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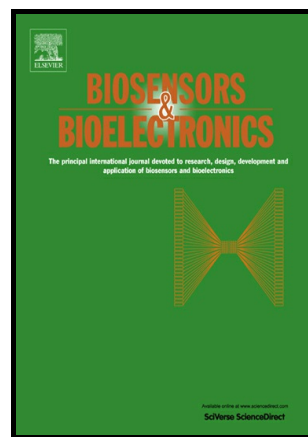
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Endonuclease Controlled Aggregation of Gold Nanoparticles for the Ultrasensitive Detection of Pathogenic Bacterial DNA

Claire McVey^a, Fumin Huang^b, Christopher Elliott^a, Cuong Cao^{a*}

^aInstitute for Global Food Security, School of Biological Sciences, Queen's University Belfast, 18-30 Malone Road, Belfast, BT9 5BN, UK

^bSchool of Mathematics and Physics, Queen's University Belfast, Belfast, BT7 1NN, UK

*Corresponding author: Institute for Global Food Security, Northern Ireland Technology Centre (NITC), 18-30 Malone Road, Belfast, BT9 5BN, United Kingdom;

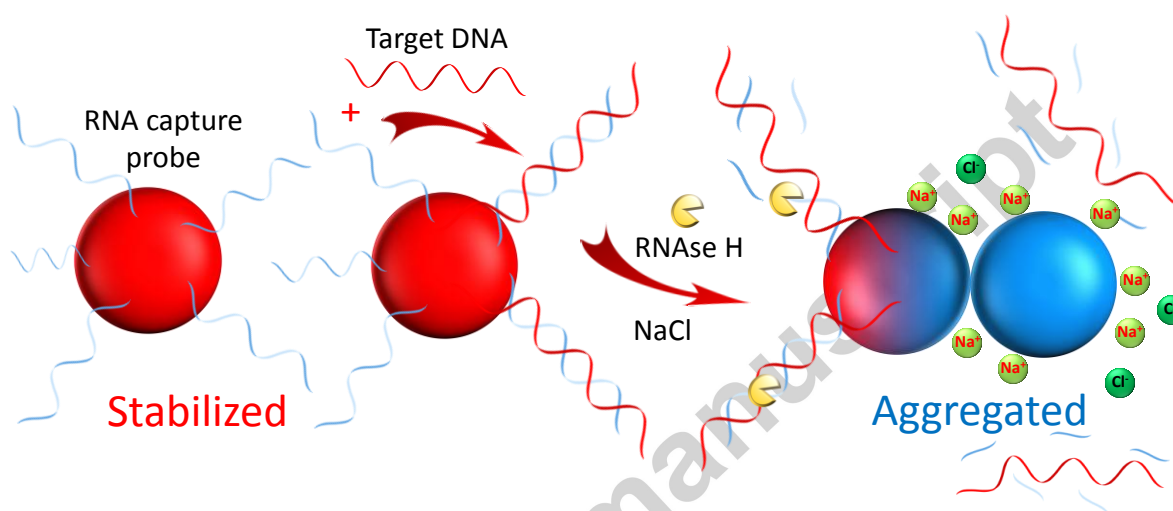
Telephone: +44 (0)28 9097 6545; Email: c.cao@qub.ac.uk

Abstract

The development of an ultrasensitive biosensor for the low-cost and on-site detection of pathogenic DNA could transform detection capabilities within food safety, environmental monitoring and clinical diagnosis. Herein, we present an innovative approach exploiting endonuclease-controlled aggregation of plasmonic gold nanoparticles (AuNPs) for label-free and ultrasensitive detection of bacterial DNA. The method utilizes RNA-functionalized AuNPs which form DNA-RNA heteroduplex structures through specific hybridization with target DNA. Once formed, the DNA-RNA heteroduplex is susceptible to RNase H enzymatic cleavage of the RNA probe, allowing the target DNA to liberate and hybridize with another RNA probe. This continuously happens until all of the RNA probes are cleaved, leaving the nanoparticles unprotected and thus aggregated upon exposure to a high

electrolytic medium. The assay is ultrasensitive, allowing the detection of target DNA at femtomolar level by simple spectroscopic analysis (40.7 pM and 2.45 fM as measured by UV-vis and dynamic light scattering (DLS), respectively). The target DNA spiked food matrix (chicken meat) is also successfully detected at a concentration of 1.2 pM (by UV-vis) or 18.0 fM (by DLS). In addition to the ultra-high sensitivity, the total analysis time of the assay is less than 3 hours, thus demonstrating its practicality for food analysis.

Graphical abstract



Keywords

RNase H, gold nanoparticles, aggregation, DNA detection, endonuclease enzyme

1. Introduction

The on-site and sensitive detection of pathogens is of critical importance to the prevention, surveillance and control of infectious diseases and their outbreak at the first onset (Rhode *et al.*, 2015;

