

# Diagnosis of rubella virus using antigen-conjugated Au@Pt nanorods as nanozyme probe

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**Background:** As nanotechnology advances rapidly, the nanozymes which could replace protein enzymes in bioanalytical assays bring a new opportunity to the development of simple and sensitive diagnostic tools.

**Purpose:** Antibody-conjugated Au-Pt core/shell nanorods (Au@Pt NRs) have been used for nanozyme probes for diagnosis of rubella virus. Au@Pt NRs, prepared by Au nanorod-mediated growth, exhibit peroxidase-like activities and could serve as an inexpensive replacement for horseradish peroxidase (HRP) in conjugation of antigen.

**Method:** Using a capture enzyme-linked immunosorbent assay for the determination of rubella virus.

**Results:** Compared with antibody-conjugated HRP, the antigen-conjugated Au@Pt NRs are more stable and more robust, but less expensive. Based on the enhanced catalytic properties of this nanozyme probe, it was found that the antigen-conjugated Au@Pt NRs-based enzyme-linked immunosorbent assay exhibited good sensitivity.

**Conclusion:** Our results indicate that these antigen-conjugated Au@Pt NRs represent a suitable nanozyme probe for future clinical applications under various conditions.

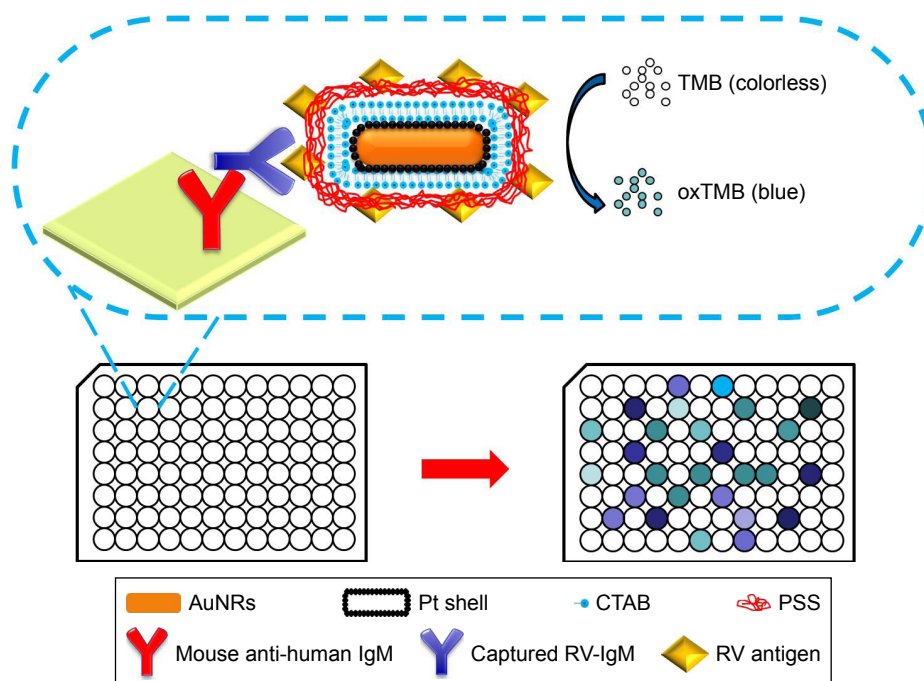
**Keywords:** gold nanorods, nanozyme, antigen conjugation, rubella virus, diagnosis

## Background

Rubella virus (RV) is the causative agent of rubella disease or so-called German measles. Although most cases of infection lead to a mild, self-limiting measles-like disease, the real threat arises when RV infects pregnant women, particularly during the first trimester, when it will cause severe fetal anomalies or even fetal loss.<sup>1–3</sup> For this reason, it is of great practical significance to establish a simple, sensitive detection method for RV. Laboratory confirmation of recently acquired rubella relies commonly on serum detection of RV-specific IgM antibodies. Enzyme-linked immunosorbent assay (ELISA) is an important method for the detection of various proteins in immunologic reactions, finding applications in numerous disciplines including the diagnosis of infectious diseases.<sup>4–6</sup> Reports have demonstrated the efficiency of detecting rubella-specific IgM or IgG antibodies in human serum or plasma using ELISA techniques in indirect or capture format.<sup>7,8</sup> However, the most commonly used enzymes in immunoassay which lose their enzymatic activities gradually over long-term storage include horseradish peroxidase (HRP). To overcome these limits, various studies on replacing natural enzymes have been reported.<sup>9–12</sup> As nanotechnology advances fast, the nanozymes which could replace protein enzymes in bioanalytical assays bring a new opportunity to the development of simple and sensitive diagnostic tools.<sup>13–16</sup> Nanozymes comprise many types of materials, iron oxide, vanadium oxide, cerium oxide, manganese oxide, gold, platinum, palladium, gold–silver alloys, carbon nanotubes, graphene, metal chalcogenides, and other

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**Figure 1** Schematic representation of antigen-conjugated Au@Pt NRs-based ELISA system.

**Abbreviations:** ELISA, enzyme-linked immunosorbent assay; CTAB, cetyltrimethylammonium bromide; NR, nanorod; PSS, poly(sodium-4-styrene sulfonate); TMB, 3,3',5,5'-tetramethylbenzidine; RV, rubella virus.

materials.<sup>17–26</sup> Compared with natural enzymes, nanozymes are advantageous in several aspects, such as low cost, ease of mass production, robustness to harsh environments, high stability, long-term storage, large surface area for further modification and bioconjugation and so on.<sup>27–32</sup>

