An aptasensor for ampicillin detection in milk by fluorescence resonance energy transfer between upconversion nanoparticles and Au nanoparticles

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ABSTRACT

This paper reports a portable fluorescence resonance energy transfer (FRET) aptasensor for ampicillin (Amp) detection using upconversion particles (UCNPs) as energy donors and Au nanoparticles (AuNPs) as energy acceptors. The optimal parameters of the detection system were investigated. Under the optimal conditions, it had a good linear relationship between the fluorescence intensities and Amp concentrations, a high coefficient of determination (R^2) of 0.9939, a wide detection range of 10–100 ng/mL, and a low limit of detection (LOD) of 3.9 ng/mL; meanwhile, the aptasensor had high selectivity for Amp against the interference of other antibiotics, and had good recovery and repeatability. Also, its detection performance had been successfully validated by milk samples. Therefore, the developed aptasensor based on FRET between UCNPs and AuNPs has a good prospect for Amp on-site detection in milk with a portable upconversion detection instrument.

1. Introduction

Ampicillin (Amp) is a kind of β-lactam antibiotic, which inhibits bacterial transpeptidase to hinder the synthesis of the cell wall (Luo, Ang, & Thompson Jr, 1997). As a human and a veterinary drug, it has a strong bactericidal activity and a wide range of applications. However, Amp cannot be completely absorbed and metabolized, and a part of Amp remains in animal’s body. It then can enter the human body through the food chain, which is harmful to human health. Partial Amp is excreted with feces and pollutes water sources and soil, through which it will be accumulated in food plants and will partially enter the human body (Eilmola & Chaudhurt, 2011).

Amp accumulation in the body will induce bacterial resistance and intestinal flora imbalance. In addition, Amp, as a hapten, can bind tissue protein in the body to become a complete antigen, which will cause hypersensitivity reactions for a few special people (Abouzied, Sarzynski, Walsh, Wood, & Mozola, 2009). The European Union mandates the maximum residue limits (MRL) of Amp in milk and animal meat as 4 μg/L and 50 μg/L, respectively, and the Food and Drug Administration (FDA) stipulates Amp MRL in milk is 10 μg/L (Song, Jeong, Jeon, Cho, & Ban, 2012). Presently, several methods have been used for antibiotics detection in food, such as high-performance liquid chromatography (HPLC) (Luo, Ang, & Thompson Jr, 1997), enzyme-linked immunosorbent assay (ELISA) (FitzGerald, O’Loan, McConnell, Benchikha, & Kane, 2007), and electrochemistry (Li, Zhu, Li, Liu, & Kang, 2020), etc. However, expensive instruments and professional users are required to implement the analysis work, making them unable for a large number of detection tasks. Researchers have made great efforts to develop biosensors for portable and simple Amp detection. It has been documented that the gold-nanoparticles-based dual-fluorescence colorimetric aptamer sensor was developed for the trace detection of Amp in milk and water (Song, Jeong, Jeon, Cho, & Ban, 2012). A portable paper-based fluorescence immunoassay was reported for the trace detection of norflaxacin in milk (Zong, Jiao, Guo, Zhu, Gao, Han, et al., 2019). A colorimetric-based point-of-care testing method, in which the fluorescence image of the sample was obtained through a smartphone camera, was developed for streptomycin detection in milk, it had the advantages of good selectivity, simple operation, and on-site detection (Liu, Yu, Cao, Guo, Zhu, Dai, et al., 2018). A fluorescent-internal-filter-based test strip was designed for the rapid on-site visual detection of tetracycline, it has the advantages of simple operation and easy portability (Gan, Hu, Xu, Zhang, Zou, Shi, et al., 2021). Although the above portable detection methods did not require expensive instruments, they all used traditional fluorescent dyes, which were often excited by ultraviolet light and can induce sample autofluorescence, therefore, false positives were easy to occur. UCNPs have anti-Stokes luminescence properties, and can absorb long-wavelength low-energy light and emit short-wavelength high-
energy light (Guo, Xie, Huang, & Huang, 2016; Xu, Chen, & Song, 2017). Because the long-wavelength light is weakly absorbed by biological samples, it can avoid the interference of sample autofluorescence. Accordingly, UCNPs are better than traditional down-conversion fluorescent dyes (Wu, Duan, Ma, Xia, Wang, Wang, et al., 2012) for the complex samples matrices, such as foods and biomedical samples (Gao, Zheng, Liu, Han, Li, Luo, et al., 2021; Li, Wang, Huang, & Chen, 2019; Yang, Li, Wu, Zhang, Feng, Yu, et al., 2021). Due to its high Stokes shift, narrow emission band, high photostability, and low autofluorescence (Long, Li, Zhang, & Yao, 2015), UCNPs, in this study, were used as the fluorescent probe for Amp detection. In addition, a portable upconversion fluorescence instrument was applied, it had the advantages of small size, low weight, and simple operation (Xu, Chen, & Song, 2017), compared to the traditional upconversion measurement instruments based on photomultiplier tubes and monochromators.

