Gold nanonetwork film on the ITO surface exhibiting one-dimensional optical properties

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Abstract

A network of gold nanostructures exhibiting one-dimensional gold nanostructure properties may become a prospective novel structure for optical, electrical and catalytic applications benefited by its unusual characteristics resulting from the collective properties of individual nanostructures in the network. In this paper, we demonstrate a facile method for the formation of high-density gold nanonetwork film on the substrate surface composed of quasi-1D nanoparticles (typically fusiform) with length ca. 10 nm - via reduction of gold ions in the presence of nanoseeds attached surface, binary surfactants of cetyltrimethylammonium bromide and hexamethyleneteramine and Ag+ ions. The length of the nanonetworks can be up to ca. 100 nm, which corresponds to the aspect ratio of ca. 10. The quasi-1D gold nanostructures as well as the nanonetworks were found to be sensitive to the binary surfactants system and the Ag+ ions as they can only be formed if all the chemicals are available in the reaction. The nanonetworks exhibit unique 1D optical properties with the presence of transverse and longitudinal surface plasmon resonance absorption. Owing to their peculiar structures that are composed of small quasi-1D nanoparticles, the nanonetworks may produce unusual optical and catalytic properties, which are potentially used in surface-enhanced Raman scattering, catalysis and optical and non-linear optical applications.

Keywords: Nanonetworks, Quasi-1D gold nanoparticles, Seed-mediated growth, 1D nanostructures

Background

One-dimensional gold nanostructures, such as nanorods, nanowires, etc., have continued to be the objective of the research in the controlled-shape synthesis nowadays due to its notable catalytic, electrical, magnetic and optical characteristics [1-4]. Their optical absorption properties, in particular, are characterised by the presence of two unique strong localised-surface plasmon resonance bands near the 500 (transverse mode) and 650 nm (longitudinal mode) but tunable within the region of ca. 400 to 550 nm and of ca. 600 to 800 nm, respectively. This enables them to produce peculiar optical properties for use in a broad range of applications including nanophotonics [6-11], light-emitting diode [12], photovoltaics [13-17] and photosynthesis efficiency enhancement [18,19] and optical sensing applications [20]. For example, nanorods demonstrate a unique profile and enhanced energy of electron-wave scattering upon exposure to the electromagnetic irradiation resulting from their anisotropic localised-plasmonic excitation character [21,22], placing them as a potential agent for photodynamic therapy of cancer cells [19,23-26], optical contrast agent and for diagnostic imaging applications [20,27,28]. As gold itself has high compatibility to a wide range of biomolecules and its peculiar surface morphology is bounded normally by highly-energetic planes ([110] and [100]) [1,3,29-31], these nanostructures become an attractive candidates for catalysts in bio-organic reactions [32,33] and chemical sensing applications [20,23].

A wide range of methods are currently available for the preparation of gold nanorods with controlled shape, yield and aspect ratios in the solution. The synthetic procedure as well as the mechanism of the formation has also been well-summarised in many recent reports [1,3,5,29]. To mention a few examples, cetyltrimethylammonium bromide (CTAB) surfactant-assisted seed growth method by Murphy et al. appeared as a versatile approach for the synthesis of gold nanorods with a controlled aspect ratio in the solution [5,29,34]. A much...
improved technique was later demonstrated by El-Sayed and co-workers, in which, by modifying the seeding and adding of Ag⁺ into the growth solutions that was used by Murphy, nanorods with a yield of nearly 100 % could be achieved [35]. Effective facet-selective gold deposition process in the presence of silver ions in the growth solution was considered as the key driving factor for the high-yield gold nanorod formation. However, in contrast to such outstanding achievement in solution phase, their growth directly on the surface has not yet been developed. Since many applications, such as optoelectronic, catalyst and surface-enhanced Raman scattering (SERS), required that they are attached onto the surface, the technique for growing the nanorods directly on the surface should be developed. It is true that the solution phase prepared gold nanorods may be transferred onto the surface via, for example, Langmuir-Blodgett [36] or drop and spin casting assemblies [37]; however, they are likely intricate and inadvertently expose the nanostructures to possible properties disruption during the process. Therefore, to develop a method that enables the shape-controlled growth directly on the surface is crucial.