Previously, we found Au–Pt core/shell nanorods (Au@Pt NRs) have intrinsic multiple enzyme mimetic capability, for example, oxidase, peroxidase, and catalase mimetic activity.<sup>33–37</sup> The as-synthesized Au@Pt NRs are able to catalyze color reactions in immunoassay and, therefore, can be used to replace the enzymes in conventional ELISA. In this work, we designed a novel conjugate based on antigen-conjugated Au@Pt NRs, which was used as nanozyme probe for virus serodiagnosis. We chose Au@Pt NRs based on the following reasons. Small Pt nanoparticles often show high catalytic activity. However, they are also easy to aggregate and a proper support is often needed to keep them in a well-dispersed state. Au NRs are chosen as the support for the Pt nanoparticles due to their well-developed synthesis method. Owing to large cohesive energy, Pt exhibits island growth mode on gold surface. Thus, well-dispersed Pt dots can be obtained using Au NRs as a support and the ligand effect can further enhance the catalytic activity of the Au@Pt NRs. For the surface modification method, rubella antigen is generally conjugated on the surface of the nanozyme to provide specificity toward antibody molecules. Significantly, Au@Pt

NRs with high surface areas offer a possibility to design probes for signal amplification. Using the reaction system of 3,3',5,5'-tetramethylbenzidine (TMB) and  $H_2O_2$ , peroxidase-like activity of antigen-conjugated Au@Pt NRs was investigated. Moreover, compared with antigen-conjugated HRP, antigen-conjugated Au@Pt NRs are more stable and possess an almost unchanged catalytic activity over a wide range of temperatures and pH. Replacing antigen-conjugated HRP, the functionalized nanoprobe were employed to capture ELISA and realized highly sensitive detection (Figure 1). Another study was carried out side by side to compare the performance of the antigen-conjugated Au@Pt NRs-based ELISA to traditional HRP-based ELISA. At the same time, the specificity of the antigen-conjugated Au@Pt NRs-based ELISA for RV-positive serum was also investigated by comparing with other infectious virus. As a new kind of nanozyme probe, antigen-conjugated Au@Pt NRs could be applied in various potential areas of catalysis and nanobiotechnology.

## Materials and methods

### Materials

Sodium borohydride ( $NaBH_4$ ), CTAB, poly(sodium-4-styrene sulfonate) (PSS), chlorauric acid ( $HAuCl_4 \cdot 3H_2O$ ), potassium tetrachloroplatinate(II) ( $K_2PtCl_6$ ), silver nitrate ( $AgNO_3$ ), L-ascorbic acid (AA), 30%  $H_2O_2$ , and TMB were all purchased from Alfa Aesar (Ward Hill, MA, USA) and used as received.

Rubella antigen, mouse anti-human IgM antibodies coated plate, HRP-antigen conjugates, and positive and negative serum samples (ELISA kit) were purchased from Kerunda Bioengineering company (Shenzhen, China). Milli-Q water (18 M $\Omega$  cm) was used for all solution preparations.

## Synthesis of gold NRs

Au NRs were synthesized using a seed-mediated growth procedure. CTAB-capped Au seeds were synthesized by chemical reduction of HAuCl<sub>4</sub> with NaBH<sub>4</sub>. CTAB (7.5 mL, 0.1 M) was mixed with HAuCl<sub>4</sub> (100  $\mu$ L, 24 mM), diluted with water to 9.4 mL, and stirred with a magnetic stirrer. Then, ice-cold NaBH<sub>4</sub> (0.6 mL, 0.01 M) was added. The solution color immediately turned from bright yellow to brown, indicating the formation of seeds. The Au seeds were used within 2–5 hours. Then, 120  $\mu$ L seed solution was added to the growth solution consisting of CTAB (100 mL, 0.1 M), HAuCl<sub>4</sub> (2.04 mL, 24 mM), AgNO<sub>3</sub> (1.05 mL, 10 mM), H<sub>2</sub>SO<sub>4</sub> (2 mL, 0.5 M), and AA (800  $\mu$ L, 0.1 M) to initiate the growth of Au NRs. After 12 hours, the reaction was stopped. The obtained Au NRs were purified by centrifuging the solution at 12,000 rpm for 5 minutes twice. The precipitate was collected and redispersed in deionized water.

## Synthesis of Au@Pt NRs

Three samples of the purified Au NR solutions (1 mL) were mixed with 62.5  $\mu$ L of 2 mM PtCl<sub>4</sub><sup>2-</sup> aqueous solution. Then, 12.5  $\mu$ L of 0.1 M AA was added and the total solution volume was diluted to 2 mL. The mixture was shaken vigorously and then placed in a 30°C water bath for 30 minutes. Within several minutes, the color of the solution changed from pink-red to dark gray, suggesting the formation of a Pt shell. Then, 1 mL of 0.1 M CTAB was added.

## Modification of the Au@Pt NRs with PSS

CTAB-coated NR solution (1 mL, Au@Pt NRs) was centrifuged at 12,000 rpm for 10 minutes, and the precipitate was dispersed in 0.5 mL PSS aqueous solution (2 mg/mL containing 6 mM NaCl). Then, the solution was stirred magnetically for 3 hours. After that, it was centrifuged at 12,000 rpm for 10 minutes, and the precipitate was redispersed in water.

## Preparation of antigen-conjugated Au@Pt NRs

PSS-coated Au@Pt NR solution (50  $\mu$ L, 5 nM) was first dispersed into 1 mL PBS buffer (0.1 M, pH 7.4). Then 50  $\mu$ L of 10 mg/mL rubella antigen was added to the above Au@Pt NRs solution and incubated at 37°C for 96 hours. After incubation,

the mixture was centrifuged at 12,000 rpm/min for 5 minutes twice. Then, the precipitate was collected and redispersed in 100  $\mu$ L PBS buffer (0.1 M, pH 7.4). Finally, it was dispersed in PBS buffer (pH 7.4) at a concentration of 2.5 nM.

