

PAPER



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Gold nanorods derivatized with CTAB and hydroquinone or ascorbic acid: spectroscopic investigation of anisotropic nanoparticles of different shapes and sizes†

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Gold nanorods stabilized by binary ligand mixtures of cetyltrimethylammonium bromide (CTAB, primary ligand) and ascorbic acid or hydroquinone were investigated by complementary synchrotron radiation-induced spectroscopies and microscopies, with the aim to find evidence of the influence of the secondary ligand molecular and chemical structure on the nanorod shapes and size ratios. Indeed, as it is well known that the CTAB interaction with Ag(I) ions at the NR surface plays a key role in directing the anisotropic growth of nanorods, the possibility to finely control the NR shape and dimension by opportunely selecting the secondary ligands opens new perspectives in the design and synthesis of anisotropic nanoparticles.

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1. Introduction

Noble metal nanoparticles (MNPs) have been attracting a huge amount of attention from the scientific community over the last two decades thanks to their peculiar physicochemical properties arising from the maximized surface-to-volume ratio, such as the well-known surface plasmon resonance (SPR).¹ The most-exploited bottom-up approach for MNP synthesis is a chemical method involving the reduction of metal salts (AuHCl₄, AgNO₃) in a medium containing capping ligands, usually organic molecules with thiol or amine end-groups, able to graft the metal surface competing with the metal particle growth. By opportunely modulating the stoichiometric ratio between metal salts, reducing agents and capping ligands, as well as the specific synthesis conditions (time of reaction, temperature) it is nowadays possible to obtain almost monodispersed nanoparticles with desired sizes and shapes.² It is noteworthy that the opportune selection of capping ligands is a crucial step in

the MNP synthesis since on the one hand they prevent disordered growth and avoid agglomeration,³ and on the other hand they can confer to the stabilized MNPs desired properties arising from the specific organic moiety (optical properties, biocompatibility, hydrophilic behaviour, *etc.*).^{4,5}

Among others, anisotropic (*i.e.*, non-spherical) particles have attracted much attention due to their size- and shape-dependent optical properties. Rod-shaped nanoparticles are particularly interesting since they display two plasmon bands: a transverse plasmon band corresponding to an electron oscillation along the short axis of the rod, at around 510–550 nm for gold nanorods (AuNRs), and a longitudinal plasmon band, in the range 600–1200 nm.⁶ Indeed, the nanorods' aspect ratio can be modified by the strict control of the experimental parameters during their chemical synthesis, allowing the opportune tuning of the position of the plasmon bands, for example obtaining nanomaterials active in the NIR, which is of major interest for applications such as therapeutic/imaging (“theragnostic”) agents. Another characteristic of anisotropic nanoparticles in general, and specifically nanorods, that attracts great attention from the research community, is their size-related SPR tunability; for example, AuNRs can experience a 50 nm red shift of the longitudinal plasmon mode for a change in aspect ratio (*i.e.*, length/diameter) from about 2.5 to nearly 3; this is a considerable shift, if we recall that for spherical AuNPs a circa 50 nm shift in the plasmon resonance requires a diameter size modification from 10 to 100 nm (λ_{max} increases with mean diameter, from about 520 nm for smaller NPs to about 570 nm for the larger ones).⁷ Moreover, it is noteworthy that the optical

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† Electronic supplementary information (ESI) available: Table S1: experimental conditions and plasmonic peaks for the synthesised AuNRs. Table S2: XPS data analysis results; Fig. S1: EDX data; Fig. S2: C 1s XPS spectra, also reporting the in-depth description of each spectral component. See DOI: <https://doi.org/10.1039/d3na00356f>



properties (optical absorption or scattering wavelengths) of the NRs strictly depend on the rods' length and are insensitive to their diameter.⁸

All the above reported considerations concur to substantiate the appealing of gold nanorods, but they also introduce a central issue in the design and synthesis of such anisotropic nanoparticles, that is the ability to finely control AuNR aspect ratio, homogeneity and dispersion, since the plasmonic optical properties are much sensitive to these parameters.⁹ To attain this goal, a mandatory step is the comprehension as accurately as possible of the mechanism driving the anisotropic growth that is at the base of NR formation. Since "there is strong experimental evidence that every ingredient/parameter in the synthesis procedure plays a critical role in determining the shape and resulting surface chemistry",¹⁰ the comprehension of the role played by each participant in the synthetic path has been and still is subject to much experimental and theoretical research.

The synthetic procedure most used to prepare AuNRs is a two-step protocol named "seed mediated-method" developed by Murphy and co-workers.¹¹ In the first step, tetra chloroauric acid (HAuCl_4) is reduced with a strong reducing agent as for example sodium borohydride (NaBH_4) in the presence of a stabilizing agent, citrate or cetyltrimethylammonium bromide (CTAB), producing 2–4 nm quasi-spherical Au seeds. Then, in the second step, the seeds are added to a growth solution containing AgNO_3 , a weak reducing agent as for example ascorbic acid, and CTAB, producing AuNRs. Nowadays, the role played by some parameters has been fully elucidated by several research reports. There is general accordance about the formation of a CTAB bilayer on the AuNR surface.¹⁰ The roles of the seeds, silver ions and CTAB concentration have been investigated systematically,¹² the importance of bromine ion as the counterion of CTA^+ as a shape-directing agent, through inhibition of the side growth during the elongation of nanorods, has been largely discussed¹³ as well as the role played in favouring the anisotropic growth by the complex formed *in situ* between CTAB and Ag(I) ions.^{14,15}

As for the silver ions and atoms, there is still debate because while several authors observe metallic silver near the AuNR surface, for example by applying X-ray photoelectron spectroscopy (XPS) and X-ray absorption spectroscopy (XAS),¹⁴ suggesting the role of silver in directing the anisotropic growth by forming a sub-monolayer on selected AuNR side-facets (namely, under potential deposition of silver on AuNRs¹⁴), other authors point out the absence of preferential absorption of Ag(I) ions on certain facets, in contrast with this hypothesis.¹⁶ However, it is well established that the presence of silver ions is necessary to improve the AuNR synthesis yield and to control the aspect ratio, a fact that justifies further investigation of the local structure of both silver atoms and silver ions, since, to the best of our knowledge, a unique picture of Ag species distribution in the AuNR bulk and surface has still not been attained.¹⁷ Another aspect that deserves further investigation is the role played by the secondary ligands, when AuNR synthesis involves the use of a binary ligand mixture. This aspect is still poorly investigated, lacking a systematic investigation leading to correlate the

secondary ligand molecular structure with AuNR dispersion, size-ratio and self-organization.

In this work, we apply complementary microscopic (FE-SEM, TEM) and spectroscopic techniques, in particular state-of-the-art synchrotron radiation-induced X-ray photoelectron spectroscopy (SR-XPS) and X-ray absorption spectroscopy (XAS) to study the molecular, electronic and chemical structure, as well as local structure, particularly around silver and gold atoms, and the morphology of gold nanorods prepared with two different secondary ligands, the weak reducing agents ascorbic acid (AA) and hydroquinone (HQ). The two series of AuNRs, namely AuNRs-AA and AuNRs-HQ, differ in the plasmon band position, morphology and aspect ratio in a reproducible way, due to the influence of the different secondary ligands on the anisotropic growth process. The accurate investigation reported in this work will shed light on the possibility of designing AuNRs of desired aspect ratios and shapes by opportunely selecting the secondary ligand, as well as on the destiny of silver atoms and ions in the AuNRs-AA and AuNRs-HQ nanosystems.