Recently, we have grown metal nanoparticles directly on the surface using a seed-mediated growth method [38-40]. Gold nanoparticles with a variety of shapes, such as nanoplates, nanocubes, etc., have been successfully prepared directly on the surface [41,42]. As for 1D morphology, particularly in nanorods, considerable efforts have been devoted by us to synthesise them directly on the surface. Unfortunately, so far their yield was relatively low so that the unique 1D optical properties that are indicated by the presence of two separate bands, i.e. transverse and longitudinal SPRs, could not be detected on their optical absorption spectra. Actually, we have also attempted many approaches to improve the yield including the use of the well-known growth solutions for gold nanorods such as prescribed by Murphy's [3,29,34,43], El-Sayed’s [35] and Korgel’s [44] groups combined with our method. Unfortunately, again, these attempts failed. This could be the result of unique growth characteristic of gold nanostructures on the surface, which might be influenced by many factors including substrate surface-nanoseeds interaction effect, substrate surface-solution interface characteristics (e.g. wet-ability), and the nature of the precursors transport to the growing crystallite on the surface, which produce a different crystal growth compared to that of in the solution. We note that the formation of 1D gold nanoparticles directly on the surface is a remarkably challenging task, however, since the anisotropic effect of the nanostructure in the applications are very special and even superior in many aspects over other morphologies, the synthesis of gold nanoparticles with alternative morphology directly on the surface but producing properties that exhibit those in one-dimensional nanostructures is of particular interest.

Here, we report the synthesis of high-density networked gold nanostructures directly on the surface exhibiting unique one-dimensional nanoparticle characters. By combining a special effect of AgNO₃ in the formation of gold nanorods to our new growth solution that contains cetyltrimethylammonium bromide and hexamethylenetetramine using a seed-mediated growth method, high-yield networked gold nanostructures can be realised directly on the surface. Optical absorption spectrum analysis on the sample surprisingly revealed that the nanostructures exhibit one-dimensional properties, reflected by the presence of two plasmonic bands on their spectrum. Therefore, they could become a replica of gold nanorods on the surface to be used in SERS, catalysis and plasmonics enhanced optoelectronics devices.

**Methods**

Networked gold nanoparticles were grown directly on the surface via our seed-mediated growth method. This technique comprises a two-step process, namely seeding and growth processes. In typical procedure, the seeding process was carried out via *in situ* reduction of gold salt on the substrate sample [45]. Briefly, a clean ITO substrate (surface resistance ca. 30 Ω/sq, CBC Ings Co. Ltd., Tokyo, Japan) was immersed into a solution that contains 0.25 mM HAuCl₄ and 0.25 mM trisodium citrate. The substrate was kept undisturbed in the solution at 28°C for 1 h. After that, 0.5 mL of 0.1 M ice-cooled NaBH₄ was added to the solution, and immediately, the gold nanoseeds formed in the solution as well as on the surface. The sample was then kept in the solution for another hour. After that, the sample was removed, rinsed with pure water and dried with a flow of nitrogen gas. Using this approach, high-density gold nanoseeds will be formed on the surface (see Figure 1A).

The growth process was carried out by immersing the substrate that has been treated with the nanoseeds into a growth solution that contains 0.5 mL of 0.01 M of HAuCl₄, 8 mL of 0.1 M CTAB, 12 mL of 0.1 M hexamethylenetetramine, 0.3 mL of 0.1 M ascorbic acid and 40 μL of 0.01 M of AgNO₃. The sample was kept undisturbed for 4 h at 28°C for the growth process. If this condition is used, the final concentration of each reagent is 0.25, 40, 60 and 1.5 mM and 20 μM for HAuCl₄, CTAB, hexamethylenetetramine (HMT), ascorbic acid and AgNO₃, respectively. Finally, the sample was removed, rinsed with plenty of pure water and dried with a flow of nitrogen gas. The growth solution used in the present study was a new growth solution developed by our group of which was the modification of our standard solution for preparing gold nanotripod in solution [46] as our initial expectation to also realise the
formation of nanotripods on the surface if using similar growth solution. There are several modifications that have been made to the standard solution, namely the AgNO₃ was new here and the NaOH is not required for the present study as it was in the preparation of gold nanotripod.

The morphology of the gold nanostructures grown on ITO surfaces was characterised using a field-emission scanning electron microscopy (FESEM) (JSM-7400 F, JEOL Ltd., Akishima, Tokyo, Japan). The optical absorption spectrum of the samples was obtained using Perkin Elmer Lambda 900 UV/VIS/NIR spectrometer (Waltham, MA, USA).