Call *et al.*, 2003; Mothershed and Whitney, 2006). While conventional techniques such as plate culturing, polymerase chain reaction (PCR) and enzyme-linked immunosorbent assay (ELISA) have been used as the predominant detection workhorses, they are limited by either time-consuming and extensive procedure, expensive analysis and operation, or inability to be implemented at point-of-testing (Fratamico, 2003; Brooks *et al.*, 2004; Maurer, 2011; Laurie and Mariani, 2009). Significant efforts have been made to improve the limitations associated with conventional techniques (Lazcka *et al.*, 2007; Baron, 2011; Law *et al.*, 2015). Among these, gold nanoparticles (AuNPs) have emerged as an excellent candidate for biosensor design owing to their unique properties. For example, colloidal AuNPs exhibit distinct colours and strong absorption bands in the visible range of the electromagnetic spectrum that are not present in the bulk metal. This fascinating optical phenomenon of AuNPs is derived from localized surface plasmon resonance (LSPR), a collective oscillation of free electrons in tandem with the incoming photon frequency. This has provided a range of simplified transducing mechanisms for biosensor design, based on assembly, disassembly, or enlargement of the AuNPs which allow scanometric, colorimetric or even naked-eye determination (Cao *et al.*, 2011; Taton *et al.*, 2000; Lee *et al.*, 2007; Zayats *et al.*, 2005; Willner *et al.*, 2006; Pingarron *et al.*, 2008; Verma *et al.*, 2015). Mirkin and co-workers have pioneered and popularized the incorporation of nucleic acid-modified AuNPs into biological sensing platforms to provide improved sensitivity, versatility and portability. Remarkably, the nucleic acid functionalized AuNPs not only provide further functionalities such as specific programmable assembly upon hybridization with their complementary counterparts, but also allow enzymatic cleavage, ligation and extension reactions for biosensor development (Li and Lu, 2000; Palchetti and Mascini, 2008; Lu *et al.*, 2011; Zhao *et al.*, 2008; Thaxton *et al.*, 2006; Wu *et al.*, 2016).

Toward this end, studies have focused on incorporating nuclease enzymes and deoxyribozymes (DNAzyme) to cleave or link oligonucleotides to induce a colorimetric response. DNA endonuclease (DNase I), Pb²⁺-dependent RNA-cleaving DNAzyme (DNAzyme 8-17), exonuclease III (Exo III) and RNase H have been used successfully for the detection of Pb²⁺, nucleic acids and folate receptor (Zhao *et al.*, 2008; Li *et al.*, 2012; Xu *et al.*, 2009; Cui *et al.*, 2011; Yang and Gao, 2014; Kim *et al.*, 2007; Liu and Lu, 2003; Huang *et al.*, 2012; Fan *et al.*, 2012; Wang *et al.*,

2011; Kanaras *et al.*, 2007; Kanaras *et al.*, 2007). Zhao and colleagues (2008) utilized DNAzyme 8-17, which cleaves the DNA substrate with a single RNA linkage in the presence of Pb^{2+} for the detection of metal ions. In a different approach, incorporating the same 8-17 enzyme, Liu and Lu (2003) reported the cross-linking of enzyme-substrate and subsequent cleavage and dissociation of AuNPs upon the addition of target analyte (Pb^{2+}). Several groups have also focused on Exo III enzyme which catalyzes the stepwise removal of mononucleotides from blunt or recessed 3'-hydroxyl terminus of duplex DNA (Cui *et al.*, 2011; Yang and Gao, 2014; Huang *et al.*, 2012; Fan *et al.*, 2012). A universal platform was developed by Cui *et al.* (2011) for the detection of DNA based on Exo III signal amplification. Furthermore, Yang *et al.* (2014) utilized Exo III for the colorimetric detection of folate receptor, in which the target induced AuNP aggregation. The utilization of Exo III enzyme has proven highly sensitive due to repeated hybridization and hydrolysis reactions. In a different approach, Wang *et al.* (2011) modified AuNPs with EcoRI enzyme and designed a specific, double stranded DNA probe which contained an EcoRI recognition site and complementary sticky ends. AuNP aggregation occurred in the presence of the target (magnesium and phosphate ions), resulting in a colorimetric response. Although highly successful, such enzymatic approaches are limited by the need for restriction binding sites, extensive probe design, and requirement for further amplification steps. For a comparative overview of DNA detection, using enzyme-assisted, gold nanoparticles-based transducing platforms see Table SI. 1.

Herein, we present an innovative sensing technology based on the unique enzymatic activity of RNase H for the detection of bacterial DNA at concentrations down to femtomolar level. Due to the ubiquitous nature and high levels found in food, especially poultry, *Campylobacter jejuni* was chosen as the target for assay development (EFSA, 2011; WHO, 2011). The method utilizes RNA-functionalized AuNPs which form DNA-RNA heteroduplex structures through specific hybridization with target DNA. Once formed, the DNA-RNA heteroduplex is susceptible to RNase H enzymatic cleavage of the RNA probe, allowing DNA to liberate and hybridize with another RNA strand. This continuously happens until all of the RNA strands are cleaved, leaving the nanoparticles unprotected and prone to aggregation upon exposure to a high electrolytic medium. The current work overcomes previous limitations associated with enzyme-based methods in that it does not require further

amplification steps. In addition, RNase H is not active on single stranded DNA or RNA molecules and only catalyzes the cleavage of RNA within a DNA-RNA heteroduplex, thus does not require specific recognition sites for enzymatic cleavage (Tadokoro and Kanaya, 2009). Furthermore, there is greater versatility and applicability with regard to probe design and thus potential for multiplexing. RNase H has previously been used for the detection of DNA *via* RNA cleavage within a DNA-RNA heteroduplex structure and subsequent release of a fluorescence dye to generate a fluorescence signal (Kim *et al*, 2007). The method has a limit of detection (LOD) of 10 pM, which highlights the ultra-sensitivity of our method which can determine target DNA at 1 pM by naked eye, or even down to femtomolar level by spectroscopic analysis. The fluorescence-based approach is further limited by cost due to the synthesis of fluorescein conjugate and requirement for equipment capable of detecting the fluorescence signals. Our work is significantly different from previous reports as it utilizes the plasmonic properties of AuNPs to produce a red-to-blue colorimetric response, thus the signal can be visibly detected by the naked eye. In addition, DNA detection can be performed at isothermal conditions in less than three hours. These advantages provide a basis for eradicating the need for a thermal cycler, complicated sample preparation, labelled fluorophore and expensive and cumbersome read-out equipment. Finally, the application to a food matrix has also been assessed and it is evident that the sensitivity and robustness of the assay is conducive for food safety analysis.

