusters (Au NCs) under UV irradiation was studied by fluorescence, UV-Vis and X-ray photoelectron spectroscopic techniques, suggesting that the fluorescence of Au NCs is size-dependent and is associated with the presence of Au$^{(i)}$ ions in the Au NCs. The effect of polymer structure on the fluorescent Au NCs has also been discussed. These highly fluorescent Au NCs have potential applications in the fabrication of optoelectronic devices and light emitting materials.

Keywords
fluorescence, gold, clusters, prepared, photoreduction, effect, polymer, ligand, structures

Disciplines
Engineering | Physical Sciences and Mathematics

Publication Details

Authors
Luo Li, Zhen Li, Hui Zhang, Shoucun Zhang, Irfan Majeed, and Bien Tan

This journal article is available at Research Online: http://ro.uow.edu.au/aiimpapers/582
Effect of polymer ligand structures on fluorescence of gold clusters prepared by photoreduction†

Luo Li, a Zhen Li, b Hui Zhang, a Shoucong Zhang, a Irfan Majeed a and Bien Tan a

Blue emission fluorescent Au6 clusters with maximum quantum yield of 20.1% were synthesized by a simple photoreduction method using three specially designed tridentate polymer ligands. The evolution of fluorescent Au nanoclusters (Au NCs) under UV irradiation was studied by fluorescence, UV-Vis and X-ray photoelectron spectroscopic techniques, suggesting that the fluorescence of Au NCs is size-dependent and is associated with the presence of Au(i) ions in the Au NCs. The effect of polymer structure on the fluorescent Au NCs has also been discussed. These highly fluorescent Au NCs have potential applications in the fabrication of optoelectronic devices and light emitting materials.

1 Introduction

Fluorescent metal nanoclusters (NCs) as a new class of metal nanomaterials have attracted great attention in recent years for their unique properties and applications. 1–5 These metal NCs are composed of a few to hundreds of atoms and possess dramatically different optical, electrical and physical/chemical properties compared to metal nanoparticles since the continuous density of states become discrete energy levels when their size is reduced to the Fermi wavelength of electrons. 6

New synthesis protocols for preparation of fluorescent metal NCs are being extensively explored because of their potential applications in the fields of optoelectronics, bio-imaging, biological sensing, catalysis and so on. 7–9 The most common methods in this respect involve; (1) reduction of metal precursors in the presence of stabilizers such as thiols-compounds, 10–12 polymers, 13–15 dendrimers, 16–18 oligonucleotides 19–21 and proteins 22,23 by chemical reduction, 24 photoreduction 25 or microwave assistant approach 26–28 and (2) etching of larger nanoparticles into smaller NCs using various thiols and biomolecules. 29–32 In these syntheses, ligands play an important role in obtaining these ultra-small fluorescent metal nanoclusters owing to their ability to protect the metal core efficiently. 33

Among the metal nanoclusters, Au nanoclusters (Au NCs), which are less toxic, more biocompatible, and highly stable under ambient conditions, have attracted more attention for potential applications than other noble metal materials. 34 In our previous work, 35 PTM-PVAc polymer ligand was employed to synthesize fluorescent Au NCs. However, the fluorescence of PTM-PVAc itself makes it difficult to ascertain whether the fluorescence comes from the Au NCs. In this paper, three types of non-fluorescent tridentate thiocetethyl-terminated polymer ligands, poly(methyl methacrylate) (PTM-PMMA), poly(n-butyl methacrylate) (PTMP-PBMA) and poly(n-etyl methacrylate) (PTMP-PrBMA) were designed to synthesize blue fluorescent Au NCs by a simple and facile photoreduction method (Scheme 1). The highest fluorescence quantum yields (QY) of Au NCs stabilized with these non-fluorescent ligands PTM-PMMA, PTMP-PBMA and PTMP-PrBMA were found to be 3.8, 14.3 and 20.1%, respectively, which are much higher than that of red emission Au NCs prepared in our previous work. 36–38 The use of non-fluorescent ligands clearly proved the observed fluorescence originates from the Au NCs rather than the ligands themselves. Moreover, these designed ligands are very useful for studying the role of polymer structure in the formation of fluorescent Au NCs. Compared to other methods, 35–37 the photoreduction approach is simple and feasible for avoiding the use of additional reducing agents, which is good for studying the photofluorescence mechanism of Au NCs. These tridentate polymer ligands have better protective capability than monodentate polymer ligands. 39–41 Comparing to the bio-imaging applications of water-soluble red-emitting Au NCs prepared in our previous work, 36–38 the resultant blue-emitting Au, clusters are also equally important because of their applications in the field of light emitting materials and optoelectronic devices for their high QY.