FRET is a mechanism that describes the transfer of energy between two light-sensitive molecules (chromophores) (Cheng, 2006). The donor chromophore, which is initially in an electronically excited state, can transfer energy to the acceptor chromophore through nonradiative dipole-dipole coupling (Helms, 2008). The generation of FRET needs to meet two conditions: one is that the spatial distance between the donor and the acceptor is less than 10 nm, and the other is the overlap between the emission spectrum of the energy donor and the absorption spectrum of the acceptor (Wang, Hou, Mi, Wang, Xu, Teng, et al., 2009). Herein, we established a label-free aptamer sensor system based on FRET (UCNPs and AuNPs) for the simple, immediate, and sensitive detection of Amp. The detection principle is shown in Scheme 1. It consisted of UCNPs, AuNPs, Amp aptamer (Apt), Amp, and NaCl, UCNPs were used as energy donors, while AuNPs were as acceptors. When they were mixed, positively charged UCNPs and negatively charged AuNPs approached each other through electrostatic adsorption, which led to the fluorescence quenching of UCNPs by FRET. When NaCl was introduced, NaCl solution shielded the charges on the surface of AuNPs, consequently, AuNPs stayed away from UCNPs and formed aggregation, and then the fluorescence of UCNPs was restored. However, when Apt was introduced into the system, it was strongly adsorbed on the surface of AuNPs through the coordination between nitrogen atoms (in Apt) and AuNPs, therefore, the phosphoric skeleton of Apt covered the surface of AuNPs, and the electrostatic repulsion prevented from their aggregation, which enhanced the stability of AuNPs in a salt solution and led to the fluorescence quenching of UCNPs. In the measurement, when Amp was added to the system, Apt would preferentially combine with Amp. With the increase of Amp concentration, AuNPs absorbing Apt decreased, resulting in AuNPs aggregation, which led to the decrease of the fluorescence quenching effect of AuNPs on UCNPs, therefore, the fluorescence intensity of UCNPs was restored. It had a linear relationship between Amp concentration and UCNPs fluorescence intensity, therefore, Amp concentration could be measured.

2. Materials and methods

2.1. Materials

Yttrium (III) chloride hexahydrate (YCl$_3$·6H$_2$O, 99.99 %), Ytterbium (III) chloride hexahydrate (YbCl$_3$·6H$_2$O, 99.99 %), and Erbium (III) chloride hexahydrate (ErCl$_3$·6H$_2$O, 99.99 %), Oleic acid (90 %), and 1-Octacene sodium (90 %) were purchased from Shanghai Aladdin Biochemical Technology Co., ltd. (Shanghai, China). Chloroauric acid tetrahydrate (HAuCl$_4$, 99.9 %), sodium citrate tribasic dehydrate, and tetraethoxysilane (TEOS) were supplied by Sinopharm Chemical Reagent Co., ltd. (Shanghai, China). Igepal CO-520 was purchased from Rhawn Chemical Technology Co., ltd (Shanghai, China). Apt 5′-GCGGG GGTG TATAG CGG-3′ was synthesized by Sangon Biotech (Shanghai) Co., ltd. (Shanghai, China). All unspecified reagents were of analytical reagent grade and millipore milli-Q ultrapure water was used in this study.

