



Label-free and sensitive detection of microalgae protein using GNRs-based resonance light scattering system

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A novel and label-free gold nanorods (GNRs)-based resonance light scattering system assay has been developed on the basis of that the interactions between microalgae proteins and GNRs could emit strong fluorescence signal. We have prepared GNRs which were well dispersed in the solution and the microalgae protein was absorbed onto the surface of GNRs. The results demonstrated that the intensity of fluorescence has correlation with the protein concentration. The optimum pH was 5.5 and the optimum concentration of inorganic salt ions Na^+ was 0.5 mol L⁻¹; the stable time of the reaction system was 2 min. Because of the protein molecules are firmly combined with the surface of the gold particles, a protein layer is formed to prevent the aggregation of gold nanoparticles. Gold nanoparticles have a strong adsorption to proteins and other biological macromolecules and will not change their biological activity, hence it provides a number of advantages. This method offers the advantages of higher sensitivity and selectivity in microalgae protein detection and providing great potential for biology diagnosis.

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1. Introduction

Microalgae proteins, as a kind of biological macromolecule, are the material basis of microalgae which constitute the basic organic matter of microalgal cells and the main life activities of microalgae.¹ Because of the significance of the microalgae proteins, it is high priority to study the microalgae proteins in clinical medicine and biological chemistry aspects.^{2,3} A number of conventional detection methods were developed and utilized for microalgae determination, such as Kjeldahl method, Bradford method, spectrometer method, chemical luminescence method and near infrared reflectance spectroscopy. However, these methods suffer from limitations of low sensitivity, poor selectivity or weak stability in different settings.

Gold nanoparticles are tiny particles with 1–100 nm in diameter. As a new type of biological molecule probe, gold nanoparticles are simple and stable. Thus, it has been extensively studied and applied^{4–6} in different shapes and geometries such as spherical particles, nanorods, nanoshells, nanostars

and nanocages.^{6–11} The optical properties are strongly dependent on their shape and size.^{12–14} As one of the earliest studies on metal nano materials and their stability, strong and stable surface plasma resonance absorption peak appearing in the UV-visible region was utilized.¹⁵ The gold nanoparticles were used in electron microscopy for the first time in 1971.¹⁶ GNRs exhibit in two SPR bands with the transverse band around 520–600 nm and the longitudinal band in the near IR region, with the exact wavelength tunable by controlling the aspect ratio of the nanorods.¹⁷ Because of their near IR SPR band, GNRs are considered as more promising than solid spherical nanoparticles for *in vivo* bio-medical applications.

Currently more studies focused on the biological probe of the gold nanoparticles. Because of its quantum effect, macroscopic quantum tunneling effect, surface effect, good biocompatibility and unique optical properties, it can be combined with many kinds of biological macromolecules such as nucleic acid,^{16,18,19} heavy metal ions^{20,21} especially protein.^{22,23} Chang *et al.* detected a series with thiol compounds, including intracellular bioactive molecules such as cysteine and glutathione, and the positioning of the erythrocyte glutathione basing on interaction between gold and Nile red dye as a fluorescence.^{24–26} Meanwhile, the strength of interaction between gold nanoparticles and the elements such as nitrogen, sulfur in some groups of compounds to detect some heavy metal ions such as Cu^{2+} , Hg^{2+} .^{27,28} Dobrovalskaja *et al.* conducted a systematic study on blood plasma protein adsorption to citrate-protected gold nanoparticles and identified about 60 different proteins in the “protein corona” formed on the gold

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nanoparticle surface.²⁹ The local electric field enhancement of the surface of metallic nanoparticles in visible light and near infrared light is caused by the plasma resonance of several orders of magnitude of the incident electric field changes. The GNRs have a strong plasma enhancement effect in the range of 600–1500 nm.^{30,31}

Most of the proteins contain fluorescent amino acids, tryptophan and tyrosine. When the fluorescent tryptophan or tyrosine interacts with gold nanoparticles, the fluorescence properties of tryptophan or tyrosine will shift.^{29,32,33} These changes can occur as emission wavelength red-shift or blue-shift, fluorescence quenching or enhancement. Such changes have been applied to determine the binding affinity and binding constant of proteins with gold nanoparticles quantitatively. Proteins and gold nanoparticles are formed by the electrostatic attraction and hydrophobic force, which give the system a characteristic of fluorescence scattering peak that enhancing the fluorescence scattering signal of the associated nanoparticle. On the basis of the linear relationship between the fluorescence intensity and the concentration of proteins, a new method for protein determination was established. In this study, a simple, sensitive and stable method based on the combination of GNRs with the microalgae protein was explored for the microalgae protein analysis; the related pH condition was determined as well. The salt ions Na^+ and the stable time of the reaction system was also examined. A number of parameters affecting the fluorescence intensity were studied and experimental sensitivity were examined. The results demonstrated a higher sensitivity and selectivity in microalgae protein detection and a great potential for biology diagnosis can be brought by this method.