†Electronic supplementary information (ESI) available: Gel permeation chromatography and 1H NMR spectroscopy of thioether polymer ligands; DLS spectra, fluorescence spectra and TEM images of Au NCs. See DOI: 10.1039/c2nr33690f
2 Experimental

2.1 Chemicals

All chemicals are of analytical grade and used as received without any further purification, unless otherwise described. Methyl methacrylate (MMA, 99%), n-butyl methacrylate (nBMA, 99%), tert-butyl methacrylate (tBMA, 99%), hydrogen tetrachloroaurate (HAuCl₄·4H₂O, 47.6%), 2,2’-azobis(isobutyronitrile) (AIBN), anhydrous ethanol, methanol, tetrahydrofuran (THF) and 1,4-dioxane were purchased from National Medicines Corporation Ltd of China. Pentaerythritol tetrakis-3-mercaptopropionate (PTMP, 97%) was obtained from Aldrich.

2.2 Synthesis of polymer ligands

The synthesis of PTMP-PBMA is described here as an example of preparation of polymer ligands. In a typical synthesis, 40 mL 1,4-dioxane solution containing monomer tBMA (7.110 g, 50 mmol), chain-transfer agent PTMP (0.489 g, 1.00 mmol) and initiator AIBN (0.082 g, 0.50 mmol) were added to a round-bottom flask fitted with a reflux condenser under nitrogen protection with magnetic stirring. The temperature of the reaction mixture was maintained at 80 °C and refluxed for 6 h. The viscous products were isolated by precipitating into a cold mixture of methanol and water (volume ratio 3:1). A fraction of low molar mass polymer, unreacted monomer and some oligomers formed due to incomplete reaction were removed during the precipitation step. The excess solvent was removed by evaporation using a vacuum oven set at 40 °C for 24 h, and the yield was 91%. PTMP-PMMMA and PTMP-PBMA were synthesized by using MMA and BMA as monomers in a similar way; their yields were 82 and 80%, respectively. The molecular weights of these three polymers were calculated based on the ratio of monomer units to the terminal group in the 1H NMR spectra. The polymer molecular weights were further determined with gel permeation chromatography (GPC) by using polystyrene as calibration standard. Comparison of molecular weights measured by NMR and GPC are listed in Table S1 (ESI†).

2.3 Preparation of Au nanoclusters

Au NCs were prepared according to the following method. Polymer ligand (PTMP-PMMMA, PTMP-PBMA or PTMP-PbBMA) was added into 5 mL of 1 mM THF solution of HAuCl₄ and incubated for 30 min, the final concentration of polymer ligand was set to 4 mM, then subjected to an ultraviolet light source (8 W, wavelength = 365 nm) for various time intervals. The as-prepared Au NCs solution was first centrifuged at 16,000 rpm two times and then purified using a microporous filter (0.22 μm). The remaining steps followed our previous work.45

2.4 Characterization

Fluorescence excitation and emission spectra were obtained by a FP-6500 fluorescence spectrometer (Jasco, JPN). UV-Vis absorption spectra were determined using a UV2550 Spectrophotometer (SHIMADZU, JPN). Dynamic light scattering (DLS) measurements were carried out with a ZetaPlus (Brookhaven, USA), which has an effective detection capability from 0.6 to 6000 nm. TEM images were recorded with a Tecnai G2 electron microscope (FEI, USA) at an acceleration voltage of 200 kV. A drop of Au NCs solution was placed on carbon film coated copper grids. X-Ray photoelectron spectroscopy (XPS) spectra were measured with an AXIS-ULTRA DLD (Shimadzu, Japan). The energy resolution was set to 1.7 eV to minimize data. Matrix-assisted laser desorption ionization time-of-flight (MALDI-TOF) was analyzed on a Bruker Autoflex MALDI-TOF instrument. Samples (0.5 μL) were spotted on a MALDI target plate with 0.5 μL CHCA matrix (10 mg mL⁻¹ CHCA in 50% ACN-0.1% TFA/25 mM diammonium citrate) and externally calibrated with a mixture of angiotensin I (5 pmol μL⁻¹)-ACTH (5 pmol μL⁻¹)-insulin (100 pmol μL⁻¹)-cytochrome C (100 pmol μL⁻¹).