## Detection of rubella IgM antibodies by ELISA

Detection of rubella IgM antibodies by ELISA was performed in 96-well polystyrene plates. Each well of the 96-well plates was precoated with mouse anti-human IgM antibodies. Firstly, each well was blocked with 5% bovine serum albumin (diluted in PBS, pH 7.4) for 1 hour at 37°C to avoid the nonspecific interaction with the plate surface. Then, the plates were washed three times with PBS with Tween 20 (PBST) buffer (pH 7.4). After this, 100  $\mu$ L of negative control, positive control, or diluted sample was added to the plate and incubated at 37°C for 0.5 hour. The plates were washed three times with PBST buffer (pH 7.4) to remove the unbound rubella IgM antibodies. Then, 100  $\mu$ L of Au@Pt NR-antigen conjugates was added to each well and incubated for 0.5 hour at 37°C. The plates were washed five times with PBST buffer (pH 7.4) to remove the unbound Au@Pt NR-antigen conjugates. The color development was initiated by adding 100  $\mu$ L substrate solution (1 mM TMB, 50 mM H<sub>2</sub>O<sub>2</sub> in PBS buffer, pH 5) into each well. The reaction was stopped after 10 minutes using 50  $\mu$ L of 2 M H<sub>2</sub>SO<sub>4</sub>. Absorbance was read at 450 nm. The clinical serum samples were selected from patients with clinical signs of rubella infection or from the patients who had been exposed to RV. The clinical serum experiment was checked with positive control, negative control, and the blank. Buffer solution was used as the blank.

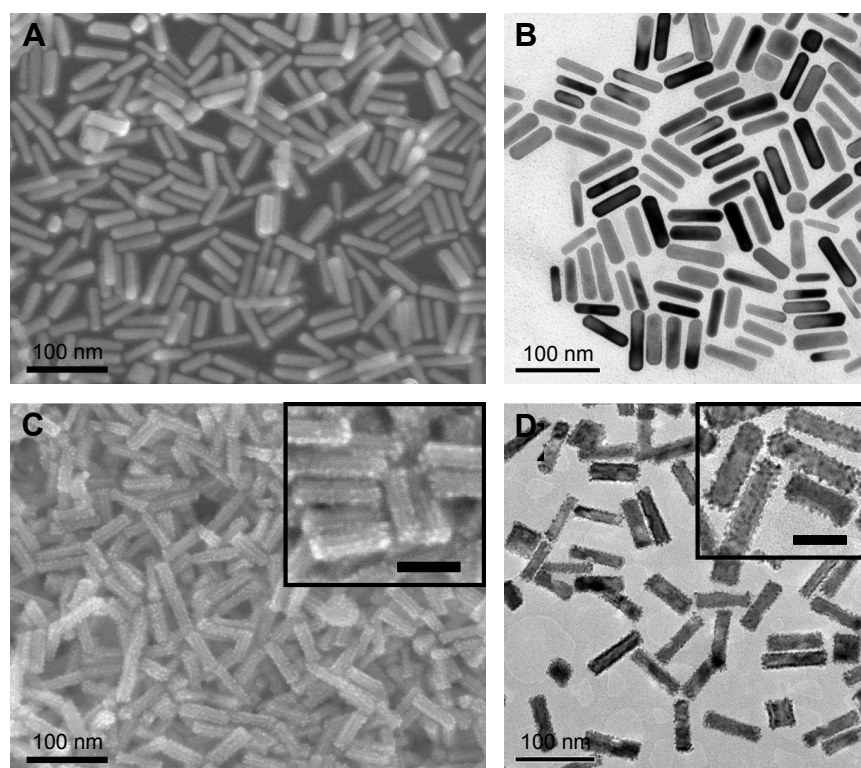
## Characterizations

Ultraviolet-visible-near-infrared extinction spectra were obtained from a Varian Cary 50. Transmission electron microscopy was performed on Tecnai G2 T20 S-TWIN (T20). The zeta potential data were obtained from Delsa Nano C (Beckman Coulter, Brea, CA, USA). ELISA was conducted on Infinite™ M200.

## Results and discussion

### Characterization of Au@Pt NRs and antigen-conjugated Au@Pt NRs

Herein, using Au NRs as seeds, we prepared Au@Pt nanostructures under mild conditions. Au NRs have an average length of 60 nm and width of 16 nm, and the aspect ratio (length vs width) of the Au NRs here is 3.8 (Figure 2). Pt nanodots with sizes of 3–4 nm were formed on the Au NRs with



**Figure 2** Typical SEM images of (A) Au NRs and (C) Au@Pt NRs. Typical TEM images of (B) Au NRs and (D) Au@Pt NRs.

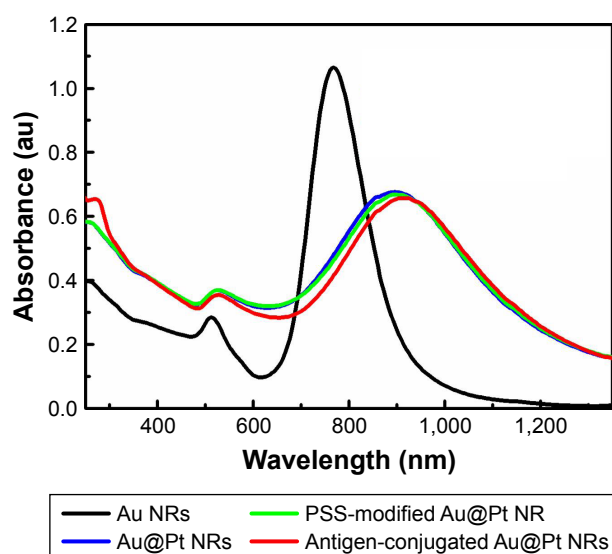
**Note:** The scale bar of insets in (C) and (D) is 50 nm.

**Abbreviations:** NR, nanorod; SEM, scanning electron microscopy; TEM, transmission electron microscopy.

a homogeneous distribution, and such a structure is desired for better catalytic activity. As shown in Figure 3, the Au NRs exhibit a strong longitudinal surface plasmonic resonance (SPR) band with a peak at 780 nm. Upon depositing Pt at a Pt/Au ratio of 0.2, Au@Pt NRs exhibit well-defined and

red-shifted longitudinal SPR bands in the visible and near-infrared region. For small Pt NPs only have SPR features in the ultraviolet region, which are difficult to employ, using Au NRs as support provides an easy way to characterize the SPR features of Pt. Also, both the amount and thickness of Pt determine its contribution to the final position and strength of the overall SPR features.