**Results and discussion**

We noted again here that the growth solution used in this project was actually a modification to the solution that was originally used to prepare gold nanotripods in solution phase [46]. Prior to modifying the solution, we actually have used such original solution to grow the attached-nanoseeds on the surface via a seed-mediated growth method. Actually, we expected that similar morphology, i.e. nanotripods, would be realised on the surface. Unfortunately, neither nanotripods nor nanorods were obtained but instead spherical gold nanoparticles were formed, reflecting unusual heterogeneous gold deposition on the nanoseed surface emerged as the results of the surface effect. Thus, shape-controlled growth on the surface often yields limited success.

The original growth solution for the nanotripods contained two special surfactants, namely CTAB and HMT. As an attempt for obtaining 1D gold nanoparticle growth from the nanoseeds on the surface and being inspired by the fact that the AgNO₃ promotes the formation of nanorods in solution [35], we added a minute amount of AgNO₃ into the original solution. After being immersed for 4 h in the growth solution, a purple-blue colour was formed on the surface. Normally, this kind of colour only appears on the gold nanostructures with one-dimensional morphology, such as nanorods or nanowires [3,29,35]. Therefore, this result signifies that one-dimensional morphology of gold nanoparticles might have formed on the surface. Optical absorption spectrum collected from the samples surprisingly showed the presence of two plasmonic bands at 520 and 680 nm. In agreement with the observed purple-blue colour, two-banded absorption spectrum is also an indication of the formation of one-dimensional gold nanostructures on the surface. We then carried out a FESEM characterisation on the as-prepared samples.

![Figure 1 FESEM images and high magnification of networked gold nanoparticles and optical absorption spectra of gold nanostructures.](image)
surprise, only high density networked string-like structures of gold nanoparticles were formed on the surface covering the entirety of the substrate surface instead of nanorods or other 1D morphologies, to which such unique two plasmonic characters can be related to. The optical absorption and the FESEM results are shown in Figure 1. However, as revealed in the high-resolution FESEM image of the samples (see Figure 1C), actually the networked structures were mostly composed of quasi-1D nanoparticles (e.g. fusiform) with size (length) approximately 10 nm that aligned side-by-side with each other. The length of the nanonetworks can be up to ca. 100 nm, which corresponds to the aspect ratio of ca. 10. Despite the fact that the spherical nanoparticles also formed the networked structures, the network is normally relatively shorter than those composed of the quasi-1D nanoparticles. Even in most cases, they seemed to be an aggregate instead of a long-range networked morphology. However, no nanorods, nanowires or other 1D nanostructures were observed on the image, confirming that one-dimensional optical properties are solely produced by the networked-gold nanoparticles on the surface. As revealed in the image, the quasi-1D structures that composed the networked structure are considerably small with average length and diameter in the range of ca. 2 to 3 nm and 10 to 15 nm, correspondingly. Such small 1D nanostructures might have prospective use in catalysis and sensing applications such as the possibility of producing peculiar properties as the results of anisotropic morphology and quantum effect.

As evident in Figure 1A,B,C, no perfect nanorods or nanowires were obtained on the surface. Therefore, the presence of one-dimensional optical properties could be directly related to the networked structure (nanonetworks) that formed linear chains on the surface. Thus, the transverse surface plasmon band, the shorter wavelength band, can be easily attributed to the oscillation of the free electron system toward the short axis of the nanonetworks. Meanwhile, the longitudinal plasmonic band, i.e. the longer band, was related to a long-range plasmonic coupling [47] amongst the nanoparticles in the networked structure (see Figure 1E). Actually, the appearance of a longitudinal plasmonic band from the networked nanoparticles here is quite unusual and is predicted as resulting from the unique 1D morphology of the individual nanoparticles in the nanonetworks. According to our earlier results [38-40], in most cases, such unique 1D plasmonic characteristics were not observable if the network was composed of spherical nanoparticles. Therefore, the appearance of LSPR in these nanonetworks in the present study indicates a peculiar dependence of long-range plasmonic coupling on shape at this length-scale regime [47]. Thus, more unusual properties will then be expected to arise by the quasi-1D but assembled in a network structure.