2. Materials and methods

2.1. Materials

Freshly bidistilled water was used as a solvent in all the procedures. All reactants required were purchased and used as received, without further purification: cetyltrimethylammonium bromide ($\text{C}_{18}\text{H}_{42}\text{BrN}$, CTAB, $\geq 97\%$ Millipore Corporation), tetrachloroauric(III) acid trihydrate ($\text{HAuCl}_4 \cdot 3\text{H}_2\text{O}$, $\geq 99.9\%$, Sigma-Aldrich), sodium borohydride (NaBH_4 , 99.99%, Sigma-Aldrich), silver nitrate (AgNO_3 , 99.9%, Carlo Erba), L-ascorbic acid ($\text{C}_6\text{H}_8\text{O}_6$, 99%, Sigma-Aldrich) and hydroquinone ($\text{C}_6\text{H}_6\text{O}_2$, $\geq 99\%$, Sigma-Aldrich). A mini spin Eppendorf centrifuge was used for the purification of AuNR samples. Uv-visible spectra were acquired with a Shimadzu 2401 PC spectrophotometer (200–800 nm), while NIR spectra were measured with a Nicolet IS50 FT-IR Thermo scientific spectrophotometer (400–1100 nm) using quartz cells with a 1 cm optical path.

2.2. Synthesis

CTAB stabilized AuNRs were prepared by a seed-growth approach as reported in the literature:¹⁸ a seed solution of quasi-spherical nanoparticles is dissolved in a growth solution containing its precursor species ($\text{HAuCl}_4 \cdot 3\text{H}_2\text{O}$ and CTAB), AgNO_3 and either ascorbic acid or hydroquinone as a reducing agent. This synthesis relies on a complex and delicate equilibrium of reactants to achieve anisotropic growth conditions: silver ions and a ligand agent show preferential absorption onto specific facets of gold and hinder reduction in these directions in the presence of a weak-enough reducing agent. The seed solution is prepared with 5 mL of $\text{HAuCl}_4 \cdot 3\text{H}_2\text{O}$ (5×10^{-4} M) and 5 mL of CTAB (0.2 M) solutions in deionized water in an inert atmosphere ($\text{Ar}_{(g)}$), to which 0.6 mL of a NaBH_4 solution (10^{-2} M) is added as a reducing agent. After 5 minutes, the solution becomes yellow-brownish and is then stored for 1 hour before use. Various experiments were carried out under different

conditions both with AA and with HQ and then the best samples were compared. The syntheses were repeated in duplicate at least three times, always giving consistent results. Table S1 (see the ESI[†]) shows the most summarized experimental data: the best results, as indicated in Table S1,[†] are the subject of this study. It is noteworthy that only anisotropic rod-shaped nanoparticles were considered for this study (neither spherical nor other non-rod-like nanoparticles were considered).

2.2.1. Synthesis of CTAB stabilized AuNRs using ascorbic acid (AA). 5 mL of a CTAB solution (0.2 M), 5 mL of a HAuCl₄·3H₂O solution (10⁻³ M) and 0.2 mL of a AgNO₃ solution (4 × 10⁻³ M) were mixed (molar ratio Au/CTAB/Ag = 1/200/0.16). Under vigorous stirring and in an inert environment (Ar_(g)), 0.07 mL of a 0.078 M ascorbic acid solution is added (molar ratio Au/AA = 1/1.092), followed by 0.024 mL of a seed solution. Gradually, up to twenty minutes after the addition of the seed, the solution turns to a reddish-violet colour. The product is left to rest for 24 hours and then purified by centrifugation (13 Krpm, 15 min, two times washing with deionized water).

2.2.2. Synthesis of CTAB stabilized AuNRs using hydroquinone (HQ). A solution containing 5 mL of a CTAB solution (0.2 M), 5 mL of a HAuCl₄·3H₂O solution (10⁻³ M) and 0.1 mL of a AgNO₃ solution (4 × 10⁻² M) was prepared (molar ratio Au/CTAB/Ag = 1/200/0.625). Under continuous stirring conditions and in an inert environment (Ar_(g)), 0.5 mL of a 0.1 M solution of hydroquinone is added (molar ratio Au/HQ = 1/10). After the full discoloration of the solution, 0.160 mL of seed solution is added to start the reaction. After 24 hours the solution appears red and the product is purified by centrifugation (13 Krpm, 15 min, two times washing with deionized water).

2.3. Microscopic and spectroscopic methods

2.3.1. Transmission electron microscopy (TEM). The ultrastructural analysis of AuNRs was performed using a TEM instrument (Philips EM 208 S, FEI Company, Eindhoven, The Netherlands) equipped with a tungsten source and magnification up to 200 K. Image acquisition was provided by a MegaView III camera (Olympus Soft Imaging Solutions). A drop of AuNR sample was deposited on a formvar carbon coated copper grid for subsequent TEM observations.

2.3.2. Scanning electron microscopy (SEM). The morphology of the samples was examined using a MIRA 3 (Tescan) SEM operated with an electron beam energy of 30 keV. The elemental composition of the samples was determined by energy-dispersive X-ray spectroscopy (EDX) using an XFlash detector (Bruker) integrated into the SEM. EDX data are reported in the ESI as Fig. S1.[†]

2.3.3. Synchrotron radiation induced X-ray photoelectron spectroscopy (SR-XPS). Experiments were carried out at the SuperESCA beamline at the ELETTRA synchrotron facility of Trieste (Italy). XPS data were collected in fixed analyzer transmission mode (pass energy = 50 eV), with the monochromator entrance and exit slits optimized at 30 and 20 μm, respectively. For the C 1s, N 1s, O 1s and Ag 3d spectral regions, photon energy of 600 eV was used; for Au 4f and Br 3d spectral regions, photon energy of 360 eV was used. The energy resolution was ΔE

= 0.25 eV. Calibration of the energy scale was made referencing the spectra to the C 1s core level signal of aliphatic carbons, found at 285.00 eV, for all samples. SR-XPS measurements were carried out on the two AuNRs-AA and AuNRs-HQ in the solid state, by drop-casting a drop of aqueous suspension onto TiO₂/Si(111) wafer surfaces. Curve-fitting analysis of the C 1s, N 1s, O 1s, Ag 3d, Br 3d and Au 4f spectra was done using Gaussian curves as fitting functions, after subtraction of a polynomial background. The Br3d_{5/2,3/2} doublets were fitted using the same full width at half-maximum (FWHM) for both components and a spin-orbit splitting of 1.04 (ref. 19) and a branching ratio (3d_{5/2}/3d_{3/2}) of 3/2. The Ag3d_{5/2,3/2} doublets were fitted using FWHM for both components with a spin-orbit splitting of 6.0 and a branching ratio of 3/2. For the Au4f_{7/2,5/2} doublets, a splitting of 3.7 eV, a branch ratio 4f_{7/2}/4f_{5/2} of 4/3 and the same FWHM values for both spin-orbit components were applied. When several different species were identified in a spectrum, the same FWHM value was set for all individual photoemission bands.

2.3.4. Near-edge X-ray absorption fine structure (NEXAFS). Measurements were performed at the BEAR beamline (bending magnet for emission absorption and reflectivity) at the ELETTRA storage ring, installed at the left exit of the 8.1 bending magnet. The apparatus is based on a bending magnet as a source and beamline optics delivering photons from 5 eV up to about 1600 eV with a selectable degree of ellipticity. The UHV end station is equipped with a movable hemispherical electron analyzer and a set of photodiodes to collect angle-resolved photoemission spectra, optical reflectivity, and fluorescence yield. In these experiments, we used ammeters to measure drain current from the sample. C and N K-edge spectra were collected at grazing (20°) incidence angles of the linearly polarized photon beam with respect to the sample surface. In addition, our carbon and nitrogen K-edge spectra have been further calibrated using the resonance at 288.70 eV, assigned to the C=O 1s → π* transition, and the resonance at 400.00 eV, assigned to the 1s → σ* transition of the positively charged nitrogen, respectively. The raw C and N K-edge NEXAFS spectra were normalized to the incident photon flux by dividing the sample spectrum by the spectrum collected on a freshly sputtered gold surface. Spectra were then normalized subtracting a straight line that fits the part of the spectrum below the edge and assessing to 1 the value at 330.00 and 420.00 eV for C and N, respectively.