Zeta potential is used to predict the surface charge and stability of the nanoparticles. In our case, the as-prepared Au NRs are positively charged because of the presence of a bilayer of cetyltrimethylammonium bromide (CTAB; Table 1), but it should be changed to negative charge to immobilize specific antigens by electrostatic interactions. CTAB facilitates electrostatic adsorption of an anionic polyelectrolyte, PSS. PSS was coated to CTAB-capped Au@Pt NRs via electrostatic



**Figure 3** UV-vis-NIR spectra of Au NRs, Au@Pt NRs, PSS-modified Au@Pt NRs, and antigen-conjugated Au@Pt NRs.

**Abbreviations:** NIR, near infrared; NR, nanorod; PSS, poly(sodium-4-styrene sulfonate); UV, ultraviolet; vis, visible.

**Table 1** Effective diameter and zeta potentials of various nanoparticles

Materials	Effective diameter (nm)	Zeta potential (mV)
Au NRs	17.5±1.3	24.2±1.4
Au@Pt NRs	28.1±0.4	20.1±0.5
PSS-modified Au@Pt	34.2±0.8	-31.9±2.1
Antigen-conjugated Au@Pt NRs	82.0±2.1	-19.1±1.5

**Abbreviations:** NR, nanorod; PSS, poly(sodium-4-styrene sulfonate).



assembly, and it endows the NRs an enhanced stability in a wide pH range. Besides, the small antigen molecule could be directly conjugated on the surface of nanozyme through electrostatic force. As shown in Figure 3, PSS modification does not lead to obvious change in the SPR features of Au@Pt NRs. Figure 3 also illustrates that conjugating rubella antigen onto the surface of Au@Pt NRs would lead to a slight red shift with slight damping in the intensity of longitudinal SPR features, originating from the longitudinal SPR properties highly sensitive to changes in the dielectric constant of the local environment of Au@Pt NR surfaces. Table 1 shows that surface charges of NRs became less negative, also confirming the successful binding of the molecule to the NRs.

In this study, a light-scattering technique (dynamic light scattering) was used to determine the effective diameter of the nanoparticles (Table 1). It is worth mentioning that the size of the as-prepared Au NRs was 60 nm in length and 16 nm in width. However, light scattering typically showed an effective diameter of 17.5 nm. The light-scattering analysis assumes that particles are spherical; hence, due to the rod shape, the diameter obtained from dynamic light scattering measurements is not the actual size of the NRs. The effective diameter is used to evaluate the relative size upon various coatings. Forming a Pt nanodots shell on Au NRs would lead to an increase in the effective diameter. Upon further coating with a layer of PSS, the effective diameter of the Au@Pt NRs would reach 34.2 nm. After antigen conjugation process, the effective diameter of the conjugate becomes 82 nm. Hence, a smooth increase in the effective diameter of the Au NRs observed by light-scattering measurements indicates an increase in the hydrodynamic radius of the particles due to adsorption of different surface modifiers, and not the aggregation of the nanoparticles.

The stability of antigen-conjugated Au@Pt NRs over storage time was evaluated using zeta potential and also

their effective diameter (Figure S1). The antigen-conjugated Au@Pt NRs exhibited stable average diameter and zeta potential over 4 weeks, further revealing their good stability.

## Optimization of catalytic conditions of antigen-conjugated Au@Pt NRs

Previous studies have proved that the as-prepared Au@Pt indeed exhibited peroxidase-like activity. Therefore, the functionalized nanoprobe, antigen-conjugated Au NRs, would have similar catalytic activity. The color reaction of TMB in the presence of  $H_2O_2$  was used to demonstrate the catalytic activity of the antigen-conjugated Au NRs.

As shown in Figure 4A and B, after mixing the antigen-conjugated Au NRs with TMB and  $H_2O_2$ , the mixture turned from clear to blue (with a peak at 652 nm in the absorbance spectra). Compared with antigen-conjugated HRP, antigen-conjugated Au@Pt NRs showed a higher kinetics due to more catalytic sites provided by the larger surface area of the nanoparticles (Figure S2). The color development was stopped by the addition of  $H_2SO_4$ , changing the color to yellow with a peak at 450 nm in the absorbance spectra. To achieve an optimal analytical performance of the antigen-conjugated Au NRs, various parameters such as concentration of TMB/ $H_2O_2$ , concentration of antigen-conjugated Au@Pt NRs, reaction time, pH, and temperature are optimized (Figure 4). The concentration of TMB is a key factor because it determines the color level of the reaction system. Thus, the influence of the concentration of TMB from 0.1 to 1 mM was investigated. Figure 4C shows that the activity of antigen-conjugated Au@Pt NRs increased up to 0.75 mM TMB, and at higher concentrations, the absorbance of the system reached the plateau of maxima. Only slight changes in light intensity were observed with further increase in concentration of TMB. The  $H_2O_2$  concentration also has a very important