As has been noted earlier, the AgNO₃ might have played a substantial role behind the formation of quasi-1D gold nanoparticles, which is the basis for the nanonetwork structure. To obtain a detailed understanding on its role in this process, we examined the growth characteristic of gold nanostructures on the surface in the presence of several AgNO₃ concentrations, namely 10, 20, 30, 40 and 50 μM, via UV-vis absorption spectroscopy and FESEM imaging. Figure 2 is the related optical absorption spectra of the samples. As judged from the curve a, neither quasi-1D nor nanonetworks were formed when the AgNO₃ was absent in the reaction. However, 1D gold nanostructures might have effectively produced when the AgNO₃ as low as 10 μM was used as the appearance of two plasmonics bands at ca. 520 (TSPR) and 630 nm (LSPR) (see curve b). The quasi-1D nanostructures as well as nanonetworks formation might be optimum when the AgNO₃ concentration used was 40 μM. It is indicated by a maximum shifting of the LSPR to red, reflecting the improvement of the aspect ratio of the nanonetworks (curve c). If the AgNO₃ concentration was higher or lower from this value, lower LSPR peak position was obtained, suggesting a decrease in the aspect ratio of the nanonetworks. The FESEM results as shown in Figure 3 further verified such growth characteristic of the nanostructures on the surface under the present treatment. For example, effective nanonetworks formation was confirmed when AgNO₃ concentration of as low as 10 μM was used in the reaction (see Figure 3A). This is probably the origin of the observed LSPR band in the spectra, formed via the presence of a long-range plasmonic coupling process in the nanonetworks. As has been noted earlier, spherical nanoparticle chains did not produce such optical properties resulting from a long-range plasmonic coupling amongst the nanoparticles in the chain. Thus, special nanostructure morphology, i.e. quasi-1D, should be present in the nanonetworks in order to support effective long-range plasmonic coupling via optimum edge-to-edge quasi-1D arrangement. The available FESEM images actually did confirm the nanostructures that composed the nanonetworks are quasi-1D morphology instead of spherical nanoparticles. It is true that high-resolution FESEM images were not available to support the claim. However, the present optical absorption spectra results that show 1D optical property, produced from effective long-range plasmonic coupling, are strong evident for the formation of such nanostructures morphology in the nanonetworks. However, a complete analysis on the detailed morphology of the individual nanostructures on the nanonetworks including high-resolution electron microscopy is being pursued and will be reported in a
different publication. Large-scale nanonetworks were effectively formed covering the entirety of the surface when optimum AgNO₃ concentration was used, namely 20 μM (see Figure 3B). One point to be noted here is that the actual length of the nanonetworks obtained using this condition was relatively the same with that shown in Figure 3A; however, their shorter axis was relatively smaller. Thus, the aspect ratio of the nanonetworks increased, shifting the LSPR to red. As has been earlier probed in the absorption spectra, further increase in the AgNO₃ added decreased the aspect ratio of the nanonetworks. Judging from the FESEM results, we pointed out the following facts as the reason for the process: The density of the nanonetworks increased with the increasing of AgNO₃ concentration. As revealed in Figure 3C, despite the fact that the short axis of the nanonetworks decrease due to the increase in the density and possible strong coupling along the short axis of the nanonetworks, further red-shifting in the LSPR was not achieved but blue-shifting instead. A different structure was obtained

Figure 2 Optical absorption spectra. Gold nanostructures on the surface prepared using several Ag⁺ concentrations, namely (a) 0, (b) 10, (c) 20, (d) 30, (e) 40 and (f) 50 μM. Other reagents were kept unchanged as that in Figure 1. Growth time was 4 h.

Figure 3 FESEM images. Selected gold nanostructures’ growth prepared using several AgNO₃ concentrations, of which their corresponding absorption spectra are shown in Figure 2. (A) 10, (B) 20, (C) 30 and (D) 50 μM. Scale bars are 100 nm.
when high concentration of AgNO₃ was added, namely 50 μM. As can be seen from the Figure 3D, nanonetworks that are composed of relatively bigger spherical nanoparticles were typical of the nanostructures product. The increase in the individual nanoparticle size might be due to a kind of ‘steric’ hindrance of surfactant adhesion onto the nanoparticles surface in the presence of high concentration of AgNO₃, accelerating the growth of individual nanoparticles.