2. Experimental Section

2.1 Reagents and chemicals

Aminated RNA probe (5'-Amino-C6-AGG UGU GGA CGA CGU CAA GUC AUC AUG-3'), complementary DNA probe (5'-CAT GAT GAC TTG ACG TCG TCC ACA CCT-3'), non-complementary DNA probe 1 (5'-CCA ACC CCC CAG AAA GAA-3') and non-complementary probe 2 (5'-TCT ATT GGT GGT AAA ACT TAC GCT GCA AGT AAA GCC GAA GGT CAC -3') were purchased from Eurofins Genomics (Ebersburg, Germany). RNase H enzyme was purchased from Takara Bio (France), and thioctic acid NHS ester was obtained from Link Technology Ltd. (UK). Sodium citrate tribasic dehydrate ($\text{HOC}(\text{COONa})(\text{CH}_2\text{COONa})_2 \cdot 2\text{H}_2\text{O}$), gold (III) chloride (HAuCl_4), dimethyl sulfoxide (DMSO) ($(\text{CH}_3)_2\text{SO}$), Tween 20, sodium chloride (NaCl), sodium carbonate

(Na_2CO_3), sodium bicarbonate (Na_2HCO_3), triethylammonium acetate buffer (TEAA), sodium phosphate (NaH_2PO_4 and Na_2HPO_4), glutathione (GSH), Tris-EDTA buffer (TE), Tris-HCl ($\text{NH}_2\text{C}(\text{CH}_2\text{OH})_3$), hydrochloric acid (HCl), potassium chloride (KCl), magnesium chloride (MgCl_2) and dithiothreitol (DTT) were purchased from Sigma-Aldrich (UK). All reagents were prepared in RNase free water (Sigma, UK). NAP-5 column was obtained from GE Healthcare (UK). Syringe filters (0.22 μm) were purchased from Merck Millipore (Germany).

2.2 Gold nanoparticle synthesis

In a typical experiment, 25 mM HAuCl_4 was dissolved in 150 mL of deionized H_2O (dH_2O) and heated to reflux under constant stirring ($t \approx 10$ minutes). 1 mL of 2.4 mM sodium citrate was quickly injected and the solution was removed from the heat upon the color changed from translucent yellow to wine red. The pH of the solution was adjusted to 6.5 using HCl.

2.3 RNA preparation

Prior to functionalisation the RNA was modified with thiotic acid at the 5' end (Dogan *et al*, 2007). The dried RNA was incubated with 30 μL of 80 mM thiotic acid (dissolved in DMSO) and 75 μL of 0.1 M $\text{Na}_2\text{CO}_3/\text{NaHCO}_3$ (pH 9.75) overnight at room temperature. The modified RNA was desalted using a NAP-5 column in 0.1 M TEAA buffer. The concentration was determined using Nanodrop 8000 (Thermoscientific, UK).

2.4 Gold nanoparticle functionalization

Gold nanoparticles were functionalized with the as-prepared RNA. Typically, 300 μL of 30 μM modified RNA was added to 1 mL of AuNP solution. The solution was incubated overnight ($t \approx 16$ hours) at room temperature and stability checked with an equal volume of 4 M NaCl to ensure successful conjugation. To improve the orientation and loading of RNA conjugated onto the surface of the AuNP, the salt concentration was slowly increased (Hurst *et al*, 2006). Firstly, phosphate buffer (60 mM) was diluted to a final concentration of 10 mM. NaCl was added in small increments (0.05 M) each hour, over six hours (0.3 M). The conjugate was incubated overnight ($t \approx 16$ hours) at room

temperature. Finally, the conjugate was centrifuged twice at 13,000 g for 1 hour to remove unbound RNA and re-suspended in TE buffer (stored at 4°C).

2.5 Determination of AuNP-RNA stability

In order to ensure the colloidal stability of the RNA-functionalized AuNPs at high salt concentrations, the conjugates were exposed to different concentrations of NaCl (Figure SI. 2). The functionalized particles exhibit excellent stability at 2 M NaCl, with no wavelength shift noted. The bare-AuNPs immediately aggregated at 0.5 M NaCl, due to the large screening effect of NaCl, causing a red-shift to longer wavelengths on the absorbance spectrum (λ_{\max} shift > 200 nm). This stability can be attributed to electrostatic repulsion or steric exclusion caused by RNA on the AuNP surface. From this analysis, successful functionalisation was confirmed and 2 M NaCl was set as the highest concentration to induce aggregation of AuNPs over varying stabilities.