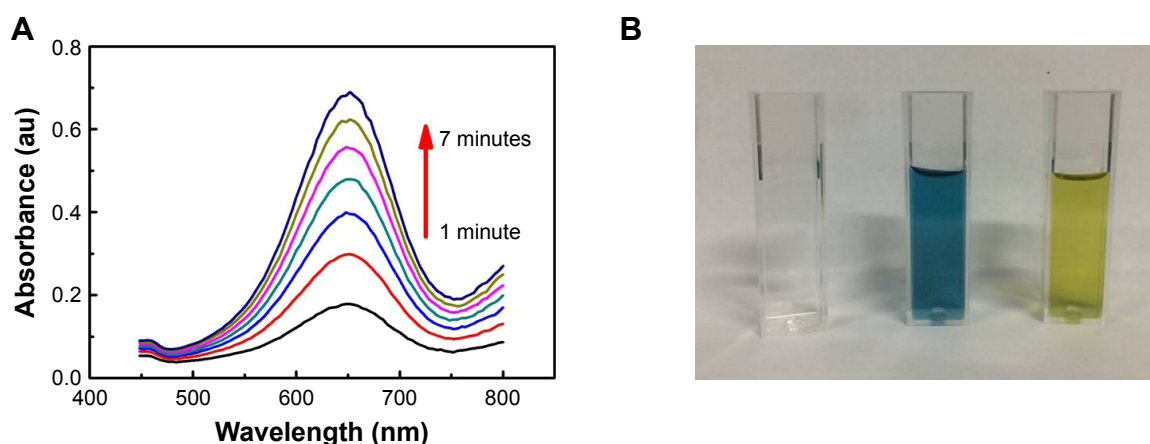
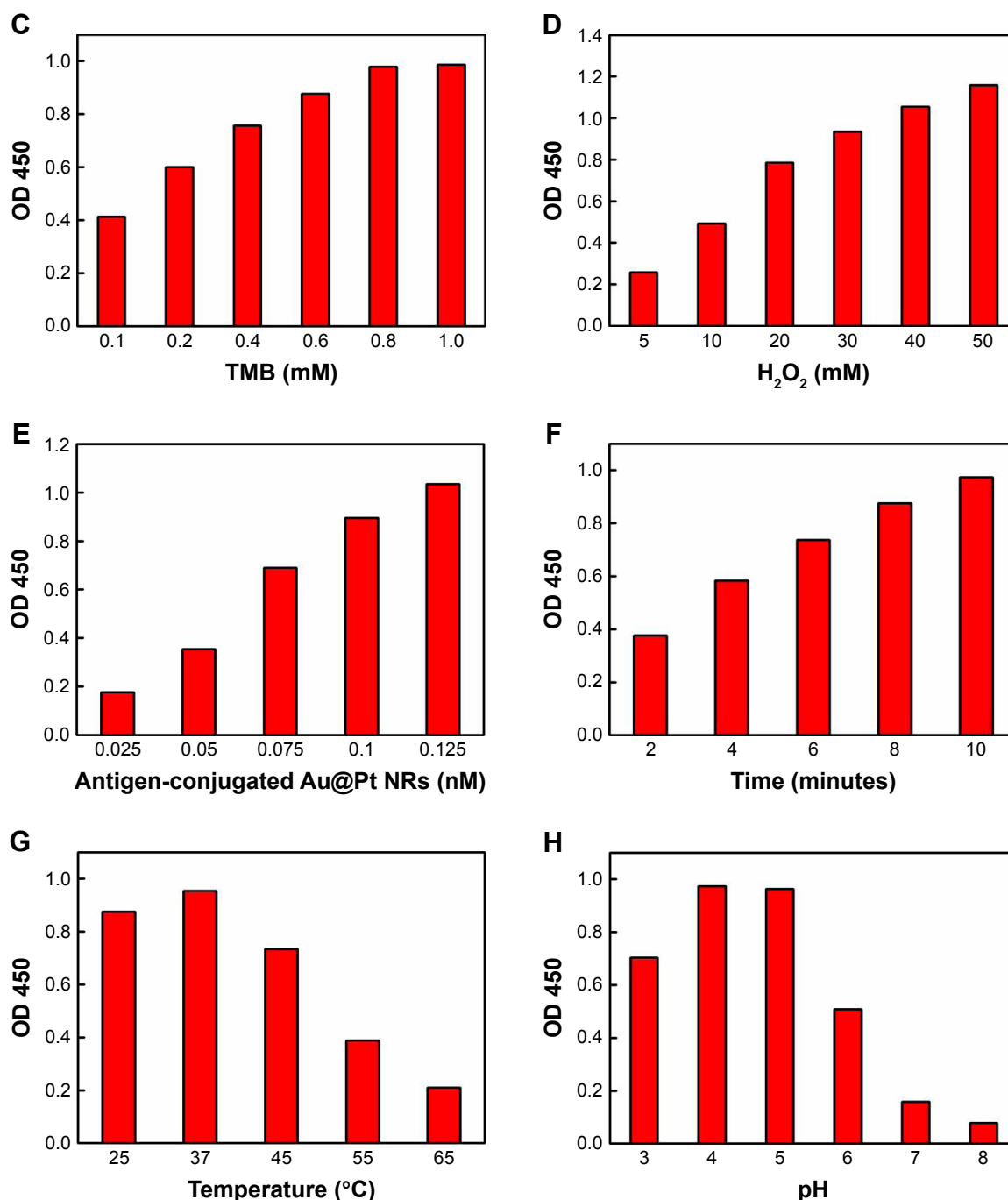


Figure 4 (Continued)



**Figure 4** (A) Time-dependent absorbance of TMB oxidation system in the presence of antigen-conjugated Au@Pt NRs. (B) Typical photographs of TMB–H<sub>2</sub>O<sub>2</sub> solution (left), TMB–H<sub>2</sub>O<sub>2</sub>–antigen-conjugated Au@Pt NRs (middle), and the reaction solutions after addition of H<sub>2</sub>SO<sub>4</sub>. Effects of (C) TMB concentration, (D) H<sub>2</sub>O<sub>2</sub> concentration, (E) antigen-conjugated Au@Pt NRs concentration, (F) reaction time, (G) temperature, and (H) pH on the catalytic activity of antigen-conjugated Au@Pt NRs.

**Notes:** Reaction conditions: (A, B) 0.125 nM antigen-conjugated Au@Pt NRs, 1 mM TMB, 50 mM H<sub>2</sub>O<sub>2</sub>, 10 minutes, 37°C, and 0.1 M, pH 5; (C) 0.125 nM Au@Pt NRs, 50 mM H<sub>2</sub>O<sub>2</sub>, 10 minutes, 37°C, and 0.1 M, pH 5; (D) 0.125 nM Au@Pt NRs, 1 mM TMB, 10 minutes, 37°C, and 0.1 M, pH 5; (E) 1 mM TMB, 50 mM H<sub>2</sub>O<sub>2</sub>, 10 minutes, 37°C, and 0.1 M, pH 5; (F) 0.125 nM Au@Pt NRs, 1 mM TMB, 50 mM H<sub>2</sub>O<sub>2</sub>, 37°C, and 0.1 M, pH 5; (G) 0.125 nM Au@Pt NRs, 1 mM TMB, 50 mM H<sub>2</sub>O<sub>2</sub>, 10 minutes, and 0.1 M, pH 5; (H) 0.125 nM Au@Pt NRs, 1 mM TMB, 50 mM H<sub>2</sub>O<sub>2</sub>, 10 minutes, and 37°C.