2.2. Apparatus

Transmission electron microscope (TEM) images were taken using a HT-7800 electron microscope (Hitachi, ltd, Tokyo, Japan, Japan). The absorption spectra of AuNPs under different conditions were recorded by a USB 4000 spectrometer (Ocean Optics, Orlando, FL., USA). The fluorescence emission spectra of UCNPs were performed using a XS11639 spectrometer with a 980 nm excitation laser and FPB-980–1.5-FS-1B probe (Shanghai Ruhai photoelectric technology Co., ltd., Shanghai, China).

![Scheme 1. The principle of the aptasensor for Amp detection.](image-url)
2.3. Synthesis and surface modification of UCNPs

UCNPs were synthesized according to the references with a little modification (Liu, Ouyang, Li, Zhang, & Chen, 2017). 1.0 mmol rare earth chloride (Y: Yb: Er = 78:20:2) was added to a 250 mL three-neck round-bottom flask containing 6 mL oleic acid (OA) and 15 mL 1-octadecene (ODE), mixed under magnetic stirring. Then, high-purity nitrogen gas was passed through for 10 min to exhaust the air in the flask. Under the protection of high-purity nitrogen gas and magnetic stirring, heat up to 160°C and hold for 30 min to form a uniform solution. Subsequently, the temperature was dropped to 50°C, and a 10 mL methanol solution containing 2.5 mM NaOH and 4 mM NH₄F was slowly added to the solution, and maintained for 30 min to remove methanol from the solution. The morphology of UCNPs TEM is shown in Figure s1 (supplementary material). They were hexagonal with uniform particle size distribution and an average particle size of 55 nm. The size of UCNPs@SiO₂-NH₂ was increased to 70 nm by coating SiO₂ on UCNPs, therefore, the thickness of the SiO₂ layer was 7.5 nm.

AuNPs were synthesized by the sodium citrate reduction method, and the size was affected by the amount of sodium citrate in the synthesis process (Frens, 1973). AuNPs were characterized by UV–vis spectroscopy and TEM. As shown in Fig. s2A, its absorption peak was at 520 nm, which was consistent with the literature (Chen, Sheng, Wang, Ouyang, Wang, Ali, et al., 2020). The TEM photo was shown in Fig. s2B, in which AuNPs were uniformly dispersed and the particle size was about 13 nm, which was consistent with the literature reports (Sun, Wang, Sun, Li, Song, Zhang, et al., 2020).

2.4. Synthesis of AuNPs

AuNPs with an average diameter of 13 nm were prepared (supplementary material) according to the referenced method (Shi, Zhao, Liu, Fan, & Cao, 2013). According to Beer’s law, the concentration of AuNPs was about 9.6 nM (Haiss, Thanh, Aveyard, & Fernig, 2007).

2.5. Analysis method construction for Amp detection

In this work, the method of Amp detection was designed as follows: 56 μL Amp with different concentrations was added to 62 μL 0.4 μM Apt solution and was mixed with vortex oscillation, and incubated for 10 min. Then, 266 μL 4.8 mM AuNPs solution was successively added and then reacted for another 8 min. Subsequently, 266 μL 90 mM NaCl was added and thoroughly shaken for 13 min. Finally, 100 μL 0.1 mg/mL UCNPs@SiO₂-NH₂ was quickly added. After vortex mixing, recorded the fluorescence spectra in the wavelength range 400–700 nm under the excitation of a 980 nm laser. In order to obtain a high performance system for Amp detection, several important parameters, including the concentrations of NaCl and Apt, and the reaction time of NaCl-AuNPs, Apt-AuNPs, and Apt-Amp, were investigated.