2.6 Colorimetric detection of pathogenic bacterial DNA

Prior to analysis, the AuNP conjugate was centrifuged (13,000 g for 30 minutes) and re-suspended in a modified Tris-HCl buffer (20 mM Tris, 40 mM KCl, 8 mM MgCl₂ and 1 mM DTT). 30 μ M GSH was also added to this buffer due to its role in aiding RNAse H enzyme activity (Lyles and Gilbert, 1991). In a typical experiment 20 μ L of AuNP-RNA was added into an Eppendorf tube with 10 μ L of target DNA or control (modified Tris-buffer) and 1.5 μ L of Tween 20. The sample was heated to 90 °C for 2 minutes, cooled slowly to 60 °C and incubated at 60 °C for 60 minutes. 0.06 U of RNAse H enzyme, prepared in modified Tris-buffer, was added to a final concentration of 0.02 U. The sample was then incubated for 1 hour at 37 °C. To induce aggregation, NaCl was added into the mixture to a final effective concentration of 2 M.

2.7 Preparation of DNA in chicken matrix

50 g of skinless chicken meat (breast) was shaken for 2 minutes in 100 mM Tris-HCl (pH 7). The subsequent matrix was filtered and diluted 1/100 in modified Tris-HCl buffer (20 mM Tris, 40

mM KCl, 8 mM MgCl₂ and 1 mM DTT). This matrix was then used to prepare a 10-fold dilution of target DNA ranging from 0 to 10 μM concentration.

2.8 Analysis instrumentation

All Ultraviolet-visible spectrophotometry (UV-vis) measurements were carried out using a Cary 60 spectrophotometer (Agilent Technologies, USA). AuNP size analysis was carried out using a Zetasizer NanoZS (Malvern, UK). Transmission electron microscopy (TEM) characterization was acquired using a Phillips CM100 (Phillips, USA) operated at 100 kV.

3. Results and discussion

3.1 Working principle of the colorimetric assay

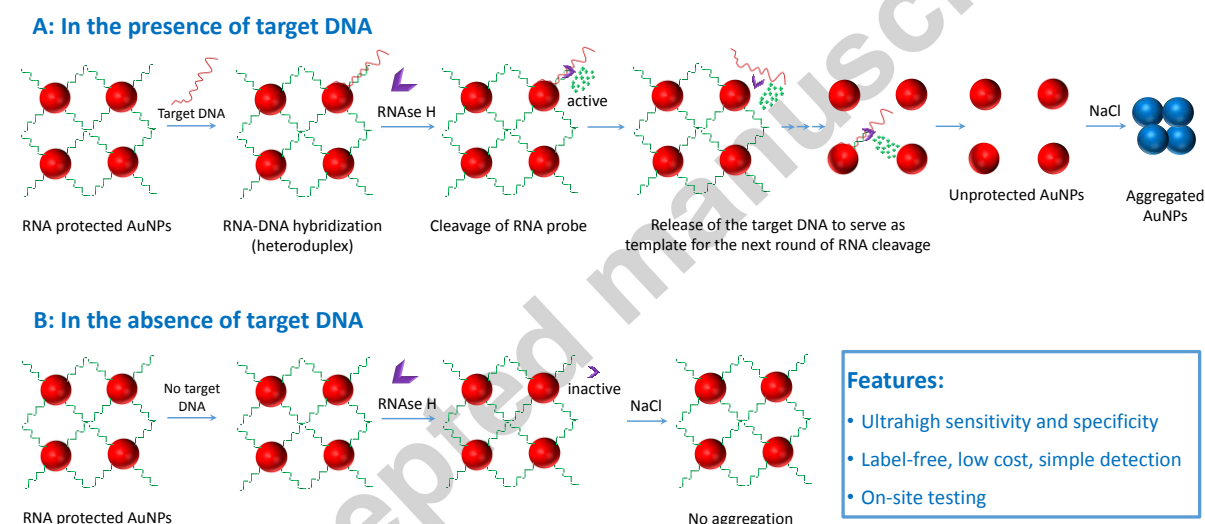


Figure 1. Overall scheme demonstrating the colorimetric detection of target DNA based on DNA-RNA hybridization and enzyme controlled cleavage and aggregation of AuNPs upon the addition of NaCl. (A) In the presence of target DNA, DNA-RNA hybridization occurs which initiates RNase H enzyme cleavage of RNA within the heteroduplex structure. The target DNA recycles until all of the RNA is cleaved, allowing for subsequent AuNP aggregation in the presence of NaCl. (B) In the absence of target DNA, no hybridization occurs, thus there is no heteroduplex structure to act on and the RNase H enzyme is inactive. Therefore the particles remain stable upon the addition of NaCl.