While AgNO₃ drives the formation of 1D crystal growth of the nanoseeds, both CTAB and HMT via their combinatorative effect are also crucial in this process. In a typical case, no 1D optical characteristic was obtained if one of the surfactants was absent in the reaction. Typical optical absorption spectra of gold nanostructures prepared using several concentration ratios of CTAB and HMT (millilitre to millilitre ratio) are shown in Figure 4. As revealed in the spectra, both surfactants must be present in the reaction to give an optimum 1D optical characteristic, which is indicated by the longest LSPR peak position when CTAB to HMT ratio is 12:8 (see curve d). By keeping the cumulative concentration of both surfactants unchanged, if the ratio was increased or decreased, the LSPR peak position was blue-shifted compared to the optimum one, a sign of a decrease in the aspect ratio of the 1D nanostructures. It needs to be noted here that the individual nanoparticles probably grew on the surface when only surfactant HMT presented in the reaction, as judged by its low optical absorbance (see curve h). This could be due to a weak capping nature of the HMT to Au⁺ so that the Au⁺ reduction occurred in the solution instead on the nanoseeds surface. Selected FESEM image of the samples as shown in Figure 5 further confirms and provides detailed pictures of gold nanostructures’ growth characteristic on the surface under several CTAB to HMT concentration ratios. For example, high-yield nanonetworks were obtained when the CTAB to HMT ratio of 12:8 was used. In good agreement with the optical absorption spectrum as shown in Figure 4, the yield was found to decrease when the ratio was increased (see Figure 5C for CTAB to HMT ratio of 10:10). Bigger-sized nanoparticle network was even obtained if the ratio further decrease to 8:12. On the basis of these results, it can be worthwhile concluding that combinative function of surfactants plays a strategic role in the formation of quasi-1D morphology and then the nanonetwork structures on the surface.

It needs to be noted here that the individual nanoparticles which compose the nanonetworks were actually mainly the quasi-1D gold nanostructures formed by a unique interplay amongst the reagents under the present condition. We hypothesised that a facet-selective surfactant adhesion, which is driven by the Ag⁺ via its unique underpotential deposition nature onto the nanoseed surface, could be the main factor for the formation of the structures. However, the exact mechanism is not clear at the moment, especially the nature of the surfactants adhesion as well as the interplay amongst them and the Ag⁺. Although the following facts could be considered: (a) Quasi-1D nanostructures were not formed when one of the surfactants was absent. The quasi-1D nanoparticles may be formed at any concentration ratio between the two surfactants but optimum at the CTAB to HMT ratio of 12:8. This reveals that the combinative function of surfactant here is crucial. (b) The quasi-1D nanostructures were also not formed when Ag⁺ is absent in the reaction. In fact, a diluted Ag⁺ concentration (ca. 10 μM) projected the quasi-1D nanoparticles growth and optimum at a concentration of approximately 20 μM. These indicated that the interplay amongst the surfactants and the Ag⁺ is necessary for the formation of quasi-1D nanoparticles. It is widely known that the Ag⁺ may adsorb onto the gold nanoseed surface and then induces a one-dimensional crystal growth in the nanoseeds, certainly in the presence of surfactant. This scheme might probably also be valid to the present condition. However, since the system was on the solid surface, the growth characteristic should be different. It is true that the bromide ions were recognised to have a key effect in the promotion of nanorod morphology growth in the solution phase via effective adsorption onto the lower-energy surface of the nanostructures, i.e. (111) and (100),

![Figure 4 Optical absorption spectra](image-url)
and play as a steric hindrance on these planes, which lead to a nanorod shape formation with growth direction toward (110) [48]. However, since in this experiment the quasi-1D morphology were not formed in the absence of Ag ion, the role of Br ion in the formation of quasi-1D nanostructures on the surface is not relevant. It is also true that the AgBr complexes may also be formed during this process. Since these complexes are relatively inactive compared to the Br ions as well as their quantity might be considerably small as the result of effective Ag ion deposition onto the Au nanoseed via effective underpotential deposition process, this chemical was believed to have minor effect on the promotion of quasi-1D gold nanostructures.