2.5 Quantum yield (QY) measurements

The QY of a compound is defined as the fraction of molecules that emit a photon after direct excitation by the source.68 This quantity is not the same as the total number of emitted photons which escape a bulk sample divided by the total number of absorbed photons, although in many instances the two quantities are nearly equal. The measurement of QY employed the comparison method which is described below (eqn (1)):

\[
\Phi_{\text{unk}} = \left( \frac{A_{\text{std}}}{A_{\text{unk}}} \right) \left( \frac{F_{\text{unk}}}{F_{\text{std}}} \right) \left( \frac{\sigma_{\text{unk}}}{\sigma_{\text{std}}} \right) \left( \frac{n_{\text{unk}}}{n_{\text{std}}} \right)^2 \Phi_{\text{std}}
\]

where \( \Phi \) = quantum yield; unk = unknown sample; std = standard; \( n \) = refractive index of solvent; \( A \) = absorption at the selected excitation wavelength; \( F \) = integrated fluorescence signal in the emission region. To calculate the QY of Au NCs, we
measured a series of the samples and the standard 9,10-
diphenylnaphthalene (Φ = 0.91 in ethanol). All the samples
were diluted to ensure optical densities less than 0.02 measured
by a Lambda 35 UV-visible spectrophotometer (Perkin-Elmer,
USA) in order to reduce the error. The emission spectra were
recorded on a FP-6500 fluorescence spectrometer (Jasco, JPN)
under excitation of 375 nm light at 25 °C.

3 Results and discussion

3.1 Synthesis and characteristic of tridentate polymer
ligands

Three types of thioether polymer ligands PTMP-PBMA, PTMP-
PBMA and PTMP-PBMA were prepared by a facile chain-
transfer radical polymerization method using pentamerthritol
tetraakis-3-mercaptopropionate (PTMP) as chain-transfer agent,
methyl methacrylate (MMA), n-butyl methacrylate (BMA) and
terr-butyl methacrylate (TBMA) as monomers, respectively
(Scheme 1a). The molecular weights and structures of polymer
ligands were confirmed by GPC and ^1H NMR analyses (Fig. S1–
S4 and Table S1, ESP). None of the polymer ligands showed
fluorescence under ultraviolet light and no emission spectra
were observed (Fig. S5, ESP).

3.2 Synthesis of blue emitting highly-fluorescent Au NCs

Au NCs were prepared by irradiating a THF solution of HAuCl4
in the presence of the polymer ligand with 365 nm light
(Scheme 1b). As shown in Fig. 1, three as-prepared Au NCs
samples Au NCs@PTMP-PMMMA, Au NCs@PTMP-PBMA and Au
NCs@PTMP-PBMA exhibited the same fluorescence profile
with two emission peaks at 412 and 433 nm and with excitation
maxima of 375 nm. Their double emission peaks could be
attributed to the surface properties of Au NCs influenced by the
solvent, which lead to the electronic property splitting and elec-
tron redistribution of the cluster surfaces. These Au NCs were
well dispersed in THF and formed a colorless suspension in
ambient light, while showing blue fluorescence under a UV
lamp (Fig. 1d). The samples were kept at −5 °C and their fluo-
rescence could be retained for more than two months. The
highest fluorescence quantum yields (QY) of Au NCs in THF
solutions were measured to be 3.8, 14.3 and 20.1% for Au
NCs@PTMP-PMMMA, Au NCs@PTMP-PBMA and Au
NCs@PTMP-PBMA, respectively.

The effect of the molar ratio between Au nanoparticles and polymer
ligands on the fluorescence of Au NCs was also investigated
(Fig. S6, ESP). The optimal concentration of polymer ligands
in order to obtain the brightest fluorescence was found to be
4 mM, which was the same for all the three polymer ligands.
The variation in polymer concentration has little effect on the
emission peak position of resulting Au NCs, but very low poly-
mer concentration (<1 mM) led to non-fluorescent species. This
may be due to the insufficient polymer ligands which might not
be able to effectively protect the Au NCs. When the polymer
ligand concentration was increased from 4 to 8 mM, the
declared decrease in the emission intensity can be attributed
to limitation of the growth of Au NCs caused by excessive polymer
ligands.

3.3 Fluorescence mechanism with the evolution of Au NCs

Fluorescence emission and absorption of the Au NCs were also
studied at different UV irradiation times in order to explore the
relationship between fluorescence properties and the growth of
Au NCs. Before irradiation, none of the Au-precursor solutions
showed fluorescence (Fig. 2a–c) and only the Au nanoparticle
absorption peak was found at 340 nm (Fig. 3). The initial Au solutions
gradually changed from light yellow to colorless in 60 min
accompanied with the evolution of emission peaks. A gradual
increase in fluorescence intensity of Au NCs and a decrease in
Au nanoparticle absorption intensity were observed along with
the increase in irradiation time. This indicates the observed fluo-
rescence originates from small Au NCs formed by the reduction
of Au nanoparticles by UV light. The maximum fluorescence intensity
appeared at 480 min with no absorption in the wavelength
range from 400 to 700 nm. DLS studies indicated the diameter
of as-prepared Au NCs were ultra-small (Fig. S7a–c, ESP).