**Abbreviations:** NR, nanorod; TMB, 3,3',5,5'-tetramethylbenzidine.

influence on the performance of the antigen-conjugated Au@Pt NRs. So, experiments on optimizing H<sub>2</sub>O<sub>2</sub> concentration were performed and the results are shown in Figure 4D. It reveals that the absorbance at 450 nm shows an almost linear increase with H<sub>2</sub>O<sub>2</sub> concentration 5–50 mM. And then,

different concentrations of antigen-conjugated Au@Pt NRs were tested with optimum substrate concentration. Figure 4E shows an increase in absorbance with increasing antigen-conjugated Au@Pt NR concentration. The optimized reaction time of antigen-conjugated Au@Pt NRs

with substrate was also tested for varying time periods (2–10 minutes). Figure 4F shows an increase in absorbance with increasing time of reaction. Finally, the catalytic activity of antigen-conjugated Au@Pt NRs was determined over a range of temperature and pH. The temperature- and pH-dependent response curves are shown in Figure 4G and H. The phenomenon indicated that the catalytic activity of the antigen-conjugated Au@Pt NRs in the reaction system of TMB and  $H_2O_2$  was strongest at 37°C and pH 4–5, which is consistent with the natural enzyme.

To sum up, the optimum conditions selected in the ELISA were as follows: the concentration of TMB was 1 mM, the  $H_2O_2$  concentration was 50 mM, the antigen-conjugated Au@Pt NRs concentration was 0.125 nM, the reaction time was 10 minutes, the reaction temperature was 37°C, and the pH of the reaction buffer was 5.

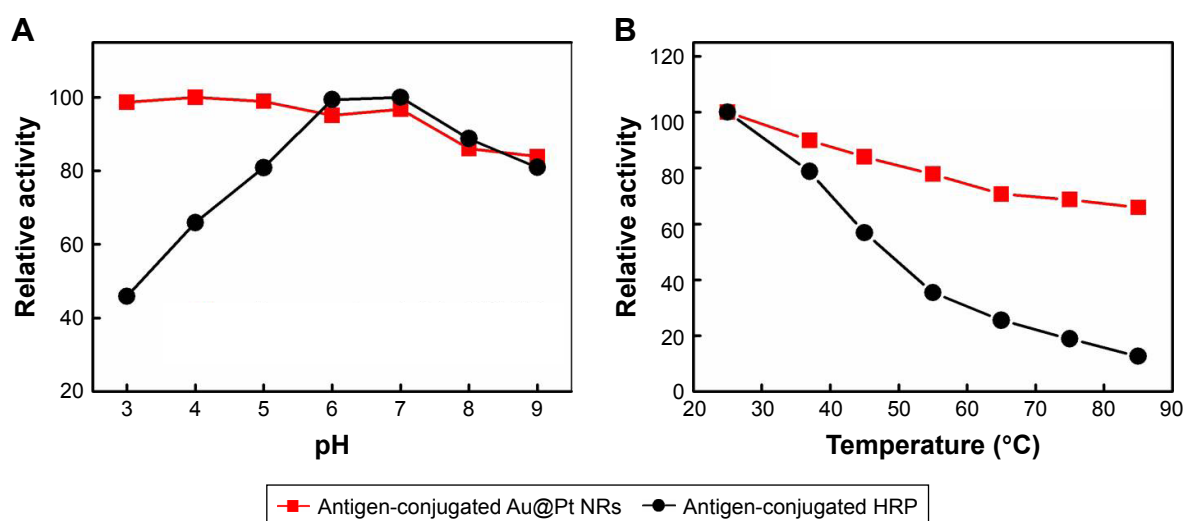
### Comparison of catalytic stability of antigen-conjugated Au@Pt NRs and antigen-conjugated HRP against temperature and pH

The stability of enzyme probe under different harsh chemical environments is critical to extend their applications. To examine the relative stabilities of antigen-conjugated Au@Pt NRs and antigen-conjugated HRP, both of them were pretreated at a range of pH values (3–9) and temperatures (25°C–85°C) for 3 hours, and then their peroxidase-like activity was carried out under the standard conditions of pH 5.0 and temperature 37°C (Figure 5). Antigen-conjugated Au@Pt NRs showed

similar activity over the pH range studied; however, the reactivity of antigen-conjugated HRP would be affected under acidic conditions (Figure 5A). Figure 5B shows that antigen-conjugated Au@Pt NRs retained the ability at all temperatures studied after 3 hours pretreatment, while antigen-conjugated HRP lost the majority of its activity above 55°C. The robustness of antigen-conjugated Au@Pt NRs makes them suitable for a broad range of applications in the biomedicine and environmental chemistry fields under various temperature and pH conditions.

### Application of biomedical assay

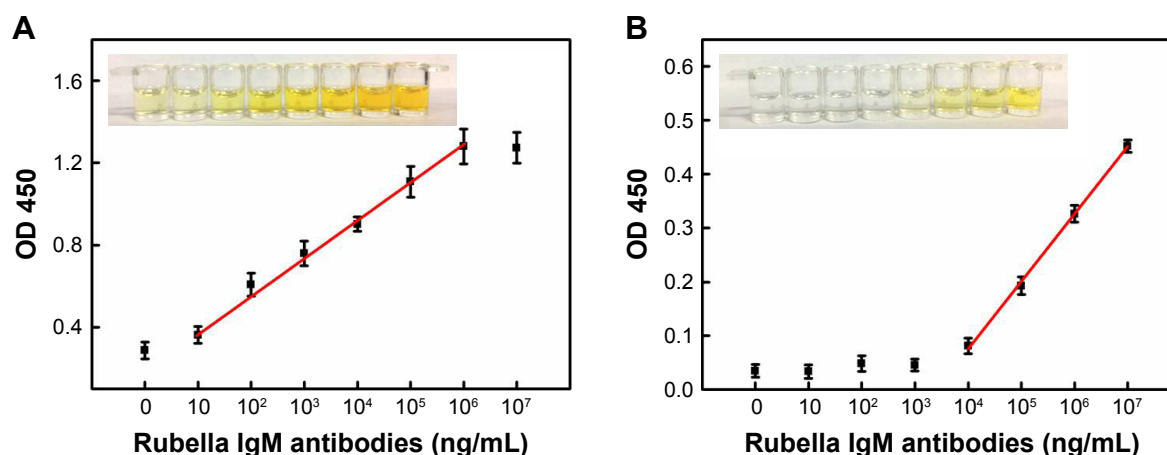
The ELISA kits for the detection of rubella-specific IgM are of the capture type. First, the assay employs anti-human IgM antibody attached to a solid phase. After washing and removal of anti-human IgM antibody, diluted test serum is added and incubated with the immobilized anti-human IgM antibody. The rubella-specific IgM antibodies present in the serum bind to anti-human IgM antibody. Then, the unbound antibodies are removed by washing and antigen-conjugated Au NR is added and further incubation is carried out. The quantity of the nanozyme probes that bind to each well is proportional to the concentration of the rubella-specific IgM antibodies present in the serum. The plates are then washed and substrate is added which results in color development. The enzymatic reaction is stopped after a short incubation period, and OD is measured by an ELISA reader instrument (Figure 1). A dose–response curve drawn for different concentrations of rubella IgM antibodies' standard solutions under optimum