2.6. Real sample detection

To validate the feasibility of this method, milk samples purchased from local supermarkets were tested. The samples were pretreated according to the national standard method (NY/T 829–2004, China). 4 mL milk was added to 20.0 mL 75 % acetonitrile, oscillated by vortex for 5 min, and then centrifuged at 8000 r/min for 10 min to remove the proteins in milk, and the supernatant was passed through a 0.22 μM membrane to remove the lipids. Subsequently, the detection was carried out with the developed method in section 2.5 and the reference method (HPLC), respectively. Three spiked samples were tested, and three parallel measurements were performed for each sample.

3. Results and discussion

3.1. Characterization of UCNPs and AuNPs

The morphology of UCNPs TEM is shown in Figure s1 (supplementary material). They were hexagonal with uniform particle size distribution and an average particle size of 55 nm. The size of UCNPs@SiO₂-NH₂ was increased to 70 nm by coating SiO₂ on UCNPs, therefore, the thickness of the SiO₂ layer was 7.5 nm.

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3.2. FRET between UCNPs and AuNPs

In this work, the UCNPs-AuNPs FRET system was constructed based on AuNPs induced UCNPs fluorescence quenching. FRET process was affected by three factors, the distance between the donor and the recipient, the spectral overlap of the donor and the receptor, and the relative orientation of the transition dipole (Jin, Wang, Lin, Jin, Zhang, Cui, et al., 2017). AuNPs generated by citric acid synthesis had negative charges and were uniformly stabilized in an aqueous solution due to the stronger electrostatic repulsion. Surface amination-modified AuNPs carried positive charges, so negatively charged AuNPs and positively charged UCNPs were close to each other through electrostatic interaction, resulting in FRET. As shown in Fig. 1A, the TEM photo of UCNPs@SiO₂-NH₂/AuNPs successfully verified the interaction. As shown in Fig. 1B, the absorption spectrum of AuNPs significantly overlapped with the emission spectrum of UCNPs, demonstrating that effective FRET could occur between UCNPs and AuNPs.

3.3. Principle of the FRET aptasensor for Amp

To verify the feasibility of this aptasensor for Amp detection, the effects of several related factors were investigated. UCNPs were incubated with Amp, NaCl, Apt, and Amp-Apt under the same conditions, respectively. As shown in Figure s3, their fluorescence intensities were as same as that of water-soluble UCNPs, so these factors can not affect the fluorescence of UCNPs.

The interactions of the components in the proposed Amp detection system are shown in Figure s4. UCNPs had a strong fluorescence signal at 540 nm. When mixed with AuNPs, the fluorescence intensity of UCNPs was significantly reduced, suggesting that FRET occurred between UCNPs and AuNPs. When AuNPs, NaCl, and UCNPs were mixed together, the fluorescence of UCNPs partially recovered, it was because the NaCl (salt) solution could shield the electrostatic repulsion between AuNPs, resulting in AuNPs aggregation, which greatly reduced the efficiency of FRET between UCNPs and AuNPs. When Apt, AuNPs, NaCl, and UCNPs were mixed together, UCNPs fluorescence was quenched again, the reason was that the Apt could be strongly adsorbed on the surface of AuNPs through the coordination between the nitrogen atoms (Apt) and AuNPs, enhancing the stability of AuNPs against the salt-induced aggregation (Chang, Chen, Wu, Yang, Lin, Mondal, Ramalal, Lavu, & Kingston, 2018). When Amp, Apt, AuNPs, NaCl, and UCNPs were mixed, the fluorescence of UCNPs was restored again, its mechanism was that Apt and Amp were banded and Apt could not stabilize AuNPs, causing AuNPs to aggregate again. Therefore, the system of Apt, AuNPs, NaCl and UCNPs could be used for the detection...
Fig. 1. TEM image of UCNPs/AuNPs (A) and their corresponding emission fluorescence/absorption spectra (B).

Fig. 2. Optimization of sodium chloride salt solution concentration (A), Apt concentration (B), and Optimization of reaction time of NaCl-AuNPs (C), Apt-AuNPs (D), and Apt-Amp(E).
of Amp.