2.3.5. X-ray absorption spectroscopy (XAS). Measurements were carried out at the Au L_{III}-edge (11 919 eV) and at the Ag K-edge (25 514 eV) respectively, at the 11.1R-XAFS beamline at ELETTRA synchrotron radiation facility²⁰ and at the LISA-BM08 beamline²¹ at ESRF (European Synchrotron Radiation Facility). The AuNR samples suspended in water were mixed with cellulose, dried in vacuum, and pressed to obtain homogeneous pellets suitable for handling. Both beamlines were equipped with double crystal, fixed exit monochromators with harmonic rejection mirrors. The Au L_{III}-edge XAS spectra were acquired in transmission geometry keeping the samples in vacuum at liquid nitrogen temperature, with two gas-filled ionizing chambers measuring the incident (*I*₀) and transmitted (*I*₁) X-ray beam intensities. Pure Au foil was placed in vacuum after the *I*₁ chamber, and the transmitted intensity (*I*₂) was measured with

a third ionization chamber placed right after the reference foil. The absorption signals of samples 2 (AuNRs-AA) and 4 (AuNRs-HQ), calculated as $\alpha_{\text{exp}} = \ln\left(\frac{I_0}{I_1}\right)$, and the one of the reference

foil, equal to $\alpha_{\text{ref}} = \ln\left(\frac{I_1}{I_2}\right)$, were acquired, aligned and averaged to improve the data statistics. The Ag K-edge XAS spectra were measured in fluorescence geometry, using a high purity germanium (HP-Ge) multidetector (13 elements, (ORTEC)). The Ag $K\alpha$ fluorescence signal for each detector I_f^i was electronically selected from the total fluorescence yield using the analyser's multichannel electronics. The absorption spectrum of pure Ag foil placed after the sample was measured and used to check the energy calibration during data collection and eventually align the energy scale of the spectra. The spectra of all detector channels (except those with a lower signal-to-noise ratio) were summed, and the absorption signal was calculated as $\alpha_{\text{exp}} = \frac{\sum I_f^i}{I_0}$. Due to the weak Ag content, multiple spectra (19 for AuNRs-AA and 8 for AuNRs-HQ samples) were measured, checked for energy calibration, and averaged to obtain data statistics suitable for quantitative analysis.

To extract EXAFS structural signal χ_{exp} , the experimental spectra, α_{exp} , were treated according to standard procedures²² including the linear pre-edge subtraction ($\alpha' = \alpha_{\text{exp}} - \alpha_{\text{pre}}$), bare atomic background (α_{b}) subtraction, and normalization, to extract EXAFS structural signals: $\chi_{\text{exp}}(k) = \frac{\alpha' - \alpha_{\text{b}}}{\alpha_{\text{b}}}$. The edge energy E_0 , origin of the photoelectron wavenumber $k = \hbar^{-1}\sqrt{2m_e(E - E_0)}$ (m_e being the electron mass), was defined at the first inflection point (maximum of the first derivative) of the pre-edge subtracted spectra α' .

The quantitative analysis of the EXAFS signals was carried out fitting the k^w -weighted theoretical curves $k^w\chi^{\text{th}}$ to the raw experimental data $k^w\chi_{\text{exp}}$, applying a non-linear least-square procedure implemented in the program FiteEXA.²² The Au-L_{III} data analysis has been carried out in the 3–12 (AuNRs-AA) and 3–15 \AA^{-1} (AuNRs-HQ) k range with $w = 2$. The Ag-K edge spectra of AuNRs exhibit a weaker signal-to-noise ratio, which is attributed to either the lower Ag content (that affects the count statistics due to the reduction of the fluorescence signal) or the more disordered/lighter local coordination chemistry (decreasing the amplitude of the EXAFS structure signal). The Ag EXAFS spectra were analyzed in the 3–8 (AuNRs-AA) and 3–12 \AA^{-1} (AuNRs-HQ) k range with $w = 1$. The theoretical curves $\chi^{\text{th}}(k)$ were calculated as a sum of partial contributions χ_i , calculated using a Gaussian pair distribution function model and the standard EXAFS formula^{23,24} with the Gaussian disorder model. The theoretical photoelectron scattering amplitude and phase functions were calculated using the FEFF8 program.²⁵

3. Results and discussion

3.1. AuNR morphology: TEM and FE-SEM images

Transmission electron microscopy analysis revealed different AuNR dimensions depending on the secondary ligand used

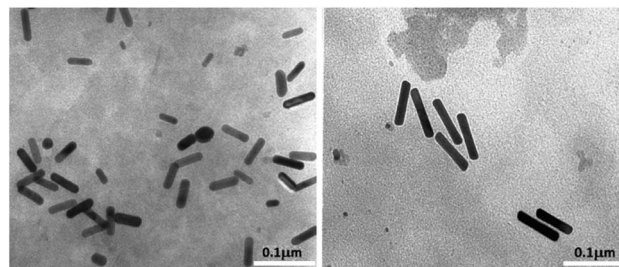


Fig. 1 TEM images of AuNRs: (a) AuNR-AA; (b) AuNR-HQ.

during the synthesis. AuNR-AA (Fig. 1a) was shorter (mean length = 40 ± 6 , mean width = 10 ± 2 nm) than AuNR-HQ (Fig. 1b, mean length = 100 ± 8 , mean width = 14 ± 4 nm).

The variation in the dimensions of AuNRs resulting from the use of different secondary ligands during synthesis is further confirmed by scanning electron microscopy, as illustrated in Fig. 2. Specifically, the AuNRs-AA sample at different magnifications in Fig. 2a and c exhibited smaller nanorods, whereas the AuNRs-HQ sample in Fig. 2b and d is composed of significantly larger nanorods. To highlight the comparison, Fig. 2e summarizes the corresponding histograms of the nanorod size distribution, yielding a mean width of 10 ± 4 nm and length of 41 ± 5 nm for the AuNRs-AA sample (average aspect ratio 4.1) and a mean width of 22 ± 4 nm and mean length of 106 ± 14 nm for the AuNRs-HQ sample (average aspect ratio 4.8).

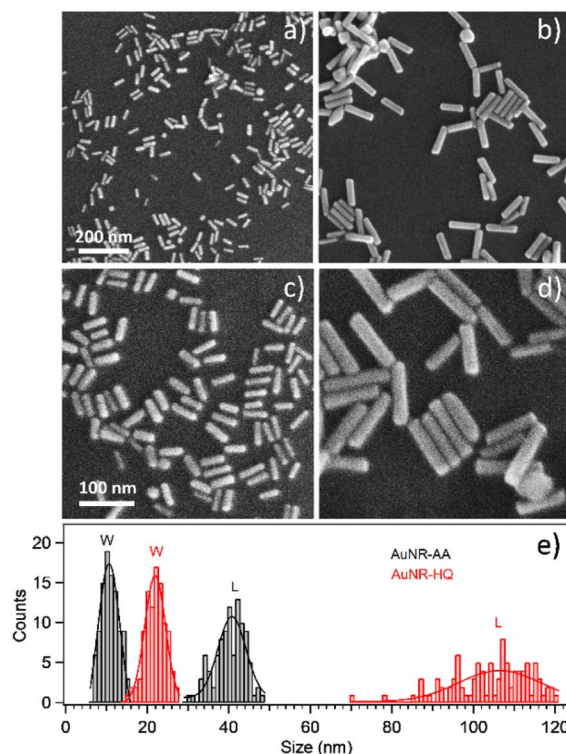


Fig. 2 SEM images of: (a) and (c) AuNR-AA; (b) and (d) AuNR-HQ; (e) corresponding histograms for their size distribution (W stands for width, L stands for length), obtained by analyzing 100 nanorods for each sample.