Meanwhile, on the network structures, their formation is assumed as an attempt to minimalise the high surface energy of the quasi-1D nanostructures that formed under the present condition. Actually, the initial nanoseeds on the surface were randomly distributed without any specific orientation (see Figure 6A). After being grown in the growth solution, adjacent nanoseeds grew into a quasi-1D morphology and aligned with each other forming the nanonetwork structures (see Figure 6B,C). Meanwhile, the nanoseeds that were far from the nanonetworks may have grown into bigger spherical nanoparticles or simply dissolved into the bulk solution via the Oswald annealing process and then supported the growth of bigger nanostructures on the surface. As revealed in the FESEM images, the nanonetwork structures have been observed on the sample that was grown for 30 min. In spite of that fact, their formation may be earlier. The formation of these structures via aggregation amongst the nanostructures is not applicable in this process. It is because the nanostructures were grown from the nanoseeds attached to the surface so that the possibility of the nanostructures migration on the surface could be minimalised. The aggregation of the nanostructures from the bulk solution and then attached onto the surface is also irrelevant here since the mature nanostructures are only on the nanoseeds that are attached to the surface.

Conclusions

We have demonstrated a facile formation of gold nanonetworks from the attached nanoseeds on the surface that exhibit a 1D structure property via a combinatorial effect of CTAB and HMT surfactants and Ag+ in the growth solution. In a typical case, the nanonetworks were mainly composed of small quasi-1D gold nanostructures that probably aligned due to surface energy minimalisation. The quasi-1D nanoparticles as well as the nanonetworks were very sensitive to the CTAB, HMT and Ag+ in the reaction and neither quasi-1D nanostructures nor nanonetworks were formed if one of
the three chemicals was absent. At this stage, the dimension control of both the quasi-1D structures and the nanonetworks, in particular the aspect ratio, was not yet achieved. An intensified further attempt to achieve that control is underway. Owing to its unique structure that composed of small quasi-1D gold nanostructures, the nanonetworks may produce unusual optical and catalytic properties that make them highly demanded in photonic, catalysis and surface-enhanced Raman scattering applications.

Competing interests
The authors declare that they have no competing interests.

Acknowledgement
We acknowledge the support from the Universiti Kebangsaan Malaysia and Ministry of Higher Education of Malaysia under research grants UKM-GUP-NBT-08-25-086, UKM-GUP-2011-377 and UKM-RRR1-07-FRG0037-2009.

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Authors’ contributions
AA carried out the nanostructure preparation and characterisation. AAU designed the concept and experiment, analysed the results and drafted, revised and finalised the manuscript. IL participated in the data analysis. MMS provided the facilities and discussed the results. MO provided the concept of the growth process of the nanostructures. All the authors contributed to the preparation and revision of the manuscript. All authors read and approved the final manuscript.

Received: 12 February 2012 Accepted: 15 May 2012 Published: 15 May 2012

References
1. El-Sayed MA: Small is different: shape-, size-, and composition-dependent properties of some colloidal semiconductor nanocrystals. Accounts Chem Res 2004, 37:326–333.
2. Ringe E, McMahon JM, Sohn K, Cobley C, Xia Y, Huang J, Schatz GC, Marks LD, Van Duyne RP: Unraveling the effects of size, composition, and substrate on the localized surface plasmon resonance frequencies of gold and silver nanocubes: a systematic single-particle approach. J Phys Chem C 2010, 114:12511–12516.
3. Murphy CJ, Gole AM, Hunyadi SE, Orendorff CJ: One-dimensional colloidal gold and silver nanostructures. Inorg Chem 2006, 45:7544–7554.
4. Wu J, Lee S, Reddy VR, Manasreh MO, Weaver BD, Yates MK, Furrow CS, Kunets VP, Benamara M, Salamo CJ: Photoluminescence plasmonic enhancement in InAs quantum dots coupled to gold nanoparticles. Mater Lett 2011, 65:3605–3608.
5. Murphy CJ, Sau TK, Gole AM, Orendorff CJ, Gao J, Gou L, Hunyadi SE, Li T: Anisotropic metal nanoparticles: synthesis, assembly, and optical applications. J Phys Chem B 2005, 109:13857–13870.
6. Ktyk M, Plucinski KJ, Eboth J, Umar AA, Oyama M: Control of the plasmon absorption of gold nanoparticles with a two-color excitation. J Appl Phys 2005, 98:1–4.
7. Munechika K, Chen Y, Tillack AF, Kulkami AP, Plante UL, Munro AM, Ginger DS: Spectral control of plasmonic emission enhancement from quantum dots near single silver nanoprisms. Nano Lett 2010, 10:2598–2603.
Kang MG, Xu T, Park HJ, Luo X, Guo LJ: Circular acoustogyratory effect on gold nanoparticles grown on indium tin oxide. Appl Opt 2005, 44(9):605–609.

