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# Fluorescence nanoprobe bearing low temperature-derived biochar nanoparticles as efficient quenchers for the detection of single-stranded DNA and 17 $\beta$ -estradiol and their analytical potential†

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Bagasse-derived biochar nanoparticles obtained under a low pyrolysis condition (400 °C) were first revealed to be capable of highly efficiently quenching the fluorescence of 6-carboxyfluorescein, with a significantly improved quenching rate constant over that of other quenchers and high-temperature prepared ones, and were designated as bagasse-derived quencher nanoparticles (BQNP). The BQNP are suitable for the construction of fluorescence nanoprobe, taking advantage of their various beneficial properties, including low cost, environmental friendliness, high dispersibility, and rich functional groups that allow their easy and versatile molecular modification. They were demonstrated to be capable of stably binding single-stranded oligonucleotides through both adsorption and covalent interactions and were utilized for the construction of both BQNP/DNA and BQNP/aptamer probes. The BQNP/DNA probe had strong resistance against degradation by deoxyribonuclease I and showed high precision and selectivity for the detection of single-stranded DNA, with a limit of detection of 1.04 nM. Moreover, the BQNP/aptamer probe demonstrated the rapid and sensitive detection of 17 $\beta$ -estradiol (E2) with a limit of detection of 0.4 ng mL<sup>-1</sup> with no cross-reactivity with the analogues, and it was also applied for real environmental sample detection and demonstrated reasonable signal recoveries. Benefiting from their strong quenching ability, low cost, and great dispersibility, the BQNP show great potential for the development of cost-effective and sensitive fluorescence sensors.

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## Introduction

Nucleic acid fluorescent probes are known to be sensitive and reliable and are widely used in biological and environmental analyses, as well as in clinical and medical diagnostics.<sup>1–3</sup> These sensors typically use fluorescence resonance energy transfer (FRET) or quenching mechanisms to detect signals.<sup>4</sup> They consist of a DNA probe molecule with a fluorophore and quencher molecule.<sup>5</sup> Nowadays, various kinds of nanomaterials have been employed to serve as quenchers, including carbon nanomaterials, gold nanoparticles, metal–organic frameworks, upconversion nanoparticles, and polymer nanostructures.<sup>3,6–8</sup> Nevertheless, it would still be worthwhile to explore novel

fluorescence nanoquenchers, featuring excellent quenching ability, good biological stability, easy preparation, and low cost, for constructing high-performance and cost-effective fluorescence sensing platforms.

As an emerging green and environmentally friendly carbonaceous material, biochar obtained from the pyrolysis of waste biomass has the properties of good source abundance from a wide range of sources, extremely low cost, easy synthesis, large specific surface area, porous structure, rich functional groups, and good stability. Recently, some additional promising characteristics of biochar have improved its attractiveness, such as high level sp<sup>2</sup> carbon structure representative of a good graphite structure, great defect structure, and electron transfer ability, which have broadened its application potential to not only soil amendment and chemical adsorption but also to advanced oxidative catalytic processes, electrocatalytic reactions, and capacitor and electrode constructions.<sup>9,10</sup> Based on its physical and chemical properties, it is likely that rationally synthesized biochar may present fluorescence-quenching capabilities, yet, to the best of our knowledge, this has been rarely examined to date. Accordingly, it would be constructive and meaningful to explore

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the fluorescence-quenching properties of biochar to develop low-cost and alternative fluorescence nanoquencher probes.