3.2. Molecular and electronic structure: SR-XPS and NEXAFS results

SR-XPS measurements were carried out on the two AuNRs-AA and AuNRs-HQ in the solid state, as described in the “Materials and methods” section. All SR-XPS data were analyzed by following a peak-fitting procedure, allowing deconvolution of the components arising from chemical elements with different atomic environments; all peak position BEs (binding energies), FWHM values, atomic ratios (relative intensities) and assignments are reported in ESI Table S2.†

C 1s, N 1s, Br 3d, Ag 3d and Au 4f core level signals of AuNRs-AA and AuNRs-HQ were collected and analyzed. C 1s spectra are reported in Fig. 3a (AuNRs-AA) and Fig. 3d (AuNRs-HQ). Both spectra appear composite and consist of at least four components corresponding respectively to aliphatic C–C groups of CTAB and AA or aromatic carbons of HQ (BE = 285.0 eV), to C–N (CTAB) and C–O (AA, HQ) signals (286.5 eV), which can't be deconvoluted with the used resolution, to the C=O carbons of AA, with a contribution from adventitious carbon (287.8 eV)¹⁷ and finally to carboxylic groups of contaminants (289.2 eV).²⁶ (The in-depth description of each spectral component of C 1s spectra is reported in the caption of Fig. S2†).

N 1s spectra (Fig. 3b and e) are made of two groups of components: the peaks at lower BE (400 eV in AuNRs-HQ, 399 eV + 400 eV in AuNRs-AA) are associated with amine-like N atoms of CTAB (400 eV);²⁷ the low-intensity peak at 399 eV in AuNRs-AA indicates amine groups of CTAB shifted at lower BE by ion-ion and/or ion-dipole CTAB-AA electrostatic interactions, as suggested in;²⁸ the reproducible N 1s signal observed in both spectra at around 402.8 eV can be attributed to neatly

protonated nitrogens of the $\text{Ag}^+/\text{Br}^-/\text{CTA}^+$ coordination compound. As observed by some authors, the positive shift in N 1s binding energy can also be interpreted as due to the gold surface vicinity.¹⁵ Correspondingly, Br3d spectra (Fig. 3c and f) show a couple of spin-orbit pairs at lower BE (Br3d_{5/2} BE = 68.2 eV) attributed to unperturbed CTAB molecules^{15,29} pointing out of the surface in the CTAB bilayer (see Scheme 1), while the Br 3d signal at higher BE (Br3d_{5/2} BE = 69.1 eV) is assigned to less negatively charged bromine anions in the “sandwich” structure of the $\text{Ag}^+/\text{Br}^-/\text{CTA}^+$ at the interface between CTAB and AuNR surface.³⁰ This structure is, at least from the SR-XPS point of view, independent of the identity of the secondary ligand, since HQ and AA-containing nanorods display very similar N 1s and Br3d photoelectron spectra, indicative of analogous electronic structures of nitrogen and bromine atoms. This finding suggests that the presence of HQ or AA as a secondary ligand does not affect the CTAB double layer arrangement, as well as the nitrogen and bromide local structure. The very low-intensity signal observed at high BE (70 eV Br3d_{5/2}) in the Br3d spectra (4–6% of Br species, almost at the detection limit of the technique also considering the very complex sample matrix³¹) is attributed to bromine atoms covalently bonded to C,^{32,33} and its presence is probably due to some degradation arising under the X-ray beam, or to contaminants.

As for the gold and silver electronic structure, Au4f and Ag3d core level spectra are reported in Fig. 4. Au4f signals (Fig. 4a and c) are the composite of two spin-orbit pairs (Au4f_{7/2}, Au4f_{5/2}), coherent with literature findings on noble metal nanoparticles of several shapes, either spherical or anisotropic.³⁴ The most intense Au4f_{7/2} signal at about 83.7 eV is assigned to metallic Au(0) atoms in the NR bulk, while the less intense doublet

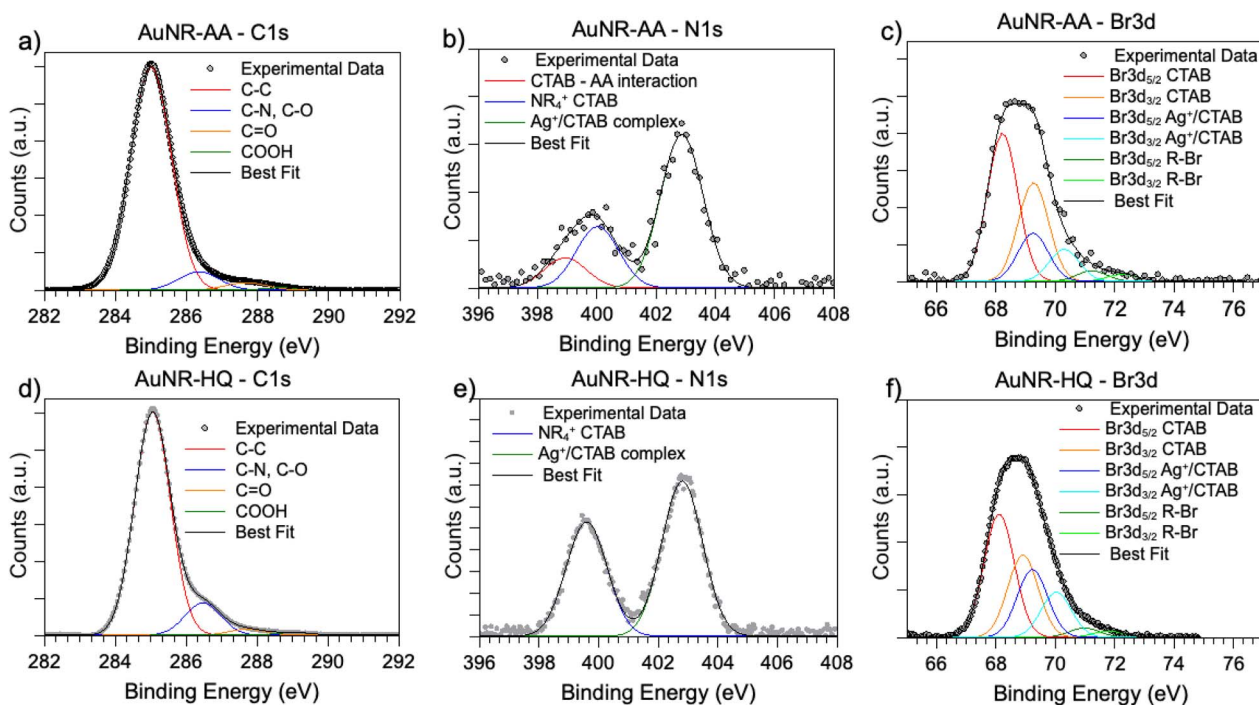
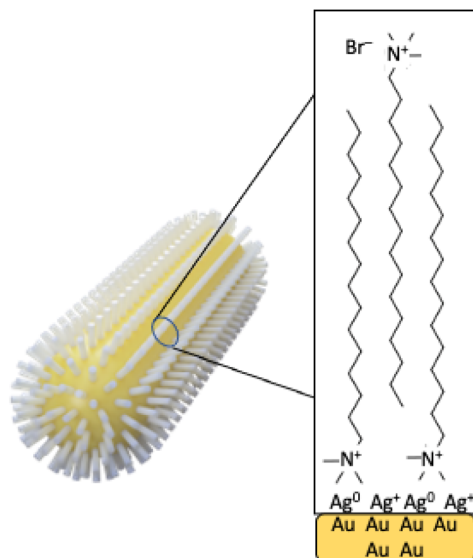


Fig. 3 SR-XPS C 1s, N 1s and Br 3d spectra of AuNRs-AA (a–c) and AuNRs-HQ (d–f).



