Advanced AuNMs as nanomedicine’s central goals capable of active targeting in both imaging and therapy in biomolecules

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Introduction

Au based nanomaterials (AuNMs) are known to possess many attractive features such as unique electrical, optical and catalytic properties as well as excellent biocompatibility. These outstanding characteristics make them promising candidates as the signal reporters, enhancement materials or others involved with bioassay, food safety and environmental monitoring.

Physics, chemistry and biology of Au based nanomaterials (AuNMs) have emerged as a broad and new subdiscipline in the community of colloids and surfaces. The specific size and shape dependent physiochemical properties and remarkable bio/chemical inertness of AuNMs have made themselves the ideal candidates for both fundamental and technical study including crystal growth, electron-transfer mechanism, localized electro-magnetic theory, catalysis, DNA assay, bioimaging and therapy, and so on [1]. Among those rich properties, the optical characteristics originated from the giant electromagnetic field near the surface of AuNMs are particularly intriguing and thus broadly applied in analytical science, e.g. colorimetric assay, surface enhanced Raman spectroscopy (SERS) and surface Plasmon resonance (SPR) spectroscopy, as well as bioimaging. Another interesting optical property of AuNMs is that fluorescence appears with their size shrinkage to below 2 nm, which allows development of luminescence-based analysis. Also, electrochemical (EC) sensors could be constructed based on the redox feature of Au NMs (Figure 1).

Analytical science has been playing a primary role in our daily life, for instance, food safety control [2], biomedical diagnosis [3], medicolegal appraisement [4], anti-terrorism alert [5], and environmental pollution monitoring [6]. Efficient detection technologies depend greatly on the analytical method in terms of analysis time, signal-to-noise (S/N) ratio, selectivity, and limit of detection (LOD) [7]. Therefore, a great of efforts should be paid to develop new analytical methods with the advantages of rapid determination, high analysis flux, satisfactory sensitivity and good robustness. Thanks to the above-mentioned optical and electrochemical features, many functionalized AuNMs have been successfully synthesized [8-13] for improving performances of analytical techniques.

Small organic molecules and metal ions

For the sensitive detection of small organic molecules, the AuNMs play an important role in improving sensitivity of the EC sensors. Especially, AuNMs composited with conducting polymers or carbon nanomaterials have been used to increase electro-chemical efficiency. It deserves noting that the decorated carbon nanomaterials with metal NPs are of special interest due to the integrated property of two components with better catalytic activity and enhanced electrical conductivity [14,15]. Zhu et al. [16] fabricated EC sensors by using spherical AuNPs combined with conducting polymer (poly-[2, 5-di-(2- thienyl)-1H-pyrrole-1-(p-benzoic acid])). To further improve electron transfer, synergistic effect of multiple nanocomposite layers, namely, chitosan-AuNPs, graphene-AuNPs and multi-walled carbon nanotubes (MWCNTs)-cobalt phthalocyanine, was exploited by Sun et al. [17]. Magalhães et al. [18] fabricated a sensitive and selective EC aptasensor for detection of cocaine based on the conformational change of the aptamer-functionalized AuNPs onto CNTs-based nanocomposite.

Detection of organic small molecules and metal ions is very important for controlling food quality, environmental protection and antiterrorism. In order to realize detection of trace amount of analytes at extremely low concentrations, various efforts for increasing the sensitivity of SPR biosensors have been paid. Most of these methods

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Figure 1. Application of AuNMs in analytical science.
are based on changing the refractive index at the sensor surfaces by means of a variety of (bio) chemical or NP agents, which are captured by the sensor surfaces subsequent to the binding of the target analyte to the primary biorecognition elements. These methods include the use of secondary and tertiary antibodies, antibodies labeled with enzymes, and dielectric or metallic NPs [19]. In particular, AuNMs of diameters ranging from 5 to 40 nm have been widely used to enhance the response of SPR biosensors [20,21].