3.4. Optimization of the detection conditions

To obtain a highly sensitive Amp sensor, several parameters, including NaCl and Apt concentrations, reaction time for NaCl-AuNPs, Apt-AuNPs, and Apt-Amp, were investigated, respectively.

The performance of the aptasensor largely depends on the number and distribution of donor (UCNPs) and receptor (AuNPs). NaCl concentration directly affected AuNPs aggregation, so it was necessary to optimize NaCl concentration. AuNPs were added to equal volume of NaCl solution (60, 70, 80, 90, 100 mM). After 10 min mixture, their absorption spectra were collected, and the results are shown in Fig. 2A. With the concentration increase of NaCl, the absorption peak at 520 nm of AuNPs decreased gradually, and its value tended to be stable at 90 mM NaCl, so the optimal NaCl concentration was 90 mM.

The Apt can be strongly adsorbed on the surface of AuNPs, improving the stability of AuNPs against the salt-induced aggregation, therefore, the concentration of Apt was also an important factor affecting their binding. Different concentrations (0.1, 0.2, 0.3, 0.4, 0.5 μM) Apt were mixed with 4.8 nM AuNPs for 5 min, and 90 mM NaCl solution was added and mixed. 10 min later, the spectra of the solution were collected. The results are shown in Fig. 2B, with the increase of Apt concentration, the absorption value of AuNPs at 520 nm increased, which demonstrated that the protection of AuNPs aggregation in the salt solution was enhanced. When Apt concentration reached 0.4 μM, the absorption value tended to be stable, therefore, 0.4 μM Apt was selected for the aptasensor system.

In addition, the three reaction time, NaCl-AuNPs, Apt-AuNPs, and Apt-Amp, were optimized. Firstly, the time optimization for NaCl-AuNPs was investigated. As shown in Fig. 2C, with the reaction time of AuNPs and NaCl extended, the absorption peak of AuNPs at 520 nm gradually decreased, indicating that AuNPs aggregated. 13 min later, the absorption value tended to be stable, therefore, 13 min was the optimal reaction time. Secondly, the influence of reaction time on the binding of Apt and AuNPs was discussed. As shown in Fig. 2D, with the time increased, the absorption peak of AuNPs at 520 nm increased, indicating that the Apt’s protection capability for AuNPs aggregation increased. When the reaction time was 8 min, Apt could be adsorbed on AuNPs well and the absorption value tended to be stable, therefore, 8 min was applied for the detection system. Finally, the specific binding time of Apt and Amp was optimized. As shown in Fig. 2E, with the increase of reaction time, the absorption peak of AuNPs at 520 nm decreased, indicating that AuNPs aggregated. When the reaction time was 10 min, the AuNPs absorption value tended to be stable, suggesting that Apt and Amp completed the specific recognition and binding, thus 10 min was applied in this detection system.

3.5. Amp detection

Fig. 3A shows the fluorescence spectra of the UCNPs/AuNPs/Apt/NaCl mixture under the presence of different concentrations of Amp ranging from 0 to 150 ng/mL. As shown in Fig. 3A, with the concentration of Amp increased, the fluorescence intensity at 540 nm of UCNPs gradually increased.

The plots of Amp concentration vs fluorescence intensity (540 nm) are shown in Fig. 3B. For 10–150 ng/mL Amp, the linear fitting equation was as follows:

\[ y = 47.392x + 259.142 \]

where \( y \) and \( x \) represented fluorescence recovery and Amp concentration, and coefficient of determination (R\(^2\)) was 0.9939, which demonstrated that they have a good linear relationship. The limit of detection (LOD) was calculated to be 3.9 ng/mL (S/N = 3), which was lower than the safe level for milk (10 μg/L). Thus, this aptasensor can quantitatively detect Amp at a low concentration.

A comparison of the different methods for Amp assay is summarized in Tab. 1. Compared with the HPLC and fluorescence colorimetry, the FRET aptasensor constructed in this work has a wide detection range. Although the sensitivity was lower than those of the HPLC and fluorescence colorimetry, its sensitivity was close to that of the ELISA method and better than that of the electrochemical method. Moreover, compared with others, the present method had the advantages of simple operation and portable, and it could meet the requirements of animal husbandry for the rapid detection of milk.