Scheme 1 Schematic representation of the AuNR surface with the evidenced CTAB bilayer.

($\text{Au}4f_{7/2}$ BE = 84.2 eV) is attributed to a low percentage of gold atoms at the NR surface and interacting with the ligands. The Ag3d spectra (Fig. 4b and d) are also very similar for AuNRs-AA and AuNRs-HQ and still composite allowing individuation of two spin orbit doublets. The most intense signal at lower BE ($\text{Ag}3d_{5/2}$ BE = 367.3 eV) is assigned to Ag^+ ions at the NR surface leading to the $\text{Ag}^+/\text{Br}^-/\text{CTA}^+$, according to the literature on analogous systems.^{15,17} It is noteworthy that this signal percentage is about 90% of all the silver sampled by XPS, *i.e.*, in the first nm of the AuNR surface (XPS sampling depths of a few

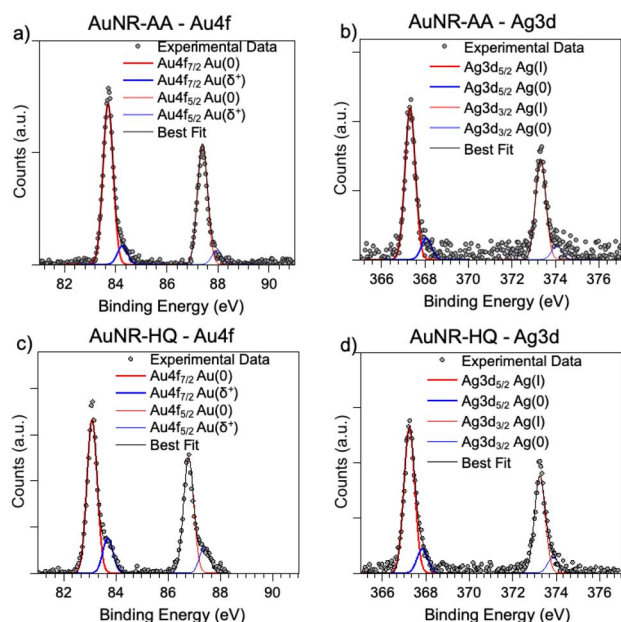


Fig. 4 SR-XPS Au4f and Ag3d spectra of AuNRs-AA (a, b) and AuNRs-HQ (c, d).

nanometers³⁵). The $\text{Ag}3d_{5/2}$ signal at higher BE (nearly 368 eV) is indicative of the presence of metallic $\text{Ag}(0)$ atoms,¹⁷ also located at the NR surface. In the literature there is still an ongoing debate about the presence of metallic silver atoms in AuNRs and their location. The finding presented here locates a low amount of $\text{Ag}(0)$ on the nanoparticle surface, and the XAS data discussed in the following will help to shed light on the local structure of silver atoms and ions also in the NR bulk.

NEXAFS spectra collected on AuNRs-AA and AuNRs-HQ in the solid state, prepared in the same manner as the samples for SR-XPS measurements, are reported in Fig. 5. The spectra of the two samples are very similar, therefore they will be discussed together. In the C K-edge spectra of AuNRs-AA and AuNRs-HQ, reported in Fig. 4a and c, respectively, several resonances due to electronic transitions from the C 1s core level to antibonding molecular orbitals can be observed, and assigned by comparison with literature data as follows: the sharp feature at about 288.7 eV is associated with the C 1s $\rightarrow \pi^*$ transition of C=O molecular orbital, the shoulder at about 288 eV to a σ^* resonance by the C-H groups and Rydberg features, additional features around 293 and 303 eV can be assigned to 1s $\rightarrow \sigma^*$ transitions by C-C and C=O molecular groups, respectively.²⁰ The pre-edge peak at 285.1 eV in the spectrum of AuNRs-HQ and appearing as a shoulder in the spectrum of AuNRs-AA is probably an artifact related to carbon contamination of the beamline mirrors. In N K-edge spectra (Fig. 5b for AuNRs-AA, Fig. 5d for AuNRs-HQ) only one sharp peak appears at 402 eV, assigned to the N 1s $\rightarrow \sigma^*$ transition of positively charged nitrogen as expected from the CTAB molecular structure, and coherently with SR-XPS findings. The low intensity peaks around 395 eV in Fig. 5d are spurious peaks related to problems with noise and background subtraction for the low intensity N K edge signal.

3.3. Local structure around Au and Ag: XAS analysis

The XAS spectra of gold and silver were analysed in both XANES and EXAFS regions, which provide complementary information on the local ordering around the average absorber. The XANES features provide information on the average valence state of the

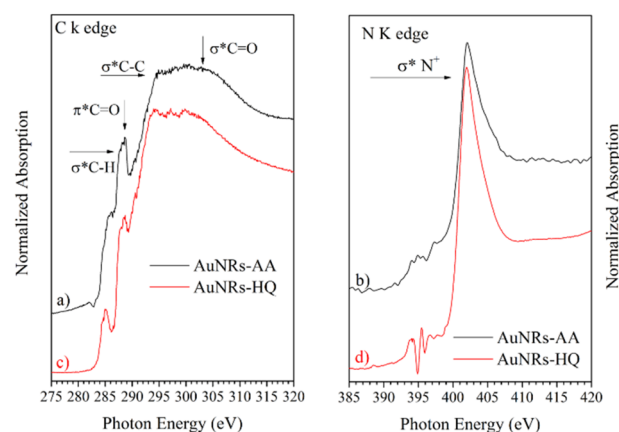


Fig. 5 Carbon K-edge and nitrogen K-edge spectra of AuNRs-AA (a, b) and of AuNRs-HQ (c, d).

absorber, the density of empty states near the Fermi level, and the symmetry of the coordination environment.³⁶ In the near edge region, the effect of structural disorder is smaller than in the extended region; therefore, the analysis of the XANES region is particularly suitable for revealing weaker electronic and/or structural differences between samples. Analysis of the EXAFS data provides further details on the average atomic coordination around the absorber in terms of the average distance, number, and mean squared relative displacement (MSRD) of the neighbouring shells.²³

3.3.1. XANES analysis. The normalized Au-L_{III} XANES of the AuNRs-AA, AuNRs-HQ and Au reference foil are presented in Fig. 6a. All XANES spectra of the samples are like that of the Au reference foil. This suggests that the larger fraction of Au in the samples is in the bulk-like environment, as expected for relatively large nanoparticles with low surface-to-volume ratios.

The first shoulder in the Au-L_{III} XANES at about 11 920 eV is commonly referred to as the “white line” of gold and is related to the photoelectron transitions from the 2p_{3/2} state to the 5d state, the intensity of the white line being proportional to the density of free states in the 5d valence state.³⁷ The very similar shape of the XANES measured on AuNRs and on Au reference foil suggests that the larger fraction of Au consists of zero-valent Au⁰ atoms located in the inner volume of the NR, with negligible

fraction of Au atoms located at the NR surface. This is consistent with relatively large nanostructures having a sharp surface.

Looking at the Ag-K edge (Fig. 6a) differences between the XANES measured on AuNR samples and on silver reference foil become apparent indicating that the Ag coordination chemistry in these samples is definitively different from metallic silver. In AuNPs, the rise of the Ag K edge XANES signal around 25 520 eV is similar to what was observed in Ag⁺ composites (*i.e.*, AgNO₃ or AgCl) so it can be attributed to the rise of the Ag valence state³⁸ in our samples. Notice that the AgO XANES features are definitively different so the presence of Ag²⁺ can be excluded.