**Synthesis of AuNMs**

Traditionally, the trisodium citrate-based thermal reduction has been broadly recognized as an universal approach for preparing spherical Au nanoparticles (AuNPs) of varied sizes ranging from 15 to 147 nm possessing strong SPR signals [22]. Although this preparation method simply involves controlling the ratios of citrate to Au ions [23], the reaction often needs to be carried out on a relatively large scale at the temperature of boiling water and lacks the reproducible preparation of uniform AuNPs with rather large sizes. After the breakthroughs reported by Schmid [24,25] and Brust et al. [26], a variety of methods have been developed to synthesize AuNMs, and many excellent reviews [27] are available now. In general, the reported synthesis methods of spherical AuNMs can be categorized as followed: (1) citrate reduction; (2) Brust-Schiffrin phase transfer synthesis; (3) other sulfur-containing ligands method; (4) soft templates including micro emulsion, reversed Micelles, surfactants, membranes and polyelectrolytes; (5) seed-mediated growth; and (6) physical methods [28]. In addition, non-spherical nanoparticles (NPs) such as Au nanorods (AuNRs) [29], Au nanocages [30], Au cubes [31], Au triangular prisms [32], as well as other unconventional structures like hollow tubes and even branched nanocrystals [33] have attracted significant research attention in the past two decades, since the properties of AuNMs are known to be strongly dependent on the size and shape of the particle [34]. Recently, a lot of excellent reviews have summarized how to achieve shape- and size-defined AuNMs with different techniques including solve thermal synthesis, seed-mediated growth, homogeneous nucleation and electrochemical method [35,36]. In this review, we mainly introduce the latest and important advancement of synthesis of the AuNMs, especially on how to control the morphology of AuNMs. A great of efforts have been paid to achieve high-quality and shape-controlled AuNMs for the purpose of highly-sensitive analysis. Unfortunately, most of the synthetic methods are empirical, and understanding their growth mechanism remains a challenging task. Interestingly, introduction of Ag+ in growth solutions significantly improves the yield of AuNRs, and Kim et al. [37] prepared AuNRs with controlled aspect ratio by using photochemistry in the presence of Ag+.

Benefiting from their high symmetry, the Au nanostars exhibited much better single-particle SERS performance compared with asymmetric Au nanostars, in terms of both intensity and reproducibility. Apart from large-sized Au particles, Au nanoclusters (AuNCs) with the sizes of less than 2 nm formed in dendrimers displays visible luminescence of high quantum yield (QY) [40]. Ying et al. reported a simple, one-pot and “green” synthetic route, based on the reduction and stabilization capability of bovine serum albumin (BSA) proteins. The AuNCs prepared at the physiological temperature (37°C) showed red emission with a QY of ~6% [41]. In order to further control the luminescence property, synthesis of atomically precise AuNCs has ignited numerous research efforts [42,43]. In 2014, Yu et al. [44] synthesized Au55 (glutathione)18 clusters with a luminescence QY greater than 60% by rigidifying their Au shell with tetraoctylammonium cations. This study presents an effective strategy to enhance the luminescence efficiencies of AuNCs by optimizing the shell structure.

In order to precisely tune the size and shape of AuNMs, DNA is recently adopted as a powerful programmable tool to realize above goals. Lu’s group systematically summarized morphology controlling, spatial positioning and dynamic assembly of AuNMs with DNA as a powerful tool [46]. In some details, their research results demonstrated that DNA could be used to control the morphologies of AuNMs during seed-mediated growth [47]. They also developed a novel method based on DNA-encoded tuning to achieve AuNMs from AuNR seeds with controlled geometric and plasmonic properties [48]. Furthermore, they suggested that the DNA molecules played important roles via influencing diffusion of the Au precursors to the seed, and therefore the AuNMs growth might be modulated through difference in DNA desorption, density and mobility on the seed surfaces [49]. These insights into the mechanism of DNA-guided AuNMs synthesis not only provide deep understanding of the interactions between the DNA and nanomaterials but also allow better control of the shapes and surface properties of many nanomaterials. Ye et al. [50] designed a millifluidic reactor to synthesize a wide variety of AuNMs at high concentrations. The synthesized method based on millifluidic reactors inspired other researchers to use the analogous reactors for possible mechanism and application study. Recently, Krishna et al. demonstrated adoption of a simple millifluidic chip for in situ analysis of morphology- and dimension-controlled growth of Au nano- and micro-structures with a time resolution of 5 ms [51] (Figure 2). See from Figure 2, gold structures with ~3 μm in diameter that had corrugated metal-like shapes were formed after a time interval of 5 h (Figure 2d). When the time of flow was increased to 9 h, these structures further transformed into micro-hemispherical ones (Figure 2c). This study offers a feasible way to obtain dimension-and morphology-controlled AuNMs in a high yield.