2. Material and methods

Chemicals and materials

The microalgae proteins were extracted from *Microcystis aeruginosa* (FACHB 905) culture which obtained from the Freshwater Algae Culture Collection of the Institute of Hydrobiology (Chinese Academy of Sciences, Wuhan, China), hexadecyltrimethylammonium bromide (CTAB), sodium citrate, sodium oleate, $\text{HAuCl}_4 \cdot 4\text{H}_2\text{O}$, silver nitrate, sodium borohydride, L-ascorbic acid (BioUltra) and hydrochloric acid were all obtained from SinopHarm Chemical Reagent Co., Ltd. (China).

Apparatus

Fluorescence spectra were obtained from fluorescence spectrophotometer (SHIMADZU-5301, Japan). Transmission electron microscopy (TEM) measurements were made on an electron microscope (JEOL-200CX, Hitachi) equipped with a camera (Model GATAN782CCD, USA) at an operating voltage of 120 kV. The samples for TEM studies were prepared by placing one droplet of the sample deposited onto carbon-coated copper grids.

Synthesis of GNRs

Gold nanorods were synthesized by utilizing seed-growing method.³⁴ First, the seed solution was prepared for gold NR

growth: 5 mL of 0.2 M CTAB solution was added to a 20 mL scintillation vial and 5 mL of 0.5 mM HAuCl_4 was mixed with the solution, 0.6 mL of fresh 0.01 M NaBH_4 was diluted to 1 mL with ultrapure water and was a trickle to join in the mixture under vigorous stirring (1200 rpm) for 2 min. The seed solution was kept for 30 min at room temperature. Next, the growth solution was prepared: 2.25 g of CTAB and 0.31 g of sodium oleate (NaOL) were dissolved in 62.5 mL of ultrapure water under vigorous stirring for 5 min, and the mixture was left undisturbed to cool down to 30 °C, then 4.5 mL of 4.0 mM AgNO_3 solution and 62.5 mL of 1 mM HAuCl_4 were added into the mixture and the solution was kept at 30 °C for 15 min, then the solution was stirred for 90 min and became colorless. Then 0.53 mL of 12.1 M HCl was dropwise into the growth solution. After stirring 15 min at 400 rpm, 0.31 mL of 0.064 M ascorbic acid (AA) was added into the solution stirred about 30 s. Finally, 0.1 mL of seed solution was injected into the growth solution. The resultant solution was vigorously stirred for 30 s and kept undisturbed at 30 °C for 12 h for gold NR growth. The size and shape of GNRs were characterized by JEOL-200CX transmission electron microscopy (TEM).

The extraction and preparation of *Microcystis aeruginosa* protein solution

Protein extraction from *Microcystis aeruginosa*: first, the sludge was milled into powder by liquid nitrogen, and the powder was suspended in four times volume of acetone with 2 mmol L^{-1} DTT and 10% TCA, followed by precipitation for 2 h at -20 °C. Next, the solution was separated with the speed of 9000 rpm for 15 min at 4 °C, precipitation was suspended in acetone with 2 mmol L^{-1} DTT for 2 h at -20 °C, followed by centrifugal separation for 15 min at 9000 rpm. The supernatant was discarded. Repeat the above steps for several times to extract enough materials. *Microcystis aeruginosa* protein can be obtained by the precipitation of the vacuum freeze drying.

Preparation of *Microcystis aeruginosa* protein solution: aqueous stock solution of *Microcystis aeruginosa* protein was achieved in a standardized way in water at room temperature and by magnetic stirring for 24 h and stored in the refrigerator. Reaction system was prepared by utilizing the protein solutions and dilution it into seven 10 mL volumetric flasks with 2 mL, 2.5 mL, 3 mL, 3.5 mL, 4 mL, 4.5 mL and 5 mL, respectively. Then 5 mL GNRs solutions were added into each flask, and bring the volume to 10 mL with distilled water. The solutions were stirred on magnetic stirrer followed by static mode for 10 min. The solutions were centrifuged and unreacted GNRs and protein residues were removed. The solutions were ready for fluorescence intensity detection. The pH value was adjusted with minute amounts of aqueous HCl (0.1 mol L^{-1}) or NaOH (0.1 mol L^{-1}) solutions, respectively.