AuNPs (17 ± 3 nm, Figure SI. 3) were first synthesized by a method reported previously with minor alterations (Turkevich *et al.*, 1951) and exhibited a typical UV-vis absorbance band at 520 nm. Subsequently, a single-stranded RNA probe (5'-Amino-C6 -AGG UGU GGA CGA CGU CAA GUC AUC AUG -3') was designed to recognize a DNA fragment of *Campylobacter jejuni*, (NCTC 11168 = ATCC 700819 chromosome, 5'-CAT GAT GAC TTG ACG TCG TCC ACA CCT-3'). The RNA probe was successfully crafted onto the AuNP *via* an N-hydroxysuccinimidyl (NHS) ester of thioctic acid by a method reported by Dougan *et al.* (2007). The AuNP-RNA conjugate exhibits a deep red colour, and absorbance peaks at 525 nm and 260 nm (Figure SI. 4), which represents the typical optical absorption of AuNPs and RNA, respectively. The shift in LSPR peak from 520 nm (bare nanoparticles), to 525 nm with the AuNP-RNA conjugates further demonstrates the successful functionalisation. The RNA-functionalized AuNPs prepared by this method show excellent stability, their plasmonic and biological properties could be retained upto 4 weeks with storage in TE buffer, at 4°C. In the presence of target *Campylobacter jejuni* DNA, hybridization occurs with RNA functionalized onto the AuNP surface. The subsequent DNA-RNA heteroduplex becomes a target for cleavage of the RNA probe *via* RNase H enzyme, allowing the DNA to liberate and hybridize with another RNA strand (Figure 1, overall scheme). This happens isothermally and iteratively until all of the RNA probes are cleaved, leaving the nanoparticles denuded. The addition of 2 M NaCl causes the denuded nanoparticles to aggregate in solution, initiating a colour change from red to blue. The aggregation state was confirmed in Figure 2 by the transmission electron microscopy (TEM) analysis of RNA-functionalized AuNPs under assay conditions in the absence and presence (1 μ M) of target DNA. The distinct colour change generated from the assay can be detected by the unaided eye, or by simple spectroscopic analysis.

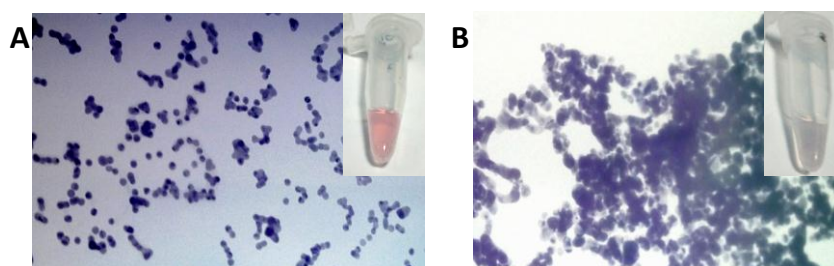


Figure 2. Transmission electron microscopy (TEM) images and photographs of the colorimetric response of AuNPs in the (A) absence and (B) presence of target DNA (1 μM) post assay conditions. Scale bar 100 nm.

3.2 Sensitivity of assay

To test the hypothesis of the assay, a 10-fold dilution of target DNA ranging from 0 to 10 μM concentration was analyzed. Figure 3A demonstrates the visible colour change of AuNP-RNA solution from red to blue or grey, with increasing target DNA concentration. The lowest target DNA concentration easily determined by the naked eye was 10^{-6} μM , as seen by a colour change from red to light blue, caused by a decrease in interparticle distance. More quantitatively, UV-vis absorption measurements were performed to determine the LSPR shift as a function of target DNA concentration (Figure 3B). The results demonstrate a shift in wavelength from a target DNA concentration of 10 fM, which continues to increase to 10 μM (Figure 3C). A good linear relationship between the LSPR shift and target DNA concentration (Figure 3C, inset) could be obtained for a range between 10 fM and 10 μM of target DNA ($R^2 = 0.93$). The limit of detection (LOD) is defined as the lowest DNA concentration with a response three times greater than the standard deviation (SD) of the blank sample. Owing to the excellent stability of the AuNP-RNA conjugate, the SD value of the zero concentration for ten measurements ($n = 10$) was as small as 1.9 nm, thus the LOD of the current assay is 40.7 fM of pathogenic bacterial DNA. The LOD is about two orders of magnitude more sensitive in comparison to a similar method utilizing fluorescein (Kim *et al.*, 2007) and other AuNP-functionalized based assays for the detection of DNA (Cui *et al.*, 2011; Fan *et al.*, 2012).

During the UV-vis analysis, we observed that the aggregated AuNPs had latched on to the wall of the Eppendorf tube or settled down to the bottom of the tube, potentially causing only free and partially aggregated particles to be measured in solution. Thus, it is evident that the LSPR shift (Figure 3B) is not as predominant when compared to the colorimetric response as seen in the colour photographs. Therefore, to accurately reflect the aggregation states of the AuNPs, dynamic light scattering (DLS) measurements were carried out to determine the average size of the AuNP-conjugates in the presence varying target DNA concentrations. DLS data is displayed as the integrated value of three measurements derived from one sample. The polydispersity index (PDI) for each sample is also

displayed which indicates the variation of nanoparticle size within a distribution. The PDI is calculated from the distribution width and mean, giving an overall indication of the non-uniformity of particles within a sample. The DLS results show an average increase in AuNP size, from 44.4 nm (± 1.2 nm) with the zero concentration sample to 854.0 nm (± 36.0 nm) at 1 μM target DNA concentration, which demonstrates increased aggregation with increasing target DNA concentration (Figure 3D). Furthermore, the size distribution charts show a shift in average size from one population, between 10 – 100 nm to a second, increasing population (100 to 1000 nm). This large size distribution can be explained by the highly sensitive nature of DLS analysis which is capable of determining the true state of particles in a given media, compared to TEM which only measures the solid state (nanoparticles are dried on grid subsequent to analysis). Furthermore, DLS measures the hydrodynamic diameter of the particle, which in this case includes the metallic core and RNA functionalised onto the surface, thus it is obvious that we should see this wide distribution (Domingos *et al*, 2009). The PDI data demonstrates broad polydispersion which increases from 0 μM to 10^{-3} μM target DNA concentration, where the index value decreases. This indicates a wide size distribution at lower target DNA concentrations, with narrower size distributions at higher concentrations. This coincides with the distribution data which indicates that there are a greater number of larger or aggregated particles at a high target DNA concentration, in contrast with lower target DNA concentrations which demonstrate a larger size frequency over several distribution sizes (i.e. 10-100 nm and 100-1000 nm). Figure 3E demonstrates the linear fitting of AuNP size increase as a function of target DNA concentration ($R^2 = 0.98$). This data provides a dynamic range of between 1 fM and 100 pM of target DNA. From this data we can determine a LOD of 2.45 fM which is not clearly discriminable to the naked-eye or detectable using UV-vis spectrophotometry. A baseline has been included in Figure 3E to remove background signal as determined by specificity and selectivity analysis.