Noteworthy, some novel synthesis strategies, such as biosynthesis, 2D immiscible oil/water interface, levitated leidenfrost drop, sunlight-induced synthesis and so on, have been reported, and such green and economic synthesis methods for preparation of uniform and controlled AuNMs provide the opportunity for development of low-cost and sustainable analytical techniques. However, the above technologies need to be further improved and standardized.
Under light excitation, a lot of “hot spots” were created between the Au
sandwiched between the capture antibody and the detection antibody.
(Au@Ramanreporter@SiO2 sandwich NP). The antigen (analyte) was
chip, while the detection antibody was conjugated with the SERS probe
the capture antibody was immobilized on the Au triangle nanoarray
“hot spot” groups for protein microarrays. Li
et al. [51]. Owing to the large surface to volume ratio, modification and
functionalization of AuNMs surfaces with biomolecules, DNA or
chemicals becomes imperative for their application in analytical
science.

It is well known that free AuNMs have high surface energy and tend
to aggregate and fuse. As a result, the intriguing properties observed
for the AuNPs would disappear, and the difficulty arises for long term
storage, processing, and applications. Therefore, great efforts have
been devoted to develop novel strategies to stabilize AuNMs [57], and
the most common approach is to coat AuNMs with either organic or
inorganic shells. These shells not only endow AuNPs with high
stability but also offer them additional functionalities. As an example,
in addition to good stability and biocompatibility, the mesoporous
silica shells that are currently broadly used have high surface area and
tunable pore size and volume, which can accommodate analytes and
drug molecules [58].

**Biomolecules**

AuNMs have been widely used in SERS based immunoassays of
biomolecules such as DNA, protein and cell. However, challenges still
remain with amplification of SERS signals due to the extremely small
cross-section of Raman scattering [59]. Yan et al. [60] introduced a new
strategy based on nano rolling-circle amplification (nanoRCA) and
nano hyperbranched rolling circle amplification (nanoHRCA) to increase
“hot spot” groups for protein microarrays. Li et al. [61] constructed a
SERS immune sensor for detection of biomarkers. In their study, the
capture antibody was immobilized on the Au triangle nanoarray chip, while the detection antibody was conjugated with the SERS probe
(Au@Ramanreporter@SiO2 sandwich NP). The antigen (analyte) was
sandwiched between the capture antibody and the detection antibody.
Under light excitation, a lot of “hot spots” were created between the Au
triangles in nanoarray, and thus the developed SERS immunoassay
could be used for sensitive biomarker detection. Luo et al. [62] reported
a facile immunoassay for porcine circovirus type 2 (PCV2) based on
SERS using multi-branched AuNPs (mb-AuNPs) as substrates. The
mb-AuNPs in the immunosensor act as Raman reporters, which were
prepared via Tris base-induced reduction and subsequent reaction with
p-mercaptobenzoic acid. The modified mb-AuNPs were covalently
conjugated to the mono-clonal antibody (McAb) against the PCV2
cap protein to form SERS immuno nanoprobes. These were captured in
a microtiterplate via an immunoreaction in the presence of target
antigens. Compared to conventional detecting methods such as those
based on PCR, the method was demonstrated to be rapid, facile and very
sensitive. Tang et al. developed a simple and reproducible SERS chip
for highly sensitive and selective screening of active ricin in complex
matrices [63]. They fabricated single strand oligodeoxynucleotides
modified AuNPs in order to amplify recognition and reaction by virtue
of their cooperative binding property. To improve sensitivity of this
method, a Ag nanoshell was deposited on post-reacted poly (21dA)-
AuNPs, which lowered LOD to 8.9 ng mL⁻¹.

The utilization of SERS-based techniques in targeting specific DNA
and RNA sequences is generally realized by combination with AuNMs and
Raman-active molecules [64,65]. Wang et al. [66] raised a concept
to use enzyme controlled plasmonic coupling as SERS nanosensors for
DNA demethylation. The nanosensors were constructed by decorating
AuNPs with Raman reporters and hemimethylated DNA probes.
The enzymatic degradation of DNA substrate probes was utilized
to induce aggregation of AuNPs, so the reproducible and sensitive
SERS signals were achieved from biological recognition events. Development of SERS-barcoded particles, which contain multiple
Raman-active molecules and are insensitive to photobleaching, is
particularly intriguing for high-throughput and multiplex DNA
screening [67]. Recently, Chen et al. [68] prepared a SERS-barcoded
nanosensor to detect Bacillusthuringiensis (Bt) gene transformed rice
expressing insecticidal proteins. The barcoded sensor was designed by
encapsulation of AuNPs with silica and conjugation of oligonucleotide
strands for targeting DNA strands. The transition between the cry1A
(b) and cry1A(c) fusion gene sequence was used to construct a specific
SERS-based detection method with a LOD of 0.1 pg/mL. In order to
build the determination models to screen transgene, a series mixture of
Bt rice and normal rice were prepared for SERS assay, and the LOD was
0.1% (w/w) was explored for transgenic Bt rice relative to normal rice.
The sensitivity and accuracy of the SERS-based assay were comparable
with real-time PCR. Recently, Ye et al. [69] proposed an asymmetric
signal amplification method for simultaneously detecting multiple
biomarkers with significantly different levels. The biobarcode consisted
of a large number of signals DNA (Cy3-DNA for Cy3-bio barcode or
Rox-DNA for Rox-biobarcode) and capture DNA immobilized onto
the Au NPs. The AuNPs simultaneously act both as a Raman-
signal-enhancing substrate and a Raman signal carrier. Using these
bifunctional probes, a linear amplification mode was obtained with
high-concentration markers, whereas quadratic amplification mode
was responsible to low-concentration markers.