3.3.2. EXAFS analysis. The Fourier transform moduli (|FT|) of the EXAFS k^w -weighted signals for the analyzed spectra are shown in Fig. 6b. The |FT| gives a more intuitive picture of the average local atomic structure around the absorbers, where the |FT| features represent the pseudo radial distribution function around the absorber minus the correction for the phase shift of the photoelectrons (which is about -0.5 \AA) and the effect of the photoelectron scattering amplitude (which affects the |FT| peak shape). The local structure around Au in AuNR samples is like that of Au foil, with the lower intensity of |FT| related to the expected greater structural disorder in nanostructured samples. In contrast, the local structure around Ag in AuNRs is definitively different from that of Ag foil according to what was observed in the XANES region. To obtain quantitative details, EXAFS data analysis was performed using a multiple-shell data refinement procedure.

The quantitative analysis of Au L_{III}-edge EXAFS spectra has been carried out selecting the relevant contribution from the Au-fcc bulk structure (crystallographic information file from the crystallography open database, COD #9008463 (ref. 42)), the $k^2\chi(k)$ experimental data and best fit curves for AuNR samples are shown in Fig. 6c, the single (SS) and multiple (MS) shell contributions used for the analysis and the residues (experimental data minus best fit curves) are shown for the sake of comparison.

To select the relevant contribution and reduce the correlations among the fitting parameters we applied constraints based on the Au crystallographic structure.⁴³ After an initial trial and error procedure 4 single scattering (SS) and 3 multiple scattering (MS) were found as a suitable set to refine the Au EXAFS experimental data. The paths used in the EXAFS fit describe the coordination shells of metallic gold up to the 4th neighboring atoms (around 5.8 Å). Considering the absorber at the unit cell cube, the 4 single scattering paths correspond, considering the absorber at the unit cell cube edge, to 12 Au₁ nearest neighbours at the center of the fcc face (SS₁), 6 second neighbours Au₂ along the fcc cube edge (SS₂), 24 third neighbours Au₃ on the opposite fcc face center (SS₃), and 12 fourth neighbours Au₄ along the fcc face diagonal (SS₄). The MS contributions correspond to the triple scattering (MS₁: Au-Au₁-Au₁) between two first neighbors as well as the triple (MS₂: Au-Au₁-Au₄) and quadruple (MS₃: Au-Au₁-Au₄-Au₁) scattering paths along the fcc face diagonal. We fixed the multiplicity numbers of the SS and MS contributions to crystallographic values, the coordination lengths of the Au-Au bond (R_{SS_i}) were left free, the coordination lengths of the next neighbour shells were constrained to a single free parameter $a = R_{SS_2}$ (the edge of

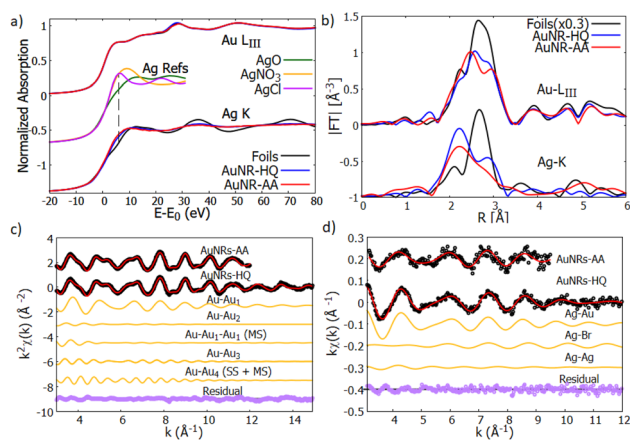


Fig. 6 (a) Normalized XANES spectra of Au-L_{III} (top curves) and Ag-K edge (bottom curves) measured on reference foils (Au and Ag, black lines) and on AuNR samples (AuNR-HQ: blue and AuNR-AA: red). The reference spectra of Ag⁺ (AgNO₃ and AgCl) and Ag²⁺ (AgO) are shown for the sake of comparison (from ref. 39–41). (b) Moduli of the Fourier transforms |FT| of k^w -weighted EXAFS spectra of Au-L_{III} (top curves) and Ag-K (bottom curves) edges for the reference foils (black lines) and AuNRs (AuNR-HQ: blue, AuNR-AA: red). (c) Results of the Au EXAFS data fitting for AuNR-HQ and AuNR-AA samples. The experimental (black dots), best fit (red lines) and the partial contributions used for the analysis (orange lines) are shown (vertically shifted for clarity). The bottom curves (purple dots) represent the residues (experimental data minus best fit). (d) Results of the Ag-K edge EXAFS fitting for AuNR-HQ and AuNR-AA samples. The black dots represent the experimental data, the red line the best fit, the orange ones represent the contribution of every path to the final fit and the purple dots represent the residual. All plots are shifted vertically for the sake of clarity. The larger noise on AuNR-AA forces restriction of the range for the analysis (see text).

the fcc unit cell): ($R_{SS_3} = a\sqrt{3}/2$, $R_{SS_4} = a\sqrt{2}$). The MSRD (σ_i^2) were refined independently, only we constrained $\sigma_{MS_3}^2 = (\sigma_{SS_3}^2 + \sigma_{MS_3}^2)/2$. The same energy shift ($\Delta E = 6.8$ eV) and the same $S_0^2 = 0.9$ (representing the passive electron reduction factor) were applied to all the paths and dataset. Table 1 resumes the best fit parameters.

The values obtained from the Au EXAFS analysis agree well with the Au bulk structure, with a weak (1%) compression of the lattice parameter; we found $a = R_{SS_2} = 4.01(2)$ Å with respect to the gold lattice parameter $a_{Au} = 4.065$ (Å), which can be attributed to thermal compression, since the measurements were performed at LN temperature.

The Ag EXAFS (Fig. 6d) spectra have larger noise with respect to the Au ones, this forces reduction of the k -range of the analysis. To individuate the suitable contributions we proceeded by trial and error selecting contributions from a shortlist of possible Ag local coordination structures, based on the sample synthesis and composition: Ag–metal, Ag–Au amalgams, and Ag–bromine (of the ionic head of CTAB). The preliminary attempts allow exclusion of a sizable amount of Ag–O contribution. Model structures used to calculate the photoelectron amplitude and phase functions were taken from structures in the Crystallography Open Database.^{44,45} After some testing S_0^2 was fixed at 0.9 for both spectra and the energy scale shift ΔE_0 , which was calculated in the analysis with the less noisy spectra (AuNP-HQ), was fixed for the analysis of the sample AuNP-AA.

The AuNR-HQ spectrum has a better signal-to-noise ratio. We found that a satisfactory best fit is obtained using a first shell Ag–Au, around 2.83(2) Å with multiplicity close to 5, which is much less than 12, expected for Ag dispersed in an Au matrix. We found two other contributions relevant for the analysis, namely Ag–Br and Ag–Ag. Due to the weakness of these contributions, the antiphase effect between Ag–Au and Ag–Br contributions and the known correlation between multiplicity number and MSRD, the obtained values are affected by quite large uncertainty. However, they demonstrate Ag bonded either to Au or Br. Assuming crisp cylindrical surfaces, we may calculate the NR's surface-to-volume ratio (S/V) at roughly 5% for AuNR-HQ and slightly less, around 4% for AuNR-AA

Table 1 The table presents all significant parameter results of the fitting of the EXAFS Au K-edge spectra of AuNR-HQ and AuNR-AA. Here R^2 is the square residual factor, N the multiplicity factor, R the interatomic distance and σ^2 the Debye–Waller factor. The symbol * highlights the constrained values. The passive electron reduction factor $S_0^2 = 0.9$ and the energy scale shift $\Delta E_0 = 6.85$ eV were kept fixed