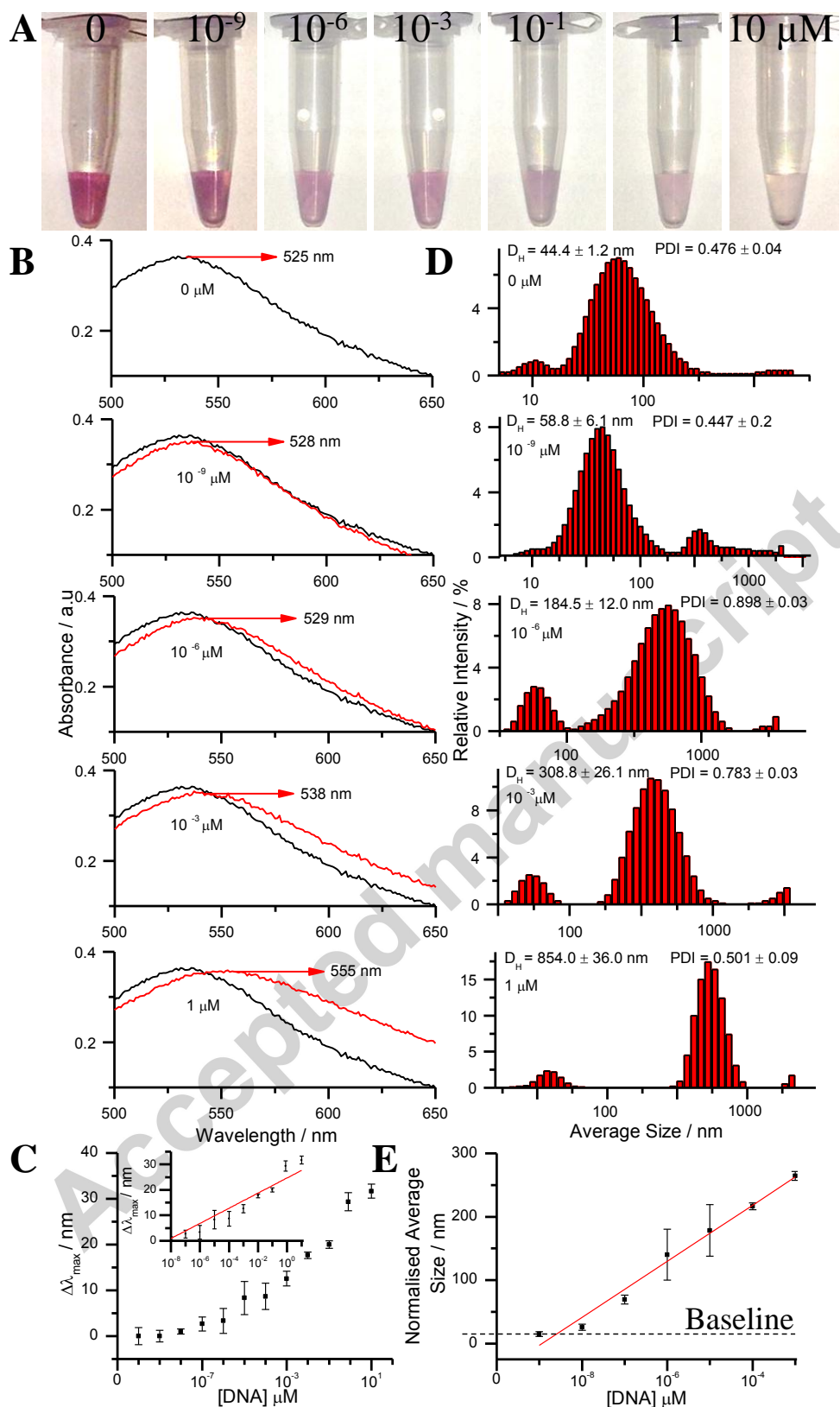


Figure 3. Analysis of AuNP-RNA conjugates in the presence of increasing concentrations of target DNA. (A) A representative colour photograph showing a visual colour change from red to blue or transparent with increasing DNA concentrations. (B) UV-vis absorption spectra demonstrating a red

shift towards longer wavelengths (525 nm to 555 nm) in the presence of increasing target DNA concentrations. (C) Wavelength shift of the LSPR peak as a function of target DNA concentration ($n = 3$). Inset: the linear relationship between target DNA concentration and maximum wavelength shift ($R^2 = 0.93$) (D) Hydrodynamic size distribution of AuNPs measured by dynamic light scattering (DLS) technique (inset PDI). (E) Linear relationship between target DNA concentration and average increase in AuNP size derived from DLS analysis ($R^2 = 0.98$).