Kityk IV, Umar AA: Circularly polarized light-induced electrogravitropism in the Au nanoparticles on the ITO. Phys E 2005, 27:420–426.

Özga K, Kawaiharauma T, Ali Umar A, Oyama M, Nouneh K, Slezak A, Fujita S, Piasceki M, Reshak AH, Kityk IV: Second order optical effects in Au nanoparticle-deposited ZnO nanocrystalline films. Nanotechnology 2008, 19(18):185709.

Geyer U, Hetterich J, Dier C, Hu DZ, Schaad DM, Lemmer U: Nanostructured metallic electrodes for plasmonic optimized light-emitting diodes. In Proceedings of SPE: topics in plasmonic. Plasmonic nanostructures and their optical properties VI. August 10 2008. San Diego. Edited by Stockman MI, Bellingham: SPIE; 2008:70320B.

Standridge SD, Schatz GC, Hupp JT: Distance dependence of plasmon-enhanced photoresponse in dye-sensitized solar cells. J Am Chem Soc 2009, 131:8407–8409.

Beck FJ, Mokkapati S, Catchpole KR: Plasmonic light-trapping for Si solar cells using self-assembled, Ag nanoparticles. Prog in Photovolt: Res and Appl 2010, 18:500–504.

Kang MG, Xu T, Park HJ, Luo X, Guo LJ: Efficiency enhancement of organic solar cells using transparent plasmonic Ag nanowire electrodes. Adv Mater 2010, 22:4579–4583.

Standridge SD, Schatz GC, Hupp JT: Toward plasmonic solar cells: protection of silver nanoparticles via atomic layer deposition of TiO₂. Langmuir 2009, 25:2596–2600.

Wu J, Makableh YFM, Vasan R, Manasek MO, Liang B, Reyner CJ, Huffaker DL: Strong interband transitions in InAs quantum dots solar cell. Appl Phys Lett 2012, 100(5):190707.

Biesboer A, Qian W, Huang X, El-Sayed MA: Gold nanoparticles surface plasmon field effects on the proton pump process of the bacteriorhodopsin photosynthesis. J Am Chem Soc 2009, 131:2442–2443.

Jain PK, Huang X, El-Sayed IH, El-Sayed MA: Review of some interesting surface plasmon resonance-enhanced properties of noble metal nanoparticles and their applications to biosystems. Plasmonics 2007, 2:107–118.

Lee KS, El-Sayed MA: Gold and silver nanoparticles in sensing and imaging: sensitivity of plasmon response to size, shape, and metal composition. J Phys Chem B 2006, 110:19220–19225.

Orendorff CJ, Baxter SC, Goldsmith EC, Murphy CJ: Light scattering from gold nanorods: tracking material deformation. Nanotechnology 2005, 16:2601–2605.

Eustis S, El-Sayed MA: Why gold nanoparticles are more precious than pretty gold: noble metal surface plasmon resonance and its enhancement of the radiative and nonradiative properties of nanocrystals of different shapes. Chem Soc Rev 2006, 35:209–217.

Jain PK, Huang X, El-Sayed IH, El-Sayed MA: Noble metals on the nanoscale: optical and photothermal properties and some applications in imaging, sensing, biology, and medicine. Accounts Chem Res 2008, 41:1578–1586.

Huang X, Jain PK, El-Sayed IH, El-Sayed MA: Plasmonic photothermal therapy (PPTT) using gold nanoparticles. Lasers Med Sci 2008, 23:217–228.

Huang X, Jain PK, El-Sayed IH, El-Sayed MA: Gold nanoparticles: interesting optical properties and recent applications in cancer diagnostics and therapy. Nanomedicine 2007, 2:691–693.

Klehtostov NG, Dykman LA: Optical properties and biomedical applications of plasmonic nanoparticles. J of Quant Spectrosc and Radiat Transf 2010, 111:1–35.

Weissleder R: A clearer vision for in vivo imaging. Nat Biotechnol 2001, 19:316–317.