In the present study, bagasse-derived quencher nanoparticles (BQNPs) obtained from the pyrolysis of bagasse with a lower temperature setting (400 °C) were first demonstrated to have an outstanding fluorescence-quenching ability toward 6-carboxyl fluorescein (FAM). The quenching efficiency of the BQNPs was evaluated by the quenching rate constant, and this was also compared to other well-known carbonyl nanoquenchers. Since BQNPs are greatly dispersible, have a large surface area, and rich functional groups, they are ideally suitable for various modifications with biomolecules. As a proof of concept, the BQNPs were utilized to construct nucleic acid-based fluorescence nanoprobe and their detection performances toward single-stranded target DNA (tDNA) and 17 $\beta$ -Estradiol (E2) were assessed, as shown in Scheme 1. It is considered that the precise analysis of nucleic acids is significant for the diagnosis of cancer, infections, and genetic abnormalities,<sup>11</sup> and E2 is the most toxic estrogenic compound that is frequently found in varying water bodies, which can cause serious health problems.<sup>12,13</sup> The single-stranded oligonucleotides could be attached on the BQNPs by either adsorption or covalent interactions. The fabricated hybridization probe and aptamer probe were demonstrated to be capable of the sensitive detection of single-stranded DNA and E2 with a low detection limit and high selectivity. Hence, the BQNPs have great potential for developing low-cost quencher sensors.

## Materials and methods

### Chemicals and reagents

17 $\beta$ -Estradiol (E2,  $\geq 98\%$ ), bisphenol A (BPA,  $\geq 99\%$ ), Trizma hydrochloride (Tris-HCl), sodium chloride (NaCl), potassium chloride (KCl), magnesium chloride hexahydrate (MgCl<sub>2</sub>·6H<sub>2</sub>O), fluorescein, DNase I (deoxyribonuclease I), and *N*-(3-

dimethylaminopropyl)-*N'*-ethylcarbodiimide hydrochloride (EDC,  $\geq 99\%$ ) were obtained from Sigma-Aldrich (Shanghai, China). The other endocrine disrupter chemicals (EDCs), including ethinyl estradiol (EE,  $\geq 98\%$ ), were purchased from Aladdin Industrial (Shanghai, China). Estrone ( $\geq 99\%$ ) was obtained from Bailingwei (Beijing, China). Testosterone ( $\geq 98\%$ ) was purchased from Yuanye Biology (Shanghai, China).

All the DNA oligonucleotides labelled with or without 6-carboxyl fluorescein (FAM) were synthesized by Sangon Biotechnology Co. Ltd (Shanghai, China). The sequences of oligonucleotides used in this work were:

DNA probe: 5'-(FAM)-  
AAGGGATGCCGTTTGGGCCCAAGTTCGGCATAGTG-3'

Target DNA: 5'-CACTATGCCGAAGTTGGGCCCAAACG  
GCATCCCTT-3'

One-base mismatched target DNA:

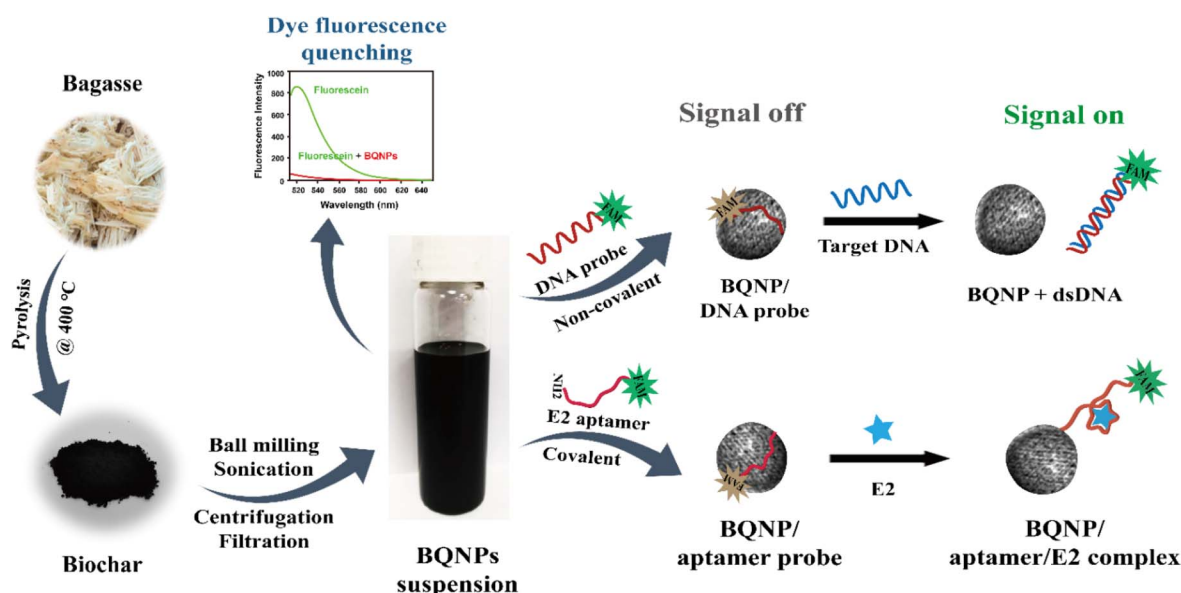
T1: 5'-CACTATGCCGAAGTTG  
GCCCAAACGGCATCCCTT-3'