Detection of circulating tumor cells (CTCs) in the blood of cancer
patients is significant for early cancer diagnosis, cancer prognosis,
evaluation of the treatment effect of chemotherapy drugs, and choice
of cancer treatment options [70]. In 2011, Wang et al. introduced
SERS technology for direct detection of target CTCs in human
peripheral blood. The LOD of this method ranged from 5 to 50CTCs
in 1 mL of blood [71]. Recently, Wu et al. [72] fabricated SERSNPs
for direct detection of CTCs in the blood with excellent specificity

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**Figure 2.** Scheme showing different stages of spatially (and time) resolved growth process of AuNMs within millifluidic chip. Printed with permission from Krishna et al. [51]. Copyright 2013 American Chemical Society.
and high sensitivity. The AuNPs were encoded with a Raman reporter molecule, 4-mercaptobenzoic acid (4-MBA), and then functionalized with reductive bovine serum albumin (rBSA) to stabilize the 4-MBA-encoded AuNPs (AuNP-MBA) and decrease the nonspecific interaction with blood cells. Then AuNP-MBA-rBSA-FA composite NPs were constructed with a targeted ligand folic acid (FA) (Figure 3 A and B). The FA on the surface of AuNP-MBA-rBSA-FA NPs was recognized by CTCs, which over expressed folate receptor alpha (FRα). The protection layer of rBSA was much thinner than that of the reported PEG, resulting in a stronger SERS signal (Figure 3C).

For detection of the biological macromolecules such as bacteria pathogen and enzyme, SPR with the advantage of label free provides a powerful platform [73,74]. There are many successful examples of SPR sensing as a medical diagnostic tool, which have been reported for biomarkers, pathogen detection and hormone analysis with high sensitivity. For example, McPhillips et al. [75] employed aligned Au nanotube arrays to strengthen performance of refractive index sensors in biomolecular binding reactions. Generally, sandwich [76] and competitive or inhibition assay [77] are two major detection approaches in SPR biosensor. In 2012, Fernandez et al. [78] demonstrated that AuNMs combined with sandwich assays could improve detection sensitivity and selectivity of protein detection. Jung et al. [79] introduced a strategy for enhancement of SPR signals by adopting AuNPs and a SiO2 layer on an Au surface. The modified surfaces showed significant changes in SPR signal when biomolecules were attached to the surface as compared with an unmodified Au surface. The LOD of AuNPs immobilized on a SPR chip was 0.1 ng mL−1 for the prostate specific antigen (PSA), a cancer marker. The enhanced shift of the absorption curve resulted from coupling of the surface and particle plasmon by the SiO2 layer and the AuNPs on the gold surface.

Now, there were a lot of reports of SPR sensors based on spherical AuNMs for detection of biomolecules [80,81]. The anti-E.coli O157:H7 polyclonal antibodies (pAb) were labeled with AuNPs, which were used as secondary antibodies. The E.coli O157:H7 was detected using direct assay and enhancing sandwich assay based on the two channels SPR biosensor. By introducing AuNPs-PAb compound, the LOD was determined to be 10 cfu/mL. The sensitivity was 100 times higher than that of direct detection [82]. Compared with spherical AuNMs, AuNRs labels are more favorable for the Au-amplified SPR biosensors because of the tunable longitudinal plasmonic peak enables an effective plasmonic coupling between sensing film and NPs. Law et al. [83] revealed the potential of applying this “perfectly matched” nanotag in a well-established SPR sensing system and immunoassay. Through detection of tumor necrosis factor alpha antigen, 40-fold sensitivity enhancement using wavelength-matched AuNRs was observed. Recently, Xiang et al. [84] developed a SPR DNA biosensor array based on target-primed rolling circle amplification (RCA) for isothermal and rapid detection of two pathogenic mycobacteria, mycobacterium tuberculosis complex (MTBC) and mycobacterium avium complex (MAC) (Figure 4). AuNPs were directly assembled on to the surface of the sensor chip via hexanethiol (HDT) for enhancement of sensitivity as a label-free detection system. Experimental results showed that the signal enhancement by the target-primed RCA together with AuNPs-embedded surface caused at least 10-fold increased sensitivity as compared with conventional RCA on bare SPR chip method.