3.6. Specificity of the aptasensor for Amp

Selectivity is a very important parameter for the performance evaluation of a sensor. In this work, six antibiotics, including erythromycin, kanamycin, penicillin G, tetracycline, oxytetracycline, and azithromycin, were subjected to the same procedure as Amp detection. As shown in Fig. 4, only Amp caused significant fluorescence changes, while the changes caused by other antibiotics were small. It demonstrated that the designed fluorescent aptasensor has a high specificity for Amp detection.

3.7. Reproducibility and stability of the aptasensor for Amp

The stability and reproducibility of a sensor were of great significance for future practical applications. Three parallel measurements were performed on the samples containing 25 ng/mL Amp, the result showed that the relative standard deviation (RSD) was 3.49 %, which demonstrated that the aptasensor had good reproducibility. After the synthesized UCNPs were stored at 4 °C for one month, three parallel tests were performed on milk containing 25 ng/mL Amp, the test result showed that the absolute error of Amp was 2.28 ng/mL, and the RSD was

![Fig. 3. Fluorescence emission spectra of UCNPs for detecting different concentrations of Amp, the original spectra (A), the regression fitting (B).](image-url)
for the milk sample, the Amp concentrations determined by the HPLC was selected as the reference method. The results are shown in Table 1, tested. According to the national standard method, the HPLC method had good stability for Amp detection. For this method, the average absorbance follows:

\[ y = 1.031x - 0.7917 \tag{2} \]

where y and x represented results from the HPLC and that from this method. R² was 0.9944, it demonstrated that this method had a good relationship with the HPLC method. For this method, the average absolute error was 2.04 ng mL⁻¹, which was small; the average relative error was 4.2 %, which was lower than 5 %. Therefore, the developed aptasensor had a good detection performance, and can be used for the detection of Amp in milk sample.

Based on the wide detection range and simple operation, the proposed method had the advantages of low detection cost, rapidity, and easy portability. Regarding the limitation of this method, the LOD was relatively high compared to other methods (Tab. s1), however, it was suitable for the detection of Amp in milk.

4. Conclusion

This study successfully developed a novel aptasensor for Amp detection based on FRET between UCNPs and AuNPs. It can be used for the accurate determination of Amp in milk. Also, it had a potential capability for Amp detection in other samples. In addition, the proposed detection system can be easily applied for the detection of other targets by replacing suitable aptamers, it provided a new way for the

applications in agriculture, environment, and biomedical areas.

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CRediT authorship contribution statement

Chong Chen: Methodology, Validation, Formal analysis, Data curation, Writing – original draft. Hong Lei: Investigation, Methodology. Nan Liu: Investigation, Methodology. Hui Yan: Conceptualization, Resources, Data curation, Writing – review & editing, Supervision, Funding acquisition.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.fochx.2022.100439.

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Fig. 4. Specificity assessment of upconversion fluorescence assays.

Table 1

| Samples | Spiked (ng mL⁻¹) | HPLC method (ng mL⁻¹) | This method (ng mL⁻¹) | Recovery (%) | RSD (%) | p  
|--------|------------------|-----------------------|----------------------|-------------|---------|-----
| 1      | 0                | N. D. ²               | -0.95 ± 0.88        | 100.5 ± 0.50 | 100.5 ± 0.49 | 0.15 |
| 2      | 10               | 9.26 ± 0.38           | 10.05 ± 0.50        | 101.2 ± 0.12 | 101.2 ± 0.50 | 0.42 |
| 3      | 45               | 47.44 ± 1.42          | 45.53 ± 2.64        | 93.3 ± 1.89  | 93.3 ± 1.89  | 0.27 |
| 4      | 100              | 90.02 ± 3.19          | 93.33 ± 1.77        | 100 ± 0.38   | 100 ± 0.38   |   – |

² N. D.: not detect.

³ RSD: (standard deviation)/mean × 100 %, n = 3.

⁴ p > 0.05, indicate: no significant difference.
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