Prolonged irradiation led to a decrease in the fluorescence
intensity with a visible color change in Au NCs solution from
colorless into light pink (Fig. S8, ESP) and the characteristic
surface plasmon resonance (SPR) peak of Au nanoparticles
appeared at 540 nm in the absorption spectra when the irradi-
ation time was extended to 600 min. For Au NCs@PTMP-PBMA,
the particle size was measured to be 2.3 ± 0.1 nm by DLS
(Fig. S7d, ESP), which also supports the observation of the SPR
absorption band. TEM images further confirmed these changes,
particles became irregular and polydisperse, their diameter
increased to more than 2.0 nm and even larger (Fig. S9a, ESP).
These drastic changes indicate that the Au NCs were aggregated.
into Au nanoparticles in this process and the fluorescence of as-prepared Au NCs is decreased. Furthermore, the NCs solution became darker in color gradually and turned into violet in one week at room temperature, which indicated these samples irradiated for a long time were not very stable and underwent slow aggregation. Interestingly, some irregular shapes of Au such as planar nanoparticles more than 20 nm long were observed in TEM images (Fig. S9b, ESI), which may be due to the Ostwald ripening process. The proposed synthesis and evolution of Au NCs is schematically summarized in Scheme 2a.

Zheng and co-workers suggested that these fluorescent Au NCs may be a complex composed of Au(0) atoms and a large percentage of Au(i) ions, and the fluorescence is associated with the fraction of Au(i) ions in the clusters. X-ray photoelectron spectroscopy (XPS) measurements were carried out to analyse the variation of gold valence states in our Au NCs. The Au sample obtained from 480 min irradiation showed that the binding energies (BE) of Au 4f7/2 and Au 4f5/2 are 84.6 and 88.3 eV, respectively (Fig. 4b, solid line). It is notable that the BE of Au 4f7/2 falls in between the Au(0) BE (84 eV) of a metallic gold film and the Au(i) BE (85 eV) of gold thiocyanate, suggesting the co-existence of Au(0) and Au(i) in the clusters. The deconvolution of the Au 4f7/2 band revealed an asymmetric band that can be fitted into two peaks at 84.1 and 85.0 eV, which were assigned to Au(0) and Au(i), respectively (Fig. 4a). According to the integrated area ratio between these two peaks, nearly 60% of Au atoms in the Au NCs were Au(i) and significantly influence the BE of Au atoms. When the UV irradiation time was extended to 600 min, the BE of Au 4f7/2 was shifted from 84.6 to 84.4 eV (Fig. 4b), indicating the decrease in Au(i) fraction. These results support that the fluorescence from Au NCs is also associated with the presence of a fraction of Au(i) ions in the clusters.

3.4 The effect of polymer ligand structures on the fluorescence of Au NCs

According to Jin and Wu, fluorescence of Au NCs could be largely facilitated by employing small molecular ligands with electron-rich atoms or groups. Since polymer ligands have longer carbon chains as compared to the small molecular ligands and small differences in molecular structure of the monomer would be greatly amplified after polymerization, introduction of high electron-donating capability could lead to a dramatic influence on the fluorescence properties of Au NCs. We assumed the Au NCs stabilized with PTMP-PBMA would show the highest fluorescence intensity while those prepared with PTMP-PtBMA would show the lowest. In order to prove this assumption, PTMP-PtMA, PTMP-PBMA and PTMP-PtBMA were designed to have similar molecular weight by adjusting the concentration of chain transfer agent. The chain numbers (repeating units) of polymer ligands were estimated at 38, 37 and 36 for PTMP-PtMA, PTMP-PBMA and PTMP-PtBMA, respectively. Au NCs samples stabilized by these three different polymer ligands were prepared using the same ligand molar ratio (4 mM) under identical irradiation conditions. The fluorescence intensity lies in the sequence Au NCs@PTMP-PtMA > Au NCs@PTMP-PBMA > Au NCs@PTMP-PtBMA (Fig. 2d), which is consistent with the QYs mentioned before, confirming our speculation. Control experiments were performed by using a mixture of monomers (PtMA, PBMA and PtBMA) and PTMP instead of polymers as ligands. However, no fluorescent species could be observed. This indicates polymers play an important role in the obtaining of fluorescent Au NCs.