3. Results and discussion

The mainly mechanism for the protein combining with the nanoparticles was shown in Fig. 1. The color of the reaction solution remained degraded pink after addition the algae protein. The protein structure containing inner base causes a strong repulsion between protein-groups and GNRs. The lack



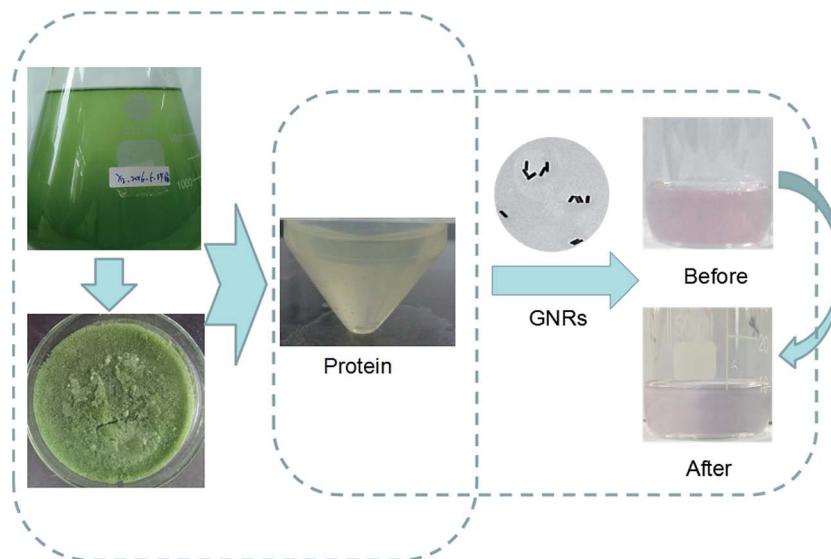


Fig. 1 Schematic representation of the strategy of algae protein combining with GNRs. Algae were collected and extracted for proteins followed addition of GNRs in containers for reactions and observations.

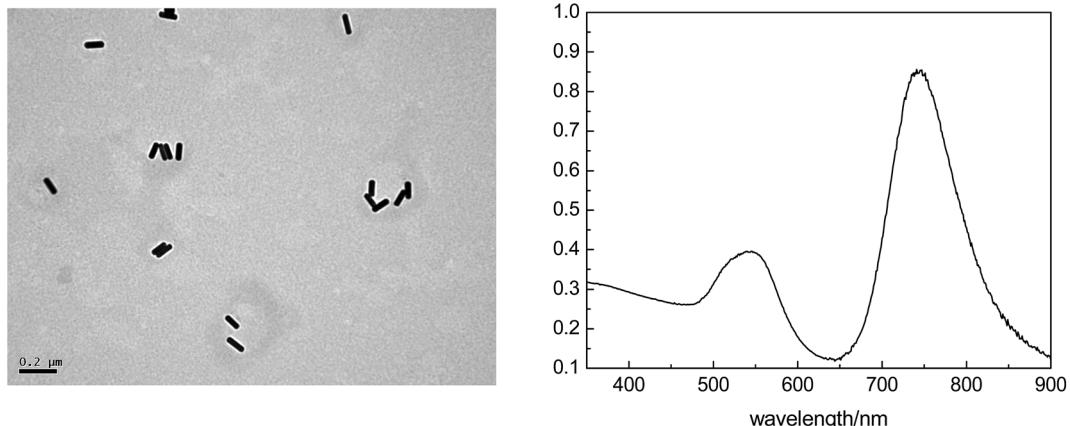


Fig. 2 TEM of GNRs (Left), and UV/vis absorption spectra of GNRs (Right).

of protection of adsorbed groups makes GNP s susceptible to aggregation, thus inducing color change of GNRs from the original pink to a degraded level.

Transmission electron microscopy (TEM) and ultraviolet absorption spectra of GNRs (UV-vis) are shown below. It can be seen its rod shape and the size is about $0.1\text{ }\mu\text{m}$ of GNR (Fig. 2).