T2: 5'-CACTATGCCGAAGTTGGTCCC  
AAACGGCATCCCTT-3'

T3: 5'-CACTATGCCGAAGTTGGACCCAA  
ACGGCATCCCTT-3'

Non-complementary target DNA:

T4: 5'-TAATACTGCCACTTCAAGACCTGGCGGCC  
ACCCGT-3'



Scheme 1 Schematic showing the procedure for synthesizing bagasse-derived quencher nanoparticles (BQNPs) and their utilization for constructing nucleic acid-based fluorescence nanoprobe to detect single-stranded tDNA and E2, respectively.



E2 aptamer probe: 5'-(FAM)-AAGGGATGCCGTTTGGGCC-CAAGTTCGGCATAGTG-(NH<sub>2</sub>)-3' (35 mer)

The stock solution of 100  $\mu\text{M}$  oligonucleotides was dissolved in deionized water and stored at  $-20\text{ }^{\circ}\text{C}$  before use. All the reagents were of analytical grade and used without further treatment. Deionized water, obtained from a Milli-Q water purification system (Millipore), was used in all the experiments. All the measurements were performed at room temperature in 100 mM Tris-HCl buffer (pH 8.0) containing 200 mM NaCl, 25 mM KCl, and 10 mM MgCl<sub>2</sub>.

### Characterization measurements

Fluorescent emission spectra were recorded using an RF-5301PC luminescence spectrometer (Shimadzu, Kyoto, Japan) with 1.0 cm micro quartz cells at room temperature. Absorption spectra were recorded on a 2600 UV-vis spectroscopy instrument (Shimadzu, Tokyo, Japan). Biochar samples were prepared in a tubular furnace (OTF-1200X, Hefeikeying Materials Technology Co., Ltd China). A planetary ball mill (Nanchang City Hengshun Chemical Equipment Manufacturing Co., Ltd) was used to grind the biochar. The zeta potential and size of the biochar were measured using a NANO ZS90 Zeta-sizer (Malvern Panalytical, UK). Transmission electron microscopy (TEM) images and high-resolution transmission electron microscopy images of the biochar were obtained by field emission transmission electron microscopy (FESEM, JEM-2200FS, Japan) on a super-thin carbon support film at an acceleration voltage of 200 kV. Fourier transform infrared spectroscopy (FTIR) was performed on an FTIR spectrometer (VERTEX-70, USA). Circular dichroism (CD) spectroscopy was performed using a J-810 circular dichroism system (JASCO, Japan), and the Raman spectra were recorded using a confocal microscope Raman spectrometer (LabRAM HR Evolution, HORIBA, France). The powder X-ray diffraction (XRD) patterns were recorded on a D8 advance diffractometer (Bruker, Germany) equipped with Cu K $\alpha$  radiation ( $\lambda = 1.54\text{ \AA}$ ) over the  $2\theta$  range of  $10\text{--}80^{\circ}$ .