Scheme 2b shows the possible interactions on the surface of Au NCs with tridentate polymer ligands PTMP-PtMA,
PTMP-PBMA and PTMP-PbBMA. Oxygen atoms from repeating units (e.g. rBMA in PTMP-PbBMA) donate delocalized electron density to the Au core while thiol groups of ligands can be attached to the Au core via strong S–Au bonds and stabilize the Au NCs. When the excited electrons are returned from the high energy state to the low energy states, they can transfer from ligands to the Au cores, which leads to an increase in fluorescence emission. Therefore, the capability of the polymer ligand to donate electrons plays an important role in improving fluorescence. The higher fluorescence intensity and QY of Au NCs@PTMP-PBMA and Au NCs@PTMP-PBMA than Au NCs@PTMP-PMMA could be attributed to the stronger electron-donating abilities of rBMA and BMA owing to their longer carbon chains. For Au NCs@PTMP-PbBMA and Au NCs@PTMP-PBMA, the differences arise due to the isomeric structures of rBMA and BMA. The tert-butyl group shows a hyper-conjugation effect, leading to stronger electron-donating abilities than the n-butyl group. Meanwhile, greater steric hindrance effect of surface PTMP-PBMA may be protects the Au core against quenching by solvent. Both effects could result in the highest fluorescence intensity and QY of Au NCs@PTMP-PBMA. On the basis of these results, we obtained highly fluorescent Au NCs by employing polymer ligands with high electron-donating ability and steric hindrance effect.

It should be noted that all these Au NCs exhibited the same fluorescence emissions at 412 and 433 nm, similar to that of peptin-mediated Au NCs. The fluorescence of metal NCs mainly depend on the particle size effect and the surface ligand effect. In order to determine the core size, we performed matrix-assisted laser desorption ionization mass spectrometry (MALDI-TOF) analysis (Fig. 5). The mass peak at m/z = 1081 is assigned to the Au clusters with a composition of Au₄S₂, whereas the other peaks could be assigned to the fragments of Au₄S₃ (m/z = 884), Au₅S₄ (m/z = 687) and Au₅S₅ (m/z = 655). No larger Auₙ clusters were found in higher mass range up to m/z 10 000. MALDI-TOF results confirmed that all of Au NCs@PTMP-PMMA, Au NCs@PTMP-PbBMA and Au NCs@PTMP-PBMA clusters have the same formulae and are composed of 5 gold atoms. Based on these results, for our Au₄ NCs, surface
ligands have little influence on the excitation and emission profile but only result in different QYs, indicating the fluorescence wavelength dominated by core size of Au NCs, following a free-electron (Jellium) model.8

3.5 The influence of temperature on the fluorescence of Au NCs

The influence of temperature on the fluorescence of Au NCs was investigated by recording fluorescence emission spectra at different temperatures in the range from 5 to 35 °C (Fig. S10, ESI†). The fluorescence intensity was increased linearly with decreasing temperature, the relative fluorescence intensity being decreased from 100% at 5 °C to 90% at 35 °C (Fig. S10 inset, ESI†). This could be explained as due to reduction in medium viscosity due to increase in temperature, which in turn leads to faster molecular movement rate and increased collision probability between the fluorescent Au NCs and solvent or other molecules and resulting in fluorescence quenching and reduction of QY.9 In addition, the fluorescence intensity decreased by 0.33% per degree centigrade, indicating that as-prepared Au NCs were only slightly sensitive to temperature.46

4 Conclusions

A simple and facile synthesis of highly emissive blue fluorescent Au NCs is reported with non-fluorescence multidentate polymer ligands by photo-reduction and the maximum QY of Au NCs was found to be 20.1%. The fluorescence properties of Au NCs under UV irradiation was studied by optical spectra, XPS and MALDI-TOF, suggesting the fluorescence of Au NCs is size-dependent and is associated with the fraction of Au(III) ions in the clusters. Our work also demonstrates that the polymer ligand structure has a greater influence on QY than on collision of Au NCs. Polymer ligands with higher electron-donating abilities and steric hindrance effect could largely promote fluorescence, which might help in the synthesis and theoretical research of fluorescent metal NCs in the future.

Acknowledgements

We thank the Analysis and Testing Center, Huazhong University of Science and Technology for their assistance in characterization of materials. This work was financially supported by the National Natural Science Foundation of China (no. 51173058).

Notes and references