The analysis of the interaction between *Microcystis aeruginosa* protein and GNRs

Fig. 3 illustrates the fluorescence spectra of the interactions between different concentrations of *Microcystis aeruginosa* proteins and GNRs. The fluorescence intensity was significantly enhanced with the increasement of *Microcystis aeruginosa* protein concentration and peak intensity at 560 nm in the system. Localized surface plasmon resonance (LSPR) of GNRs can produce a localized field enhancement effect except for GNRs with fluorescence.^{35,36} Hydrogen bonds exist between the

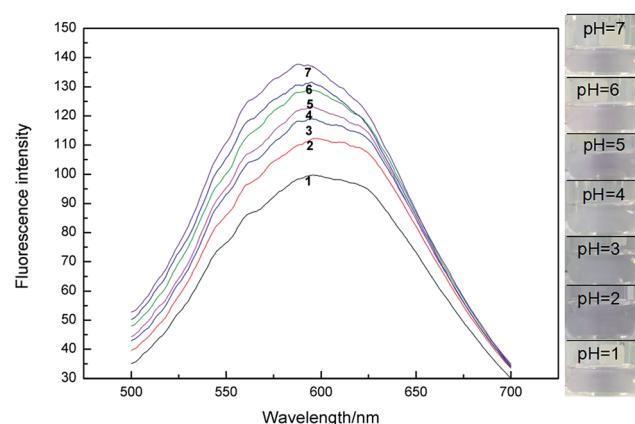


Fig. 3 Fluorescence spectra of interactions between different concentrations of *Microcystis aeruginosa* protein and GNRs.

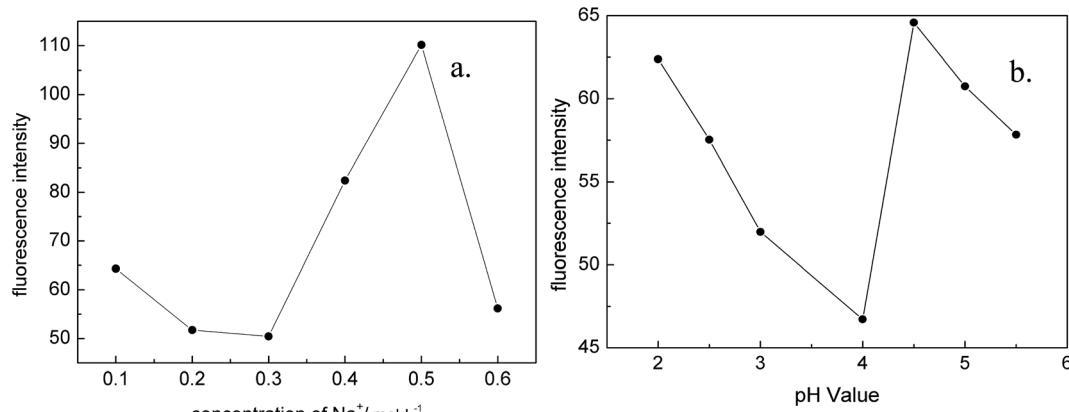


Fig. 4 (a) The effect of pH of buffer solution (pH = 2.0, 2.5, 3.0, 4.0, 4.5, 5.0, 5.5) GNRs: 5 mL; *Microcystis aeruginosa* protein 5 mL; (b) the effect of different concentration of Na^+ (mol L^{-1}) (Na^+ : 0.1/0.2/0.3/0.4/0.5/0.6).

amino groups, carboxyl groups and thiol groups in GNRs and *Microcystis aeruginosa* proteins.

On the one hand, the surface plasmon resonance of gold nanoparticles was changed, the incident light and the surface plasmon resonance of the gold nanoparticles absorb the same wavelength as the resonance frequency, and thus the characteristic absorption spectra was formed. The fluorescence enhancement includes the excitation and the emission enhancement. When the wavelength of the plasmon resonance wavelength was closed to the protein molecules, the emission intensity is enhanced and maximum the fluorescence enhancement. Ming *et al.* reported on the strong polarization dependence of the plasmon-enhanced fluorescence on single GNRs. The fluorescence from the organic fluorophores which embedded in a mesostructured silica shell around individual GNRs is enhanced by the longitudinal plasmon resonance of the nanorods. When the longitudinal plasmon wavelength and the excitation wavelength of the GNRs are almost equal, the fluorescence enhancement reaches the maximum value.³⁷ On the other hand, the covering layer of GNRs was formed on the surface of the *Microcystis aeruginosa* protein, which is much larger than that of aggregation of the GNRs. The Raman scattering signals and the interaction between GNRs and *Microcystis aeruginosa* proteins was enhanced. The result is demonstrated by Qin Long *et al.*^{38,39} that the higher the concentration of protein or amino acid, the stronger the signal intensity.