### Preparation of the bagasse-derived quencher nanoparticles (BQNPs)

The raw biomass from agricultural waste bagasse was used to prepare the biochar nanoparticles. The bagasse was washed with deionized water three times to remove residual sugar and then exsiccated for 12 h at  $80\text{ }^{\circ}\text{C}$  in a vacuum drying oven. The dried bagasse was pyrolyzed at  $400\text{ }^{\circ}\text{C}$  in a tubular furnace with a flow of N<sub>2</sub> gas to prevent organic materials from igniting.<sup>14</sup> The heating rate for pyrolysis was  $7\text{ }^{\circ}\text{C min}^{-1}$ , and the residence time was 2 h. The produced rough carbon material (biochar) was ground into fine powder using a planetary ball mill for 2 h at 300 rpm, and then dispersed in deionized water, followed by ultrasonication for 10 min. Afterwards, the mixture was centrifuged at 13 000 rpm for 3 min, and the nanoparticles in the liquid supernatant were collected by filtration through a membrane (cut-off size of  $0.22\text{ }\mu\text{m}$ ). After drying in a vacuum

oven, these nanoparticles were ready for usage and denoted as bagasse-derived quencher nanoparticles (BQNPs).

### Preparation of the BQNPs/DNA probe and tDNA detection

A solution of BQNPs ( $100\text{ }\mu\text{g mL}^{-1}$ ) prepared in 100 mM Tris-HCl buffer (pH 8) was incubated with 30 nM of DNA probe for 5 min at room temperature. After thoroughly washing off the unbound DNA with Tris-HCl buffer, the produced composites of BQNPs and DNA probe (BQNPs/DNA probe) were ready for tDNA detection.

For DNA sensing, the BQNPs/DNA probe was incubated with various concentrations of tDNA ranging from 2 to 400 nM in 100 mM Tris-HCl buffer (pH 8.0) for 60 min at room temperature. The fluorescence intensities were measured with the excitation and emission wavelength of 492 nm and 518 nm, respectively.

The stability of the sensing system was tested by treating the BQNPs/DNA probe with DNase I ( $2\text{ U mL}^{-1}$ ) for 10 min at room temperature, and after inactivating the activity of the DNase for 10 min at  $80\text{ }^{\circ}\text{C}$ , the fluorescence profile and the target analyte detection ability were evaluated.

### Preparation of the BQNPs/aptamer probe and E2 detection

Conjugates of the BQNPs/aptamer probe were synthesized by coupling the amino group ( $-\text{NH}_2$ ) at the 3' termini of the FAM-labelled E2 aptamer probe to the carboxyl group ( $-\text{COOH}$ ) of BQNPs for E2 detection. Briefly, the carboxyl groups were first activated by EDC ( $10\text{ mg mL}^{-1}$ ) in MES buffer (100 mM, pH 5) for 20 min at room temperature. Upon activation, the solution was washed three times with deionized water to remove excess EDC. Subsequently, the activated BQNPs ( $200\text{ }\mu\text{g mL}^{-1}$ ) were reacted with FAM-labelled E2 aptamer probe (200 nM) in deionized water for 2 h at room temperature with gentle shaking (200 rpm). This was followed by centrifugation (10 min, 13 000 rpm) and thoroughly washing with deionized water to remove free aptamers. Then the obtained BQNPs/aptamer probe was stored at  $4\text{ }^{\circ}\text{C}$  for further use.

For E2 detection, a stock solution of E2 was prepared in ethanol and diluted with the binding buffer (100 mM Tris-HCl buffer) to various concentrations. Next, 50  $\mu\text{L}$  of BQNPs/aptamer probe ( $10\text{ }\mu\text{g mL}^{-1}$ ) dispersed in the binding buffer was incubated with specific concentrations of E2 ranging from 1 to 500  $\text{ng mL}^{-1}$  with shaking (200 rpm) for 15 min at room temperature. Then the fluorescence intensities of the solution were measured in a micro quartz cuvette cell with the excitation and emission wavelengths of 492 and 518 nm, respectively.

For real-sample environmental water detection, the sensor was applied to the analysis of E2 levels in three real water samples (tap water, lake water, and ground water). The tap water was obtained from the laboratory water tap. The lake water was obtained from Nanhu Lake, Changchun, China. The groundwater sample was obtained from the Changbaishan Hotel, Changchun, China. Before detection, the real water samples were filtered with a  $0.22\text{ }\mu\text{m}$  filter membrane to remove the suspended microorganisms and solid matter. Then the water samples were spiked to attain specific concentrations of



E2 (5, 8, 10, 20 ng mL<sup>-1</sup>), followed by quantitative sensing with the proposed method. All the measurements were performed in triplicate.