3.3 The selectivity of the assay

Control experiments were carried out with RNase H enzyme removed to determine if nanoparticle aggregation was fully attributed to the enzymatic cleavage of RNA (target DNA concentration was fixed at 10 μ M). In the absence of target DNA but in the presence of RNase H enzyme, the average AuNP size was determined to be 44.4 ± 1.2 nm (Figure 4A) with no visible change in colorimetric response (Figure 4B) and no shift in LSPR peak (maximum absorbance peak at 525 nm; Figure 4C). In the absence of RNase H but in the presence of 10 μ M target DNA, the hybridization of the RNA-DNA resulted in an increase in the average size of the nucleotide-AuNP complex of 24.8 nm as compared to the zero (no target) sample (Figure 4A). The UV-vis results further demonstrate that the removal of RNase H enzyme results in almost no change in the LSPR peak shift. This result confirms that the addition of RNase H enzyme is essential in controlling aggregation of the AuNPs through cleavage of functionalised RNA.

Control experiments were also carried out with two non-complementary DNA sequences at a concentration of 10 μ M. The DLS results demonstrate a slight increase in particle complex size with a maximum value of 9.6 nm (Figure 4A), attributing to the background noise of the assay (also denoted as the baseline on Figure 3E). The colour images in Figure 4B along with the UV-vis results (Figure 4C) demonstrate no visual or spectroscopic change in colour or LSPR peak under the test experiments. The high specificity of the assay can be credited to the specific DNA-RNA recognition and hybridisation, as well as subsequent highly selective RNase H enzyme cleavage of the RNA in the heteroduplex.

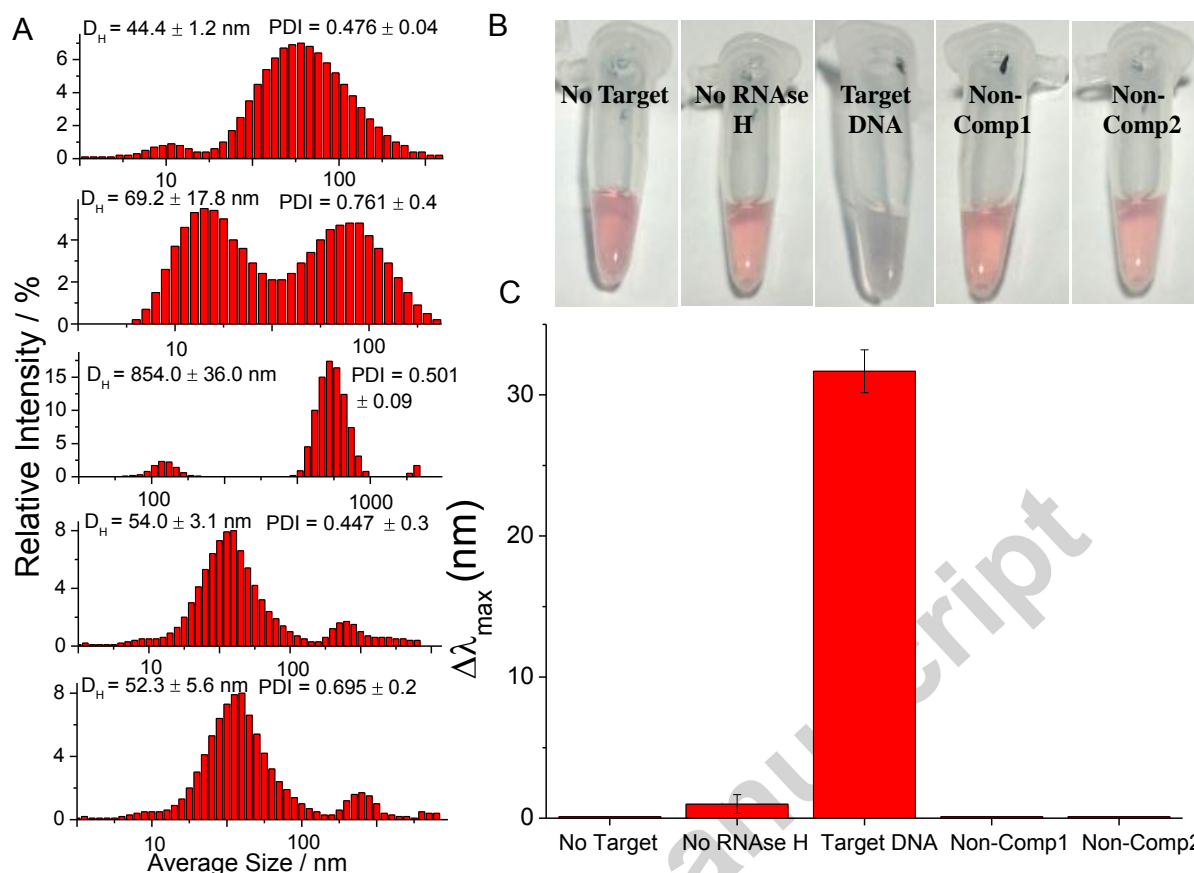


Figure 4. Illustrating the assay specificity. (A) DLS measurements demonstrating the hydrodynamic size distribution of AuNP-RNA post assay conditions. From top to bottom: RNase H enzyme and 0 μ M target DNA; no RNase H enzyme and 10 μ M target DNA; RNase H enzyme and 1 μ M target DNA; RNase H enzyme and 10 μ M non-complementary DNA sequence 1; and RNase H enzyme and 10 μ M non-complementary DNA sequence 2. (B) and (C) are the corresponding colour photograph and maximum wavelength shift of LSPR peak under different conditions ($n = 3$), respectively.