Besides, the chemical vapor deposition (CVD) method has been adopted by Ruoff’s group for large scale preparation of high-quality graphene sheets [85]. The strategy based on AuNMs combined with graphene sheets could improve the conductivity of SPR [86]. In 2013, Zhang et al. [87] employed Au-graphene oxide composite in the

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**Figure 3.** (A) Schematic Illustration for the Design of SERS Nanoparticles. (B) Characterization of the AuNPs (control) and SERS nanoparticles. TEM images of AuNPs (a) and AuNP-MBA5-rBSA2-FA3 (b). (c) Size distributions of AuNPs and AuNP-MBA5-rBSA2-FA3 in Milli-Q water at room temperature. (d) UV-vis spectra of AuNPs and AuNP-MBA5-rBSA2-FA3. (C) Detection sensitivity of the AuNP-MBA5-rBSA2-FA3 nanoparticles for MCF-7 cells in the rabbit blood. Printed with permission from Wu et al. [72]. Copyright 2015 American Chemical Society.
wavelength modulation SPR biosensor for rabbit IgG detection. In their study, the staphylococcal protein A (SPA) modified Au-graphene oxide composite was directly immobilized onto SPR chips without any additional chemical treatment. The biosensor with the SPA modified Au-graphene oxide composite as the enhanced sensing platform exhibited a satisfactory response to rabbit IgG. The LOD obtained with the composite was 16 times lower than that obtained with the SPA modified chip. This study provided a simple and effective approach for fabrication of sensitive SPR immunosensors and extended the application of the Au-graphene oxide composite in immunoassays.

Molecularly imprinted polymers (MIPs) have become a competitive tool in the field of molecular recognition, owing to their low cost, physical robustness, thermal stability and easy preparation over biological receptors and other functionalized materials [2]. High surface-to-volume ratio makes AuNPs a competitive candidate as matrix material for novel nanosized MIPs. Xie et al. [88] showed a surface molecular self-assembly strategy for MIP in electro polymerized aminothiophenol (ATP) membranes at the AuNP-modifed glass carbon electrode for electrochemical detection of pesticide chlorpyrifos. Yu et al. [89] constructed an EC sensor for determination of dopamine based on core-shell composite of AuNPs and SiO₂ MIPs through sol-gel technique. Similarly, Xue et al. [90] prepared a highly sensitive and selective biomimetic EC sensor for amperometric detection of trace dopamine (DA) in human serums by AuNPs doped MIPs (Figure 5).
The LOD of the proposed EC sensor for DA was 7.8 n mol/L. Sun et al. [91] presented an EC sensor for detection of 3-chloro-1,2-propandiol (3-MCPD) that was usually used as surfactant. They fabricated an AuNPs-modified glassy carbon electrode coated with an MIP film via electropolymerization. The LOD reached $3.8 \times 10^{-18}$ mol/L impressively, the sensor showed high sensitivity, good selectivity, excellent reproducibility and stability during the quantitative determination of 3-MCPD.