**Figure 5** Comparison of the stability of antigen-conjugated Au@Pt NRs and antigen-conjugated HRP.

**Notes:** Antigen-conjugated Au@Pt NRs and antigen-conjugated HRP were (A) treated by media with pH ranging from 3 to 9 for 3 hours or (B) treated at a wide range of temperatures between 25°C and 85°C, and then their peroxidase activities were measured under standard conditions.

**Abbreviations:** HRP, horseradish peroxidase; NR, nanorod.

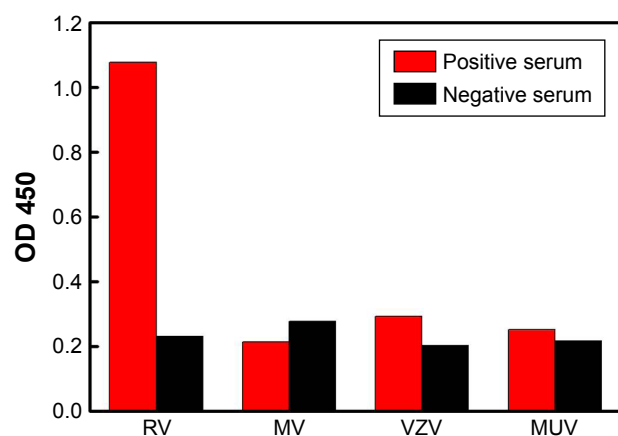


**Figure 6** Dose–response curves for different concentrations of rubella IgM antibodies standard solutions.

**Notes:** (A) Antigen-conjugated Au@Pt NRs-based ELISA and (B) antigen-conjugated HRP-based ELISA. All the error bars were calculated based on the SD of three measurements. The insets are the corresponding colors in the well.

**Abbreviations:** ELISA, enzyme-linked immunosorbent assay; HRP, horseradish peroxidase; NR, nanorod.

conditions is shown in Figure 6. The detection of rubella IgM antibodies in the linear range from 10 to  $10^6$  ng/mL, and the detection limit is as low as 10 ng/mL. Due to the amplified signals generated by the nanoprobe, the detection limit for rubella IgM antibodies increases by three orders of magnitude compared with conventional immunoassays. The increase in sensitivity is attributed to the use of Au@Pt NRs as a carrier and amplifier. First, Au@Pt NRs with high surface areas offer a possibility to load number of antigens to the surface of Au@Pt NRs. More antigens bound onto the Au@Pt NRs surface provide better detection specificity. It, indeed, shows that we can reduce the background signal by harboring more antigens on the particle surface (Figure S3). Second, the largest surface area also provides most catalytic sites, resulting in the strongest peroxidase-like activity compared to that of natural HRP enzyme (Figure S2).



**Figure 7** Specificity of antigen-conjugated Au@Pt NRs-based ELISA.

**Abbreviations:** ELISA, enzyme-linked immunosorbent assay; MUV, mumps virus; MV, measles virus; NR, nanorod; VZV, varicella-zoster virus.

Furthermore, it was found that the ELISA system retained nearly the same original optical response (OD 450) even after 4 weeks of storage at 4°C (Figure S4), revealing the good stability of the system. Compared to the detection range, the detection time, and the limitations in other rubella IgM immunoassays (Table S1), antigen-conjugated Au@Pt NR is an efficient candidate in the development of rubella IgM immunoassays.

To further investigate the specificity of the ELISA system, the immunoassay was used to detect other infectious viruses, such as measles virus, varicella-zoster virus, and mumps virus. No remarkable response was observed for the other samples in comparison to the result obtained in the RV-positive serum, indicating good specificity of the immunoassay (Figure 7). The proposed method was further validated by testing 30 clinical serum samples with reference to a commercialized ELISA method (20 RV negative and 10 RV positive). When the cutoff value was defined as three times the blank signal, the diagnosis of RV by the antigen-conjugated Au@Pt NRs-based ELISA agreed well with the commercial ELISA result, indicating the proposed immunosensor could be used for the preliminary detection of rubella-specific IgM in clinical diagnosis.

## Conclusion

In this paper, we prepared a novel nanoprobe to detect rubella IgM antibodies in sera. The rationale of detection is based on antigen-conjugated Au@Pt NRs. Compared with HRP-antigen conjugates, the obtained Au@Pt NR-antigen conjugates exhibited not only high peroxidase-like activity, but also robustness to harsh environments. Based on the



enhanced catalytic properties of this nanozyme probe, the detection limit for rubella-specific IgM antibodies in human serum was as low as 10 ng/mL, which is 1,000 times more sensitive than commercial ELISA. Hence, our experimental results demonstrate the potential feasibility of antibody-conjugated Au NRs for the highly sensitive virus diagnosis in future clinical applications under various conditions.