## Results and discussion

### Structural properties of the BQNPs

BQNPs were prepared from the biomass resources by the pyrolysis of bagasse at 400 °C under nitrogen protection, followed by grinding and size screening. The BQNPs were highly dispersive and presented nanosized and irregular structures (see TEM images in Fig. 1A). From the TEM image analysis, the average size of the BQNPs was 117.1 ± 38.4 nm (Fig. 1E). The particles from two further random batches exhibited a similar size distribution and average size value (Fig. S1†), indicating that the particle-generating approach was dependable and could be replicated. Upon further magnification under HR-TEM, many black dots smaller than 3 nm with an average size of 2.2 nm could be observed on the surfaces of the BQNPs (Fig. 1B), and almost each of them showed crystalline stripes representing a graphene-like nanosheet. Also, on the surfaces of BQNPs, aromatic cluster-like structures were visible (Fig. 1C and D), which was in accordance with the findings in rice-straw-derived biochar.<sup>15</sup> These structural characteristics would contribute to an easy and strong attachment of single-stranded DNA to the BQNPs.

The structural properties of the BQNPs were also characterized by means of Raman spectroscopy, FTIR, and XRD. The Raman spectra of the BQNPs (Fig. 1F) displayed D (~1343.2 cm<sup>-1</sup>) and G (~1595.4 cm<sup>-1</sup>) bands, which were assigned to local defects/disorders and stretching vibrations of the graphitized structure of the carbon atoms, indicating the crystallinity of the graphene nanosheets.<sup>16</sup> The intensity ratio of

the D and G bands ( $R = I_D/I_G$ ) was calculated as 0.57. The FTIR spectra of the BQNPs were obtained in the range of 400–4000 cm<sup>-1</sup> (Fig. 1G). The broad band at 3405 cm<sup>-1</sup> represented the band stretching of the hydrogen-bonded hydroxyl group (-OH), and the peaks at 1704 and 1600 cm<sup>-1</sup> were characteristic peaks of the aromatic C=C and C=O groups, respectively,<sup>17,18</sup> which contribute to the excellent water solubility and further modification capability. The XRD profile of the BQNPs exhibited two broad diffraction peaks around  $2\theta = 27^\circ$  and  $2\theta = 43^\circ$  (Fig. 1H) attributed to the crystal planes of parallel stacked graphene sheets and the presence of a honeycomb structure formed by sp<sup>2</sup> hybridized carbons,<sup>19,20</sup> consistent with the results from the HR-TEM analysis, and which could contribute to  $\pi$ - $\pi$  stacking interactions of single-stranded DNA. After DNA adsorption on the BQNPs, there was a negligible change in the corresponding diffraction peaks (Fig. S2B†), indicating there was no significant structural alteration caused by the DNA attaching.

### Fluorescence-quenching ability of the BQNPs

Fluorescence quenching denotes the process of decreasing the fluorescence intensity of a fluorophore by a quencher,<sup>21,22</sup> and it requires molecular contact between them. To investigate the quenching ability of the BQNPs, the fluorescence intensities of FAM after incubating in varying concentrations of BQNPs solution were examined. As shown in Fig. 2B, the fluorescence of FAM gradually decreased with the increase in BQNPs concentration. This observation demonstrates that the BQNPs could act as an efficient fluorescence quencher of FAM. Fluorescence quenching can occur through a static process, dynamic process, or a combination of both. Static quenching creates a non-emissive ground-state complex between the fluorophore and quencher, while dynamic quenching occurs when