**Metal ions**

Some EC sensor methods based on AuNMs have been developed for detection of heavy metal ions. For examples, Shen et al. [92] proposed an EC DNAzyme sensor based on AuNPs for sensitive and selective detection of Pb$^{2+}$. A label-free EC sensor was developed for the highly sensitive and selective detection of Hg$^{2+}$ by Tang et al. [93]. In their study, the catalytic HAuCl$_4$/NH$_2$OH reaction was utilized for formation of AuNPs as signal reporter after capturing Hg$^{2+}$ on the modified electrode surface through the specific thymine-Hg$^{2+}$-thymine (T-Hg$^{2+}$-T) coordination. Compared with the traditional metal NPs-based method, this sensor avoided the labeling of the DNA probe with NP tags, and only one unlabeled T-rich DNA sequence was needed, which greatly reduced the cost and simplified the sensing procedure. Thus, a LOD as low as 0.06 nM could be obtained for Hg$^{2+}$. Furthermore, in 2016, Wang et al. [94] presented a sensitive, selective and reusable EC biosensor for determination of Hg$^{2+}$ based on thymine modified AuNPs/reduced graphene oxide nanocomposites (Figure 6). Notably, the developed EC sensors afforded excellent selectivity for Hg$^{2+}$ against other heavy metal ions including Zn$^{2+}$, Cd$^{2+}$, Pb$^{2+}$, Cu$^{2+}$, Ni$^{2+}$ and Co$^{2+}$. Wang et al. [95] prepared an EC sensor based on graphene and AuNPs for detection of trace Cu$^{2+}$ with the anodic stripping voltammetry analysis. Yang et al. [96] constructed a highly sensitive electrochemical DNA biosensor made of polyaniline (PANI) and AuNPs nanocomposite (AuNPs@PANI) for detection of trace concentration of Ag$^+$. The presence of Ag$^+$, with interaction of cytosine-Ag$^+$-cytosine (C-Ag$^+$-C), cytosine-rich DNA sequence immobilized onto the surface of AuNPs@PANI had a self-hybridization and then formed a duplex-like structure. The prepared EC sensors could detect Ag$^+$ at a wide linear range of 0.01–100 nM with a LOD of 10 pM.

**Biomolecules**

Ultrasensitive detection of biomolecules (especially for disease markers) is very important for early disease diagnosis. To realize ultrasensitive detection of biomolecules, development of novel and robust detection methods for signal amplification of EC sensors is essential. In past work, a lot of photo- or electro-active indicators, such as ferrocene [97], methylene blue [98] and enzymes [99], were immobilized onto AuNPs to realize the detection signals amplification. Zheng et al. prepared an ultrasensitive EC sensor based on network-like thiocyanuric acid/AuNPs for detection of thrombin [100]. Zhao et al. [101] fabricated an ultrasensitive and highly specific EC aptasensor for thrombin based on amplification of aptamer-AuNPs-horseradish peroxidase conjugates. He et al. [102] reported a label-free and sensitive signal-on electrochemical assay for MTase quantification and activity analysis using AuNPs amplification. In 2016, an ultrasensitive DNA biosensor was developed through in situ labeling of electroactive melamine-Cu$^{2+}$ complex (Mel-Cu$^{2+}$) on the end of hairpin-like probe using AuNPs as the signal amplification platform by our group [103]. In this work, the highly conductive AuNPs were applied as the loading platform of the Mel-Cu$^{2+}$ tags. The proposed biosensor showed high selectivity, wide linear range and low LOD for target DNA.

Except for the aforementioned amplification strategies, rolling circle amplification (RCA) is one of the most popular and important amplification strategies. Therefore, the dual amplification strategies combined RCA with DNA-AuNPs probe could tremendously improve the sensitivity of bacterial detection. Zhu et al. [104] reported an electrochemical sensing strategy for ultrasensitive and rapid detection of Salmonella by combining the RCA with DNA-AuNPs probe. As shown in Figure 7, the target DNA could be specifically captured by probe 1 on the sensing interface. Then the circularization mixture
was added to form a typical sandwich structure. In the presence of dNTPs and phi29 DNA polymerase, the RCA was initiated to produce micrometer-long single-strand DNA. Finally, the detection probe (DNA-AuNPs) recognized RCA product to produce enzymatic electrochemical signal. The developed method was successfully applied to detect Salmonella with lower LOD of 6 CFU mL⁻¹ in real milk sample.