## Acknowledgments

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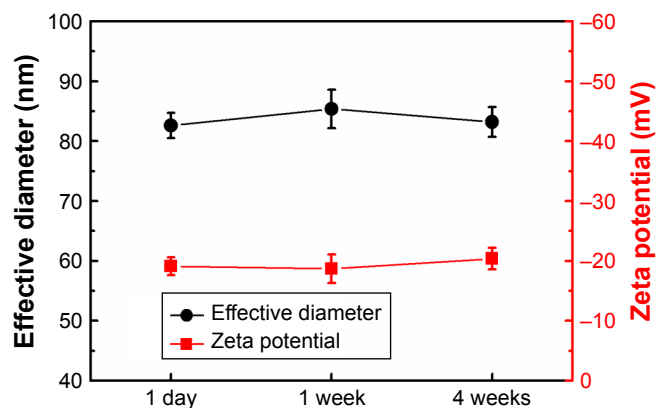
## Disclosure

The authors report no conflicts of interest in this work.

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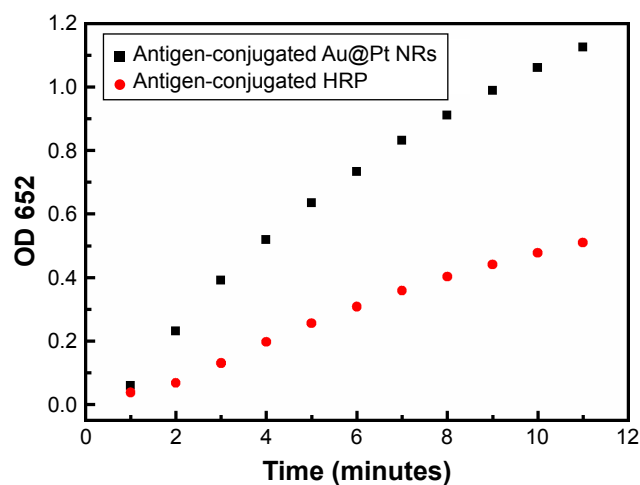
## Supplementary materials



**Figure S1** Effects of storage time on the stability of antigen-conjugated Au NRs in 0.1 M PBS solution (pH=7.4) at room temperature.

**Note:** All the error bars were calculated based on the SD of three measurements.

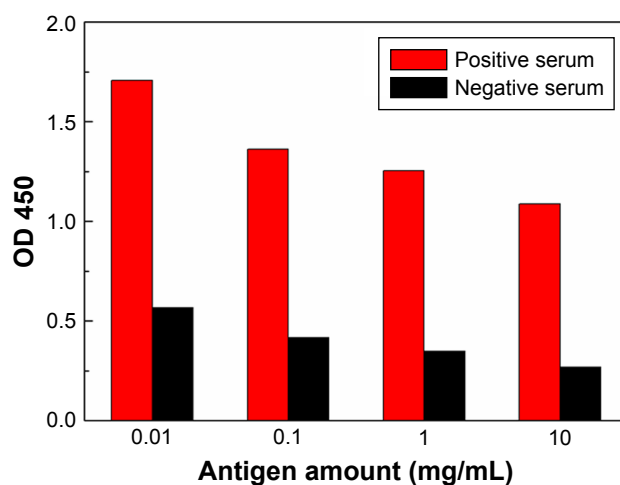
**Abbreviation:** NR, nanorod.



**Figure S2** The absorbance evolution at 650 nm over time for TMB oxidation catalyzed by antigen-conjugated Au@Pt NRs and antigen-conjugated HRP.

**Note:** Reaction conditions: 1 mM TMB, 50 mM H<sub>2</sub>O<sub>2</sub>, 10 minutes, 37°C, and 0.1 M, pH 5.

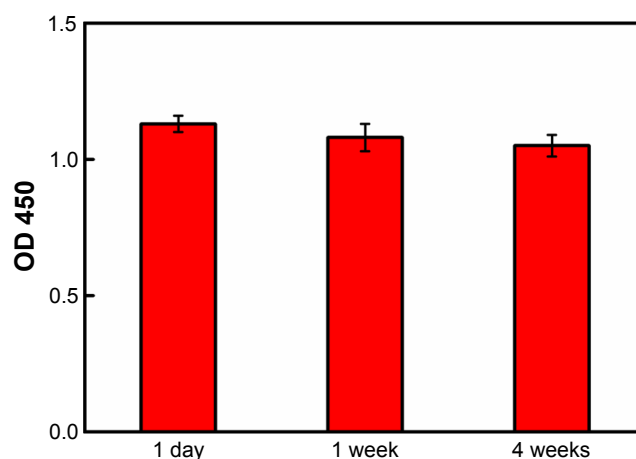
**Abbreviations:** HRP, horseradish peroxidase; NR, nanorod; TMB, 3,3',5,5'-tetramethylbenzidine.



**Figure S3** Effects of the amount of antigens on Au@Pt NR surface for the performance of the immunoassay.

**Note:** The resulting antigen-conjugated Au@Pt NRs were evaluated by detecting positive control and negative control.

**Abbreviation:** NR, nanorod.



**Figure S4** Effects of storage time on the reactivity of antigen-conjugated Au@Pt NRs in 0.1 M PBS solution (pH=7.4) at room temperature.

**Notes:** All the error bars were calculated based on the SD of three measurements. Reaction conditions: 0.125 nM antigen-conjugated Au@Pt NRs, 1 mM TMB, 50 mM H<sub>2</sub>O<sub>2</sub>, 10 minutes, 37°C, and 0.1 M, pH 5.

**Abbreviations:** NR, nanorod; TMB, 3,3',5,5'-tetramethylbenzidine.

**Table S1** Comparison of rubella IgM immunoassay

System	Detection method	Detection range (ng/mL)	Detection time (hours)	Limitations
Sigma rubella IgM (capture) (Sigma-Aldrich Co., St Louis, MO, USA)	IgM capture format assay	10 <sup>3</sup> –10 <sup>7</sup>	~2.5	Relative low stability
Sigma rubella IgM (Sigma-Aldrich Co.)	Indirect format assay	10 <sup>3</sup> –10 <sup>7</sup>	~1.5	Gives false-positive results
This work	IgM capture format assay	10–10 <sup>7</sup>	~2	No commercial assay available

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