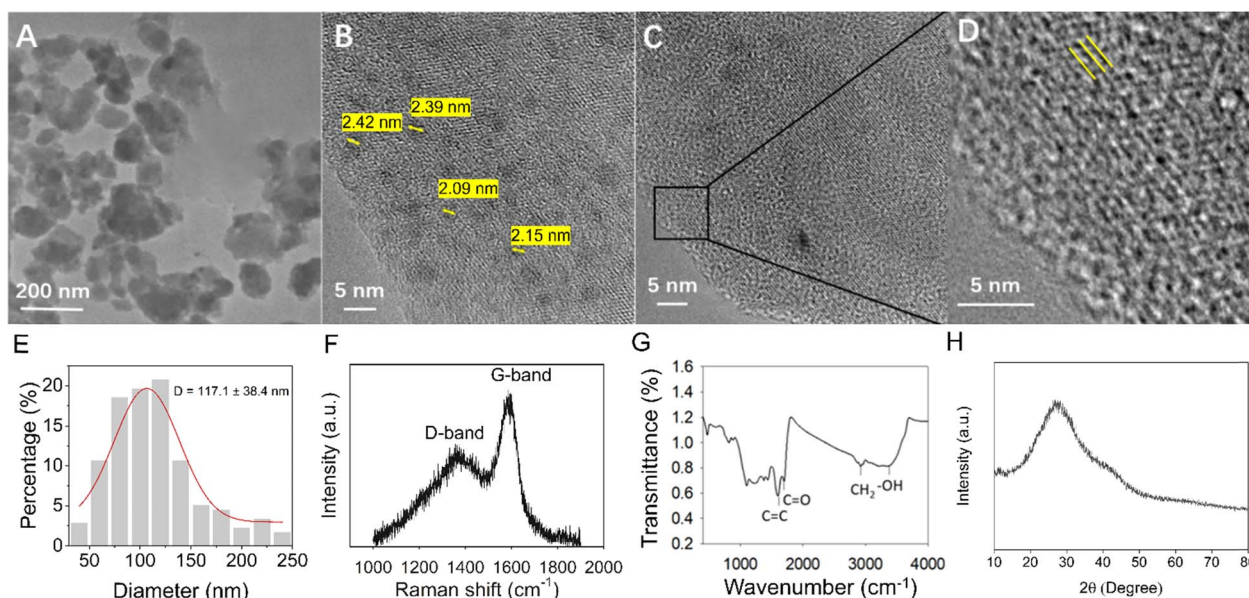


Fig. 1 Characterization of the BQNPs. (A) TEM image, (B) zoomed-in TEM image showing black dots on the surface, (C and D) HR-TEM images, (E) size distribution of BQNPs determined by quantitative analysis of the TEM images. Over 200 nanoparticles were measured. (F) Raman spectra, (G) FTIR spectra, and (H) powder X-ray diffraction pattern.



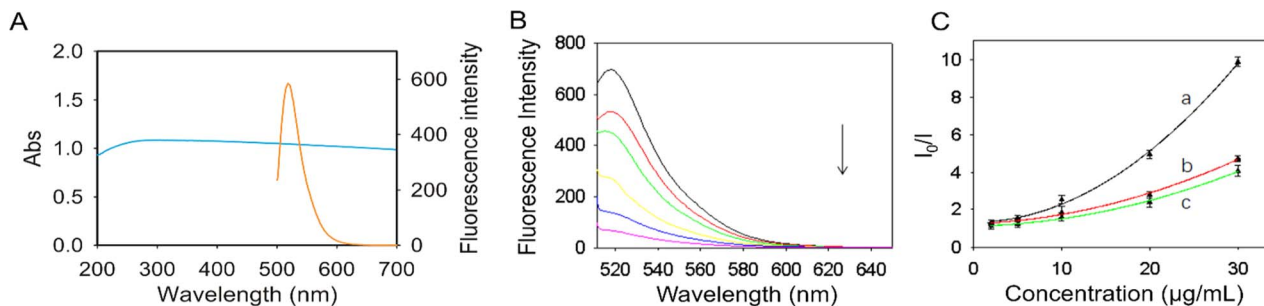


Fig. 2 (A) UV-vis adsorption spectrum (blue line) of BQNPs, and fluorescence emission spectrum of fluorescein (orange line). (B) Fluorescence spectra of fluorescein ( $1.3 \times 10^{-8}$  M) in water with different concentrations of BQNPs (0, 2, 5, 10, 20, 30  $\mu\text{g mL}^{-1}$ , from top to bottom). (C) Stern–Volmer plots of BQNPs (2, 5, 10, 20, 30  $\mu\text{g mL}^{-1}$ ) prepared at varying temperatures, 400 °C (curve a), 600 °C (curve b), and 800 °C (curve c).