Both of graphene based materials and AuNPs are the frequently used nanomaterials in the field of EC biosensors for detection of biomolecules due to their excellent electrical signal amplification and the versatile functionalization chemistry. Accordingly, hybrid nanomaterial sensors based on AuNPs distributing on the surface of graphene oxide (GO) or reduce GO (rGO) also attract much scientific interest [105,106]. The most frequently used technique refers to reaction of AuCl₃ with GO under reductive condition for in situ anchoring AuNPs to GO or rGO. This method, however, often lacks fine control over the size, uniformity and density of AuNPs on the GO sheets in the reaction process [107]. AuNPs can be decorated to GO surface based on NH-Au binding [108] or SH-Au binding [109]. DNA [110] and proteins [111] have been also used as the molecular linkers between AuNPs and GO. These methods have greatly increased the uniformity and density of AuNPs. Liu et al. [112] presented stable label-free EC sensor for detection of cardiac troponin-I (cTnI) in the early diagnosis of myocardial infarction based on AuNPs and GO nanocomposites. The EC sensor demonstrated good selectivity and high sensitivity against human-cTnI, and was capable of detecting cTnI at concentrations as low as 0.05 ng mL⁻¹, which was 100 times lower than that by conventional methods. Karaboga et al. [113] described a simple and disposable immunosensor based on indium-tinoxide (ITO) sheets modified with AuNPs to sensitively analyze heat shock protein70 (HSP70), a potential biomarker that could be evaluated in diagnosis of some carcinomas. Wang et al. [114] constructed an AuNPs/polyaniline/chitosan-graphene sheets based electrochemical DNA sensor with functionalhairpin probe for detection of BCR/ABL fusion gene in chronic myelogenous leukemia. Recently, the interest in layered transition metal chalcogenides, especially MoS₂, has been growing quickly because they share many impressive physicochemical properties of graphene. Su et al. [115] have prepared an EC sensor based on MoS₂ layer for sensitive detection of proteins. In subsequent work, they further fabricated a MoS₂ based EC aptasensor for simultaneous detection of thrombin and adenosine triphosphate (ATP) based on AuNPs-MoS₂ nanocomposites [116].
Development of novel ultrasensitive EC sensors provides a great opportunity for quick and specific sensing of cancer molecular markers (CMMs), which early detection is of great importance for the success of cancer therapy [117]. Johari-Ahar et al. [118] modified an Au electrode with mercaptopropionic acid (MPA), and then consecutively conjugated with silica coated AuNPs (AuNPs@SiO$_2$), CdSe QDs and anti-CA-125 monoclonal antibody (mAb). Successive conjugation of AuNPs@SiO$_2$, CdSe QD and anti-CA-125 mAb onto the Au electrode resulted in sensitive detection of CA-125 with a LOD of 0.0016 U mL$^{-1}$. Serafin et al. [119] presented an immuno reagent label-free strategy to construct a dual EC immunosensor using car-bon nanotube screen printed electrodes modified with AuNPs and PEDOT NPs for the multiplexed determination of human growth hormone and prolactin.

The ECL of luminol promoted by AuNMs was reported by Cui et al. [120]. The ECL intensity of luminol was found to be enhanced by ~2 to 3 orders with catalysis of AuNPs. Li et al. [121] proposed a simple and sensitive sandwich-type ECL immunosensor for detection of cancer antigen 125 on a nanoporous Au modified glassy carbon electrode. Gao et al. [122] prepared an ultrasensitive luminol ECL immunosensor by using carboxylic group functionalized MWCNTs as platform and glucose oxidase supported on AuNPs decorated MWCNTs as labels. The results indicated that this developed sensor exhibited sensitive and stable response for detection of α-1-fetoprotein (AFP), ranging from 0.0001 to 80 ng mL$^{-1}$ with a LOD down to 0.03 pg mL$^{-1}$. Gui et al. [123] constructed a simple and sensitive ECL biosensor for determination of β-lactamase with Ru (phen), (cpphen)Ru$^{3+}$-linked-ampicillin (Ru-Amp). In their work, Ru-Amp complex act not only as a specific recognition element for β-lactamase but also as the ECL luminescent reagent. The biosensor displayed excellent sensitivity of a concentration variation from 50 pg mL$^{-1}$ to 100 ng mL$^{-1}$ with a high sensitivity of 37 pg mL$^{-1}$. Beside the organic dye, noble metal NPs could greatly affect ECL behavior of semiconductor QDs [124]. The ECL enhancement by LSPR of AuNPs was investigated and termed as surface enhanced ECL (SEECL) [125]. The LSPR of AuNPs was found to increase both the excitation rate and the emission factor of luminophores, and thus the ECL intensity was enhanced greatly. Based on this SEECL effect, a method was developed for ultrasensitive detection of Hg$^{2+}$ in drinking water [126]. Moreover, a SEECL sensor was prepared for detection of carcinoembryonic antigen (CEA). In this sensor, Ru (bpy)$_2$2$^{2+}$-doped SiO$_2$ NPs (Ru@SiO$_2$) act as ECL luminophores, while AuNPs were used as LSPR source to enhance the ECL signal. Two different types of aptamers specific to CEA were modified on the surface of Ru@SiO$_2$ and AuNPs, respectively. A LOD of 1.52 × 10$^{-6}$ ng/mL of CEA in human serum was achieved [127].