Paths	AuNR-HQ			AuNR-AA		
	N	R (Å)	$\sigma^2 (\times 10^{-2} \text{ \AA}^{-2})$	N	R (Å)	$\sigma^2 (\times 10^{-2} \text{ \AA}^{-2})$
SS ₁	12*	2.85(1)	0.93(1)	12*	2.85(1)	0.83(1)
SS ₂	6*	4.01(2)	1.87(3)	6*	3.97(1)	2.35(2)
MS ₁	48*	4.32(2)	0.36(5)	48*	4.32(1)	0.11(5)
SS ₃	24*	4.98*	1.34(4)	24*	4.99(2)	1.35(2)
SS ₄	12*	5.76*	1.25(2)	12*	5.73(4)	2.96(3)
MS ₂	24*	5.76*	1.33*	24*	5.73(4)	2.34(2)
MS ₃	12*	5.76*	1.40(6)	12*	5.73(4)	1.71(5)

Table 2 The table presents all significant parameter results of the fitting of the EXAFS Ag K-edge spectra of AuNR-HQ and AuNR-AA, where N is the multiplicity factor, R the interatomic distance and σ^2 the Debye–Waller factor. The symbol * highlights the constrained values. The passive electron reduction factor $S_0^2 = 0.85$ and the energy scale shift $\Delta E_0 = 0.8$ eV were kept fixed

Paths	AuNR-HQ			AuNR-AA		
	N	R (Å)	$\sigma^2 (\times 10^{-2} \text{ \AA}^{-2})$	N	R (Å)	$\sigma^2 (\times 10^{-2} \text{ \AA}^{-2})$
Ag–Au	5.0(5)	2.83(1)	1.2(2)	4.1(6)	2.82(1)	1.8(2)
Ag–Br	0.5(1)	2.63(2)	0.5*	0.5(2)	2.69(3)	0.29(3)
Ag–Ag	0.7(1)	2.81(2)	1.5*	1.3(3)	3.03(4)	1.5*

samples; however, surface flaws and roughness may result in higher actual S/V values for the samples. Given the total Ag fraction in the samples (around 3% from EDX analysis) the EXAFS results indicate that Ag atoms likely form an Ag–Au layer on the NR surface (Ag–Au and Ag–Ag neighbours), some of the Ag atoms connecting Br from one side and the NR inner Au atoms from the other side. The hypothesis that a fraction of unreacted Ag binds only to the amine group in the CTAB head can be excluded because the washing procedure would have removed that fraction, leaving therefore only Ag bound to the gold nanorods. The EXAFS data on the AuNR-AA sample are much noisier and the results are surely less accurate, but they generally confirm the trend observed on AuNR-AA with a slightly higher fraction of Ag–Ag neighbours, which is consistent with smaller S/V in this sample. The results of the Ag–K edge EXAFS data analysis are shown in Table 2.

4. Conclusions

The investigation of the morphology, molecular, electronic structure and local structure around silver and gold atoms reported in this work sheds light on the correlation between the secondary ligand used in the synthesis path, *i.e.*, HQ or AA, and the optical properties displayed by two sets of AuNRs. The here investigated AuNRs-AA and AuNRs-HQ differ in their plasmon band position, morphology and aspect ratio in a reproducible way, and our multi-technique investigation evidenced the influence of the different secondary ligands on the anisotropic growth process, leading to the conclusion that the moderate reducing agent AA induces the formation of AuNRs of lower size ratio, while HQ allows obtaining longer rods. The destiny of silver atoms and ions was also deeply investigated, and the analogous presence of Ag(I) and Ag(0) species on the two sets of AuNR surfaces revealed by SR-XPS and XAS suggested that the observed different size ratio of AuNRs-AA and AuNRs-HQ nanosystems is induced by the secondary ligand only.

Author contributions

C. B.: conceptualization; C. B., C. M., S. A., A. L.: data curation, formal analysis; C. B., C. M., I. V., G. I.: funding acquisition; C.

B., C. M., A. L., S. A., G. I., I. K., A. S., A. C., M. C., S. M.: investigation, methodology, validation, visualization; C. B., A. L., S. A.: writing – original draft; C. B., G. I., C. M., I. V.: writing – review & editing.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- M. A. El-Sayed, Some interesting properties of metals confined in time and nanometer space of different shapes, *Acc. Chem. Res.*, 2001, **34**(4), 257e64.
- D. Maccora, V. Dini, C. Battocchio, I. Fratoddi, A. Cartoni, D. Rotili, M. Castagnola, R. Faccini, I. Bruno, T. Scotognella, A. Giordano and I. Venditti, Gold nanoparticles and nanorods in nuclear medicine: a mini review, *Appl. Sci.*, 2019, **9**(16), 3232.
- M. Grzelczak, J. Vermant, E. M. Furst and L. M. Liz-Marzan, Directed self-assembly of nanoparticles, *ACS Nano*, 2010, **4**(7), 3591e605.
- G. Pasparakis, Recent developments in the use of gold and silver nanoparticles in biomedicine, *Wiley Interdiscip. Rev.: Nanomed. Nanobiotechnol.*, 2022, **14**(5), e1817.
- L. M. Rossi, J. L. Fiorio, M. A. S. Garcia and C. P. Ferraz, The role and fate of capping ligands in colloiddally prepared metal nanoparticle catalysts, *Dalton Trans.*, 2018, **47**(17), 5889e915.
- H. Yin, N. Li, Y. Si, H. Zhang, B. Yang and J. Wang, Gold nanonails for surface-enhanced infrared absorption, *Nanoscale Horiz.*, 2020, **5**, 1200–1212.
- N. Garg, C. Scholl, A. Mohanty and R. Jin, The role of bromide ions in seeding growth of Au nanorods, *Langmuir*, 2010, **26**(12), 10271–10276.
- S. Eustis and M. El-Sayed, Aspect ratio dependence of the enhanced fluorescence intensity of gold nanorods: experimental and simulation study, *J. Phys. Chem. B*, 2005, **109**(34), 16350–16356.
- P. K. Jain, X. H. Huang, I. H. El-Sayed and M. A. El-Sayed, Noble metals on the nanoscale: optical and photothermal properties and some applications in imaging, sensing, biology, and medicine, *Acc. Chem. Res.*, 2008, **41**(12), 1578e86.
- C. J. Murphy, L. B. Thompson, A. M. Alkilany, P. N. Sisco, S. P. Boulos, S. T. Sivapalan, *et al.*, The many faces of gold nanorods, *J. Phys. Chem. Lett.*, 2010, **1**(19), 2867e75.
- N. R. Jana, L. Gearheart and C. J. Murphy, *J. Phys. Chem. B*, 2001, **105**, 4065–4067.
- D. K. Smith and B. A. Korgel, The importance of the CTAB ligand on the colloidal seed-mediated synthesis of gold nanorods, *Langmuir*, 2008, **24**(3), 644–649.
- N. Garg, C. Scholl, A. Mohanty and R. Jin, The role of bromide ions in seeding growth of Au nanorods, *Langmuir*, 2010, **26**(12), 10271–10276.
- Y. Xu, L. Chen, X. Ye, X. Wang, J. Yu, Y. Zhao, M. Cao, Z. Xia, B. Sun and Q. Zhang, Cooperative interactions among CTA⁺, Br[−] and Ag⁺ during seeded growth of gold nanorods, *Nano Res.*, 2017, **10**(6), 2146–2155.
- F. Hubert, F. Testard and O. Spalla, Cetyltrimethylammonium bromide silver bromide complex as the capping agent of gold nanorods, *Langmuir*, 2008, **24**, 9219–9222.
- S. R. Jackson, J. R. McBride, S. J. Rosenthal and D. W. Wright, Where's the silver? Imaging trace silver coverage on the surface of gold nanorods, *J. Am. Chem. Soc.*, 2014, **136**, 5261–5263.
- C. Oliveira, C. Ribeiro Chaves, P. Bargiela, M. da Grac, C. da Rocha, A. Ferreira da Silva Jos, F. Diniz Chubaci, M. Boström, C. Persson and M. Malta, Surface studies of the chemical environment in gold nanorods supported by X-ray photoelectron spectroscopy (XPS) and *ab initio* calculations, *J. Mater. Res. Technol.*, 2021, **15**, 768–776.
- S. E. Lohse and C. J. Murphy, The quest for shape control: a history of gold nanorod synthesis, *Chem. Mater.*, 2013, **25**(8), 1250–1261.
- L. Smykalla, P. Shukryna, M. Korb, H. Lang and M. Hietschold, Surface-confined 2D polymerization of a brominated copper-tetraphenylporphyrin on Au(111), *Nanoscale*, 2015, **7**, 4234–4241.
- A. Di Cicco, G. Aquilanti, M. Minicucci, E. Principi, N. Novello, A. Cognigni and L. Olivi, Novel XAFS capabilities at Elettra synchrotron light source, *J. Phys.: Conf. Ser.*, 2009, **190**, 012043.
- F. d'Acapito, G. O. Lepore, A. Puri, A. Laloni, F. La Manna, E. Dettona, A. De Luisa and A. Martin, *J. Synchrotron Radiat.*, 2019, **26**, 551–558.
- C. Meneghini, F. Bardelli and S. Mobilio, ESTRA-fitEXA: a software package for EXAFS data analysis, *Nucl. Instrum. Methods Phys. Res., Sect. B*, 2012, **285**, 153–157.
- G. Bunker, *Introduction to XAFS: A Practical Guide to X-Ray Absorption Fine Structure Spectroscopy*, Cambridge University Press, Cambridge, 2010.
- D. E. Sayers, E. A. Stern and F. W. Lytle, New technique for investigating noncrystalline structures: fourier analysis of the extended X-ray absorption fine structure, *Phys. Rev. Lett.*, 1971, **27**, 1204–1207.