Optimization of experimental conditions for fluorescence detection

Based on the principles, there are two main factors that affect the experiment: pH value and the concentration of inorganic salt ion. To further understand the optimal experimental conditions, two approaches were developed for the analysis.

In the strategy 1: we studied the effect of pH on fluorescence intensity (Fig. 4a). In the pH range of 2.0–4.0, the fluorescence intensity decreased while the pH value increased. Specifically, when pH increased from 4.0 to 4.5, the fluorescence intensity increased; when pH increased from 4.5 to 5.5, the fluorescence

intensity decreased to the minimum value. At pH 4.5, the most sensitive response was obtained and the fluorescence intensity reached its maximum value.

When the pH is low, the *Microcystis aeruginosa* protein molecules partially deformed and the fluorescence intensity was decreased. The *Microcystis aeruginosa* proteins are the amphoteric electrolyte with isoelectric points, thus the pH of the solutions can change the solubility of the protein. With the increase of pH to 4.5 in the solution, the GNRs and *Microcystis aeruginosa* proteins have strong interactions, and the fluorescence intensity was enhanced. However, in the work presented by Yuyun Zhao *et al.*,⁴⁰ a strong interaction between gold nanoparticles were stabilized by sodium citrate and electric point greater than 7 of the protein in pH 7.4 solution, and the solution appears color change or coagulation flocculation phenomenon.⁴⁰ In our work, there is no similar phenomenon, the possible reason may be the pH value of the solution in this paper is less than 7 and the different structure of the protein.

In the strategy 2: we studied the effect of inorganic salt ion concentration on fluorescence intensity. When the concentration of Na^+ was 0.1–0.3 mol L^{-1} , the fluorescence

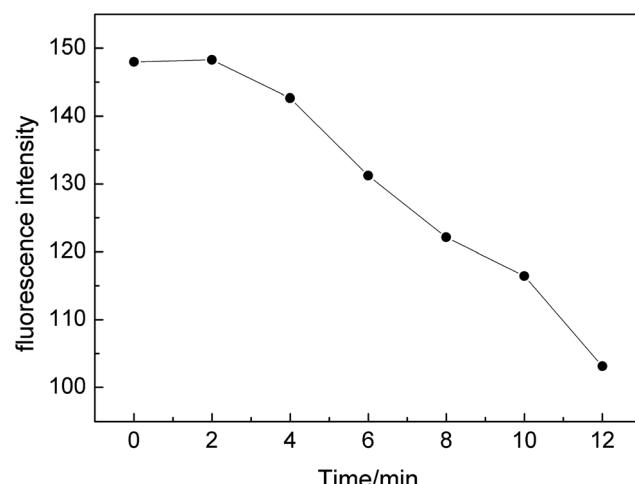


Fig. 5 The change of fluorescence intensity of different reaction time.