Electrochemical kinase analysis is of special interest in monitoring biological activity in process of life. Measurements of phosphorylation reactions based on oxidation current of electroactive species such as tyrosine [128], ferrocene [129], etc., which are conjugated on the substrate during the phosphorylation processes, have been reported [130]. To simplify the detection procedure, AuNPs labeled phosphorylation process for the kinase assay was designed by measuring the redox currents of AuNPs [131]. However, the sensitivity of the developed method was relatively low. In order to improve the analytical performance, a DNA-based strategy was also described by the chronocoulometric response of [Ru(NH$_3$)$_6$]$_{3+}$ absorbed on the DNA-AuNPs that linked with the phosphorylated peptide by Zr$^{4+}$ [132]. Despite the improvement of these methods, it still remains great challenge to develop sensitive, accurate and rapid methods for the profiling of kinase activity and inhibition. Most recently, Wang et al. [133] built a highly sensitive EC sensor for monitoring kinase activity based on DNA induced AuNPs polymeric network block signal amplification. In this strategy, the current signal of EC sensor was significantly amplified to afford a highly sensitive electrochemical analysis of kinase activity, due to its excellent electro activity and high accommodation of the DNA AuNPs polymeric network block for [Ru(NH$_3$)$_6$]$_{3+}$. Notably, the proposed EC sensor presented a low LOD of 0.03 U mL$^{-1}$ for protein kinase A (PKA) activity and excellent stability even in cell lysates and serum samples. Jing et al. [134] developed a sensitive and selective electrochemical method for detection of DNA methylation as well as determination of DNA methyltransferase (MTase) activity and screening of MTase inhibitor. In this work, methyleneblue (MB) was employed as electrochemical indicator and DNA-modified AuNPs were used as signal amplification unit because the DNA strands in this composite had strong adsorption ability toward MB (Figure 8). The differential pulse voltammetry signal demonstrated a linear relationship with logarithm of DNA methylation concentration ranging from 0.075 to 30 U/mL, achieving a LOD of 0.02 U/mL.

Overall, the LSPR feature and excellent biocompatibility of AuNMs provide great chance for improving the analytical performance of EC sensors. Currently, only spherical AuNPs have been extensively applied into fabrication of EC sensor. Certainly, other shaped AuNMs possesses specific physicochemical property, and may offer additional application option in signal amplification of ECL or EC sensors.

**Conclusion**

1. A series of novel and sophisticated synthesis methods for AuNMs have been developed, but how to precisely control the monodispersed size and morphology and to achieve high quality of AuNMs with vivid color (LSPR) is still a crucial but challengeable issue for their application in analytical science. The AuNMs with precisely-controlled vivid color possess abundant optical information and can encode biological or chemical recognition units to develop robust analytical methods in food safety, clinic diagnosis and so on. In addition, we are far from controllable assembly of AuNMs into the desirable structures of the collective properties [135].

2. AuNMs have shown excellent performance in enhancement of SERS signal to improve detection sensitivity. Therefore, AuNMs in combination with other functional metal or organic nanomaterials, such as Si, Al, MOFs and so on, to develop multimodal composite nanomaterials become a new trend in high sensitive SERS detection of various target analytes [136,137], as well as in high resolution bioimaging in the future.

3. AuNMs offer a rapid, efficient, cost-effective and robust sensing platform for detection of different chemicals and bio-markers due to their unique chemical, physical and optical properties. It is noted that the sensitive, stable and multiplex assay of target analyte is highly desirable. To meet these requirements, new multifunctional AuNMs must be developed and used as LFICA label, colorimetric sensing readout and signal amplification of EC sensor, etc.

4. AuNCs as excellent fluorescent probe materials have shown great potential in analytical science [138]. However, extensive efforts must be made for developing novel synthesis methods to achieve high quality AuNCs with remarkable QY and extraordinary stability. In addition, the coordination nature of the AuNCs-protection group complexes implies that the stability of metal NCs is affected by the presence of other strong competing ligands in aqueous solution. The intracellular stability of AuNCs hence becomes a serious issue. With
development of technologies of synthesis and surface modification, AuNMs will be widely employed as alternatives to conventional fluorophores in analytical science including biosensors, bioimaging and so on.

5- We notice that though many new analytical methods based on AuNMs have been developed in the laboratories, such materials are not used as much in an industrial setting. There must be more scope for these platforms to improve, for example, cost of preparing the AuNMs based sensors, reproducibility of different batches for sensor production, and stability of long-term storage of the AuNMs based sensors, and so on.

6- AuNMs with excellent features have prompted great development of analytical sciences and have stood at a critical juncture, with a vast amount of researches that form a solid foundation for future work. Great deals of efforts need to be paid to push on to take AuNMs-based analytical technologies from lab to market.

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