Murphy CJ, Golle AM, Hunyadi SE, Stone JW, Sisco PN, Alkilany A, Kinar DS, Hankins P: Chemical sensing and imaging with metallic nanoparticles. Chem Commun 2008, 8:544–557.

Murphy CJ, Safi TK, Golle A, Orendorff CJ: Surface-directed synthesis and optical properties of one-dimensional plasmonic metallic nanostructures. MRS Bulletin 2005, 30:349–355.

Wang ZL, Gao RP, Nikoobakht B, El-Sayed MA: Surface reconstruction of the unstable 110 surface in gold nanorods. J Phys Chem B 2000, 104:5417–5420.

Murphy CJ, Thompson LB, Alkilany AM, Sisco PN, Boulois SP, Sapvalan ST, Yang JA, Chemak DJ, Huang J: The many faces of gold nanorods. J Phys Chem Lett 2010, 1:2667–2675.

Narayanan R, El-Sayed MA: Effect of colloidal nanocrystal analysis on the metallic nanoparticle shape: the suzuki reaction. Langmuir 2005, 21:2007–2033.

Umar AA, Oyama M: Electrochemical responses on gold nanoparticles directly attached indium tin oxide electrodes fabricated using a ‘touch’ seed-mediated growth method. Indian J Chem Inorg Phys Theor Anal Chem 2009, 44:938–944.

Jana NR, Gearheart L, Murphy CJ: Wet chemical synthesis of silver nanorods and nanowires of controllable aspect ratio. Chem Commun 2001, 7:617–618.

Nikoobakht B, El-Sayed MA: Preparation and growth mechanism of gold nanorods (NRs) using seed-mediated growth method. Chem Mater 2003, 15:1957–1962.

Acharya S, Efrima S: Two-dimensional pressure-driven nanorod-to-nanowire reactions in Langmuir monolayers at room temperature. J Am Chem Soc 2005, 127:3486–3490.

Liu K, Zhao N, Kurnacheva E: Self-assembly of inorganic nanorods. Chem Soc Rev 2011, 40:6566–6571.

Umar AA, Oyama M: Growth of high-density gold nanoparticles on an indium tin oxide surface prepared using a ‘touch’ seed-mediated growth technique. Cryst Growth Des 2005, 5:599–607.

Umar AA, Oyama M: Attachment of gold nanoparticles onto indium tin oxide surfaces controlled by adding citrate ions in a seed-mediated growth method. Appl Surf Sci 2006, 253:2933–2940.

Umar AA, Oyama M: A cast seed-mediated growth method for preparing gold nanoparticle-attached indium tin oxide surfaces. Appl Surf Sci 2006, 253:2196–2202.

Umar AA, Oyama M, Salleh MM, Mallis BY: Formation of high-yield gold nanoplates on the surface: effective two-dimensional crystal growth of nanoseed in the presence of poly(vinylpyrrolidone) and cetyltrimethylammonium bromide. Cryst Growth Des 2009, 9:2835–2840.

Umar AA, Oyama M: Formation of gold nanoplates on indium tin oxide surface: two-dimensional crystal growth from gold nanoseed particles in the presence of poly(vinylpyrrolidone). Cryst Growth Des 2006, 6:818–821.

Jana NR, Gearheart L, Murphy CJ: Wet chemical synthesis of high aspect ratio cylindrical gold nanorods. J Phys Chem B 2001, 105:4065–4067.

Smith DK, Korgel BA: The importance of the CTAB surfactant on the colloidal seed-mediated synthesis of gold nanorods. Langmuir 2008, 24:644–649.

Zhang J, Oyama M: Gold nanoparticle arrays directly grown on nanostructured indium tin oxide electrodes: characterization and electroanalytical application. Anal Chim Acta 2005, 540:299–306.

Umar AA, Oyama M: High-yield synthesis of tetrahedral-like gold nanotipods using an aqueous binary mixture of cetyltrimethylammonium bromide and hexamethylenetetramine. Cryst Growth Des 2009, 9:1146–1152.

Jain PK, Eustis S, El-Sayed MA: Plasmon coupling in nanorod assemblies: optical absorption, discrete dipole approximation simulation, and excitation-coupling model. J Phys Chem B 2006, 110:18234–18253.

Garg N, Scholl C, Mohanty A, Jin R: The role of boride ions in seeding growth of Au nanorods. Langmuir 2010, 26:10271–10276.