- 25 A. L. Ankudinov, B. Ravel, J. J. Rehr and S. D. Conradson, Real space multiple scattering calculation of xanes, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1998, **58**, 7565.
- 26 V. Secchi, S. Franchi, D. Ciccarelli, M. Dettin, A. Zamuner, A. Serio, G. Iucci and C. Battocchio, Biofunctionalization of TiO₂ surfaces with self-assembling layers of oligopeptides covalently grafted to chitosan, *ACS Biomater. Sci. Eng.*, 2019, **5**, 2190–2199.
- 27 S. D. Techane, L. J. Gamble and D. G. Castner, X-ray photoelectron spectroscopy characterization of gold nanoparticles functionalized with amine-terminated alkanethiols, *Biointerphases*, 2011, **6**, 98.
- 28 D. Kwaśniewska and J. Kiewlicz, Study of interaction between cationic ligand (CTAB) and ascorbic acid/ascorbic acids derivatives by tensiometric and spectroscopic methods, *J. Mol. Liq.*, 2022, **354**, 118917.
- 29 I. Haidar, A. Day, P. Decorse, S. Lau-Truong, A. Chevillot-Biraud, J. Aubard, N. Felidj and L. Boubekeur-Lecaque, Tailoring the shape of anisotropic core-shell Au-Ag nanoparticles in dimethyl sulfoxide, *Chem. Mater.*, 2019, **31**(8), 2741–2749.
- 30 S. Jin Heo, S. Yoon, S. H. Oh and H. J. Kim, Modification of hybrid active bilayer for enhanced efficiency and stability in planar heterojunction colloidal quantum dot photovoltaics, *Nanoscale Res. Lett.*, 2013, **8**, 488.
- 31 A. G. Shard, Detection limits in XPS for more than 6000 binary systems using Al and Mg K α X-rays, *Surf. Interface Anal.*, 2014, **46**, 175.
- 32 I. Berlanga, Synthesis of non-uniform functionalized amphiphilic block copolymers and giant vesicles in the presence of the Belousov-Zhabotinsky reaction, *Biomolecules*, 2019, **9**, 352.
- 33 F. C. Loh, K. L. Tan and E. T. Kang, XPS studies of charge transfer interactions in some poly(N-vinylcarbazole)/acceptor complexes, *Eur. Polym. J.*, 1991, **27**(10), 1055.
- 34 I. Venditti, G. Iucci, I. Fratoddi, M. Cipolletti, E. Montalesi, M. Marino, V. Secchi and C. Battocchio, Direct conjugation of resveratrol on hydrophilic gold nanoparticles: structural and cytotoxic studies for biomedical applications, *Nanomaterials*, 2020, **10**, 1898.
- 35 G. Beamson and D. Briggs, *High Resolution XPS of Organic Polymers: The Scienta ESCA 300 Database*, John Wiley & Sons, Chichester, U.K., 1992.
- 36 M. Benfatto and C. Meneghini, A close look into the low energy region of the XAS spectra: the XANES region, in *Synchrotron Radiation*, ed. S. Mobilio, F. Boscherini and C. Meneghini, Springer, Berlin, Heidelberg, 2015.
- 37 C. Nayak, D. Bhattacharyya, K. Bhattacharyya, A. K. Tripathi, R. D. Bapat, S. N. Jha and N. K. Sahoo, Insight into growth of Au-Pt bimetallic nanoparticles: an *in situ* XAS study, *J. Synchrotron Radiat.*, 2017, **24**(Pt 4), 825–835.
- 38 B. A. Kyffin, D. M. Pickup, G. Mountjoy, F. Foroutan, I. Abrahams and D. Carta, Atomic-scale structural characterization of silver-doped phosphate-based glasses prepared by coacervation, *J. Phys. Chem. C*, 2021, **125**(22), 12256–12268.
- 39 E. A. Pradas del Real and G. Sarret, *Ag K Edge XAS Fluorescence of Frozen AgNO₃ Solution*, SSHADE/FAME (OSUG Data Center), 2013, Dataset/Spectral Data, DOI: [10.26302/SSHADE/EXPERIMENT_GS_20170713_007](https://doi.org/10.26302/SSHADE/EXPERIMENT_GS_20170713_007).
- 40 E. A. Pradas del Real and G. Sarret, *Ag K Edge XAS Transmission of AgCl*, SSHADE/FAME (OSUG Data Center), 2014, Dataset/Spectral Data, DOI: [10.26302/SSHADE/EXPERIMENT_GS_20170713_002](https://doi.org/10.26302/SSHADE/EXPERIMENT_GS_20170713_002).
- 41 E. A. Pradas del Real and G. Sarret, *Ag K Edge XAS Transmission of AgO*, SSHADE/FAME (OSUG Data Center), 2013, Dataset/Spectral Data, DOI: [10.26302/SSHADE/EXPERIMENT_GS_20170712_002](https://doi.org/10.26302/SSHADE/EXPERIMENT_GS_20170712_002).
- 42 R. W. G. Wyckoff, Second edition. Interscience Publishers, New York, Cubic closest packed, ccp, structure, *Cryst. Struct.*, 1963, 7–83.
- 43 C. Battocchio, C. Meneghini, I. Fratoddi, I. Venditti, M. V. Russo, G. Aquilanti, C. Maurizio, F. Bondino, R. Matassa, M. Rossi, S. Mobilio and G. Polzonetti, Silver nanoparticles stabilized with thiols: a close look at the local chemistry and chemical structure, *J. Phys. Chem. C*, 2012, **116**(36), 19571–19578.
- 44 J. Spreadborough and J. W. Christian, High-temperature X-ray diffractometer, *J. Sci. Instrum.*, 1959, **36**, 116–118.
- 45 H. Takahashi, S. Tamaki and S. Sato, Electron density distribution in Ag Br, *J. Phys. Soc. Jpn.*, 1987, **56**, 3593–3597.