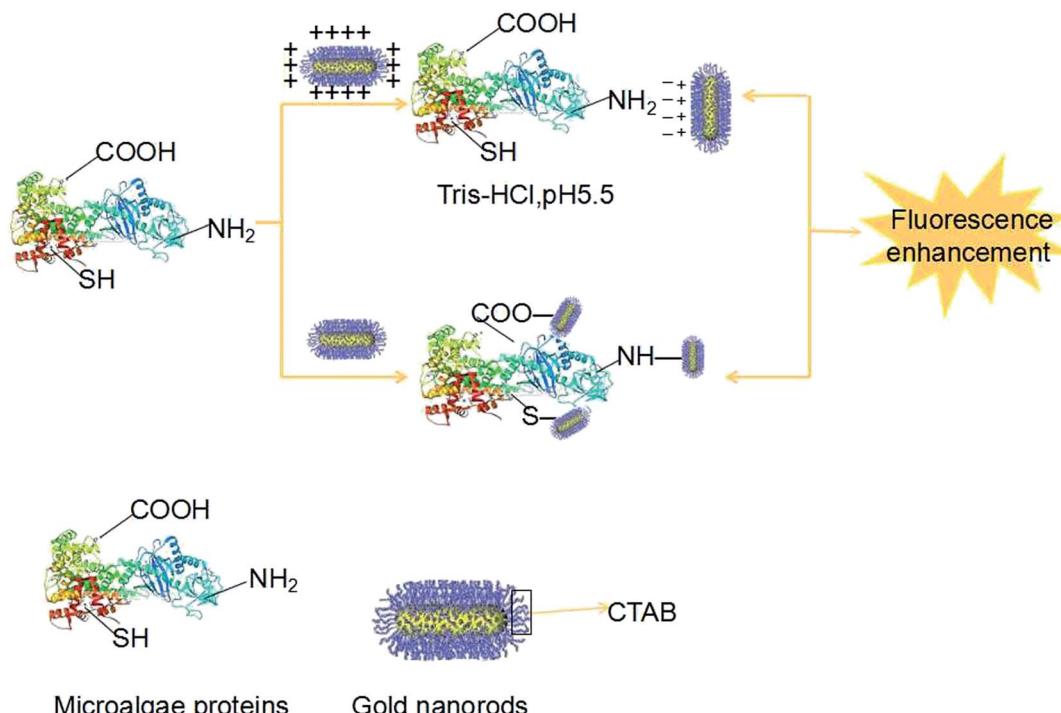


Fig. 6 Reaction mechanism of gold nanorods and protein.

intensity changed little. When the concentration of Na^+ was $0.3\text{--}0.5\text{ mol L}^{-1}$, the fluorescence intensity was obviously enhanced, and the most sensitive response appears at 0.5 mol L . When the concentration of Na^+ was above 0.5 mol L^{-1} , the intensity of fluorescence was decreased. It can be concluded that the optimal concentration of Na^+ was 0.5 mol L^{-1} as illustrated in Fig. 4b. This result was consist with the experimental studies by Pylaev T. E.⁴¹ and Javad Hassanzadeh.⁴² The interference of Na^+ , k^+ , Ca^{2+} , Ag^+ , Mg^{2+} , Zn^{2+} , Cu^{2+} , Fe^{3+} , Mn^{2+} were explored. The amounts of most potentially interfering species in serum are below their tolerable levels or can be decreased with diluting, so there would be no interferences from these species in *Microcystis aeruginosa* protein determination.

Sensitivity of experiments

The fluorescence intensity of the system reached the maximum in $0\text{--}2\text{ min}$, the combination of the GNRs and the *Microcystis aeruginosa* proteins was the highest. Subsequently, the fluorescence intensity gradually decreased in $2\text{--}12\text{ min}$ (Fig. 5). Sensitivity of the experiments have been examined in terms of detection limit and linear dynamic range. It is found that at the range of $5\text{ to }10$, the method exhibited linear gradient change of fluorescence intensity.

It is probably bound to facets at the end of GNRs *via* complex formation between its indole groups and empty orbitals on these facets about the combination of the GNRs and the *Microcystis aeruginosa* proteins was decreased. In the experimental study of Javad Hassanzadeh,⁴² the adsorption of albumin on the GNRs active surfaces inhibits their catalytic action and decreases fluorescence signal. But this result is not

consistent with the experimental results of Wu Menghui⁴³ and Jolanda Spadavecchia⁴⁴ using bovine serum protein (BSA), which may be due to differences in the structure of different proteins, resulting in differences in measurement results.^{45\text{--}48}

Discussion

The interactions between gold nanoparticles and proteins are due to the surface of the gold nanoparticles with more charges (Fig. 6). When the protein is equal to or slightly greater than the isoelectric point of protein in pH, the protein is neutral and the electrostatic interaction between protein and gold nanoparticles is small. However, the surface tension of protein molecules is very large in a weak hydration state, which is more easily adsorbed on the surface of gold nanoparticles.

Because of the protein molecules are firmly combined with the surface of the gold particles, a protein layer is formed to prevent the aggregation of gold nanoparticles. Gold nanoparticles have a strong adsorption to proteins and other biological macromolecules and will not change their biological activity, hence it has a good application prospect.

4. Conclusion

The present work has demonstrated that the combination of GNRs and proteins in *Microcystis aeruginosa* can enhance the fluorescence intensity with the increase of concentration of algae proteins. The mechanism involved in the formation of hydrogen bonds between the amino groups and carboxyl groups of the GNRs and the algae proteins which changed the surface plasmon resonance luminescence properties. When the GNRs



were combined with *Microcystis aeruginosa* proteins, the size of the polymers was much larger than that of the nanorods, which greatly enhanced the fluorescence intensity. This nanoparticle method has exhibited a high precision, sensitivity approach for the algal protein determination.

Abbreviations

GNRs	Gold nanorods
LSPR	Localized surface plasmon resonance
IR	Infrared
DTT	DL-Dithiothreitol

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