3.4 Application to real sample matrices

The applicability of the assay was tested by preparing a 10-fold dilution of target DNA ranging from 0 to 10 μ M in chicken matrix. The aim of this experiment was to determine if the assay could be applied to a food matrix, given the complex medium and thus potential interferences and inhibitors that could cause deterioration of assay parameters such as reduced sensitivity and specificity. Figure

5A shows the maximum LSPR shift, as a function of target DNA in chicken matrix. The results demonstrate good linearity ($R^2 = 0.96$) spanning from 10 fM and 10 μ M. In the complex environment, the spiked target DNA was identified at a concentration as low as 1.2 pM based on the LSPR analysis. It was noted that the LOD had increased two orders of magnitude in comparison to buffer conditions and there was an overall decrease in maximum LSPR shift (15 nm), which might be attributable to interferences of the enzymatic reaction within the sample. Furthermore, parallel DLS measurements were carried out to determine the aggregation states of AuNPs post analysis in chicken matrix. Figure 5B demonstrates the linear correlation between normalized average size (nm) and DNA concentration (μ M). The linear range of the assay was determined to be between 10 fM and 1 μ M, with a limit of detection of 18.0 fM. In addition, the commercial availability of portable and battery operated UV-Vis spectrophotometers and particle size analyzers makes on-site detection using the developed assay highly possible in case where improved sensitivity is required beyond that of naked eye detection (Ocean Optics, 2016; Xylem, 2016; TSI, 2016). Thus, the assay shows potential application to food analysis.

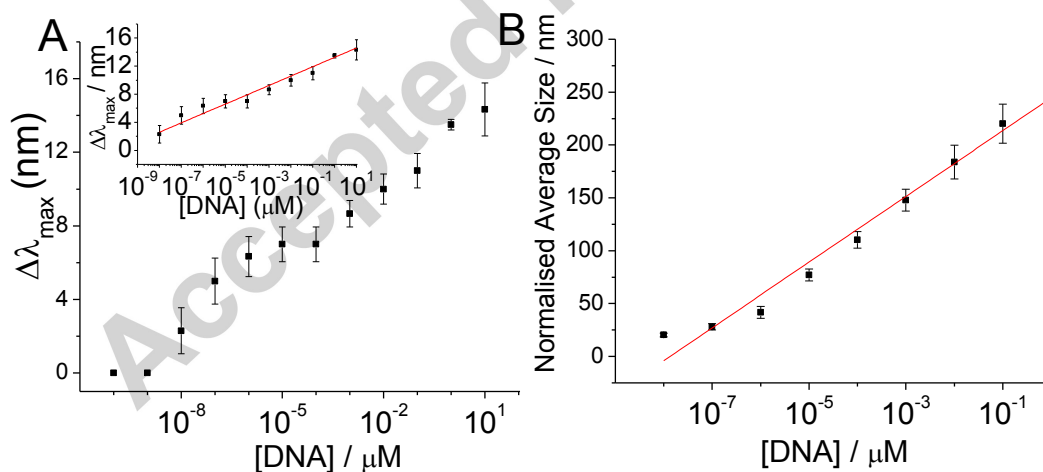


Figure 5. (A) Wavelength shift of the LSPR peak as a function of target DNA concentration analyzed in chicken matrix, demonstrating the sensitivity of the assay. Inset is the linear relationship between target DNA concentration spiked in chicken matrix and maximum wavelength shift ($R^2 = 0.96$). (B)

Linear relationship between target DNA prepared in chicken matrix and normalized, average increase in nanoparticle size determined by dynamic light scattering technique (DLS) ($R^2 = 0.98$).

4. Conclusion

In conclusion, we have presented a highly sensitive and selective method for the detection of DNA, based on endonuclease controlled aggregation of plasmonic AuNPs. RNase H enzymatic cleavage in combination with DNA-RNA hybridization provides a highly specific and ultra-sensitive assay, which can detect 1 pM target DNA concentration visibly or down to femtomolar level by spectroscopic techniques (40.7 fM and 2.45 fM as measured by UV-vis and dynamic light scattering (DLS), respectively). The detection capabilities within a food matrix show good sensitivity (1.2 pM and 18.0 fM as analyzed by UV-vis and DLS, respectively). In addition to the ultra-high sensitivity, the total analysis time of the assay is less than 3 hours, thus demonstrating its practicality for food analysis. The versatility of probe design and enzyme cleavage offers a broad range of potential applications for the detection of DNA. Future work will focus on further applications of the method and potential multiplexing capabilities for real clinical and veterinary samples.

Conflict of interest

The authors have no conflict of interest to declare

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Supporting Information

Comparative overview of DNA detection, using enzyme-assisted, gold nanoparticle-based transducing platform (Table SI. 1.); Comparison of the stability of bare and functionalized gold nanoparticle (AuNP) (Figure SI. 2); AuNP characterization by TEM (Figure SI. 3); Functionalisation of AuNPs with RNA (Figure SI. 4). Information is available from the Elsevier Online Library or from the author.

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Highlights

- RNA-functionalized AuNPs is proposed for the detection of bacterial DNA
- Successive hybridization and cleavage reactions between target DNA and RNA by RNase H
- Resulting in unprotected-AuNPs, which are susceptible to aggregation upon exposure to NaCl
- The method allows naked eye determination of target DNA at concentrations as low as 10^{-6} μM
- Ultra-high sensitivity at femtomolar level (2.45 fM) can be achieved