the quencher deactivates the excited state of the fluorophore, often through a collisional mechanism.<sup>23</sup>

The Stern–Volmer plot is a graphical representation used to study the quenching of fluorescence and evaluate the fluorescence-quenching mechanism. The Stern–Volmer equation is given by:<sup>21</sup>

$$\frac{I_0}{I} = 1 + K_{SV}Q = 1 + k_q\tau_0Q$$

where  $I_0$  and  $I$  are the fluorescence intensities of a fluorescein dye in the absence and presence of a quencher, respectively,  $Q$  is the quencher concentration ( $\text{g L}^{-1}$ ),  $K_{SV}$  is the Stern–Volmer quenching constant ( $\text{L g}^{-1}$ ),  $k_q$  is the quenching rate constant ( $\text{L g s}^{-1}$ ), and  $\tau_0$  is the lifetime of the fluorescein dye in the excited state (s).<sup>24</sup>

If fluorescence quenching is due to a purely static or purely dynamic mechanism, the Stern–Volmer plot of  $I_0/I$  vs.  $Q$  will be linear. Based on the nonlinear nature of Stern–Volmer plot, it is possible that both dynamic and static quenching may be involved.<sup>23</sup> An upward curve in the Stern–Volmer plot of the system, as shown here in Fig. 2C, illustrates a complex fluorescence-quenching process, which consists of both static and dynamic processes for quenching fluorescence.

FRET is a non-radiative process in which energy is transferred from the excited state of a donor to an acceptor through dipole–dipole interactions.<sup>25</sup> The prerequisites for its occurrence are that the distance between the donor and acceptor is within the nanometre scale (smaller than 10 nm), and there is spectral overlap between the donors and acceptors.<sup>26,27</sup>

To investigate whether FRET could occur between BQNPs and FAM, the UV-vis absorption spectrum of the BQNPs and the fluorescence emission spectrum of the fluorescein FAM were obtained. As shown in Fig. 2A, the BQNPs absorption spectrum (blue line) exhibited a wide absorption spectrum from 200 to 700 nm, which indicates a good spectral overlap with the fluorescence emission spectrum (orange line) of FAM, which has an emission peak at about 518 nm. This suggests that FRET could occur between the BQNPs and FAM, in which the BQNP could be a fluorescence acceptor and FAM could be a fluorescence donor.

In addition, the quenching efficiencies of BQNPs prepared at varying pyrolysis temperatures (400 °C, 600 °C, and 800 °C) were investigated by Stern–Volmer analysis.

According to the Stern–Volmer plots (Fig. 2C), the  $k_q$  of BQNPs prepared at 400 °C ( $33.08 \times 10^9 \text{ L (g s)}^{-1}$ ) was higher than that prepared at 600 °C ( $12.14 \times 10^9 \text{ L (g s)}^{-1}$ ) and 800 °C ( $10.31 \times 10^9 \text{ L (g s)}^{-1}$ ), indicating that BQNPs obtained at lower temperature present the highest quenching ability towards FAM. This was supposed to be related to the greater amounts of carboxyl groups distributed on the surface of BQNPs obtained at lower temperature (see Fig. 1G and S2A†), since the carboxyl groups are generally considered good electron acceptors in the non-resonance energy-transfer process by deprotonating and negatively charging. Moreover, it was found that the  $k_q$  of BQNP was also higher than that of other reported quenchers, such as graphene sheets ( $1.25 \times 10^9 \text{ L (g s)}^{-1}$ ), single walled carbon nanotubes ( $0.233 \times 10^9 \text{ L (g s)}^{-1}$ ), and graphene oxide ( $0.03 \times 10^9 \text{ L (g s)}^{-1}$ ).<sup>28</sup> This was a possibility related to the high dispersibility and large surface area of the BQNPs, which enable more sites for access to FAM and fluorescence quenching.<sup>29,30</sup> This result highlights that the BQNPs would be a good choice as an alternative nanoquencher, and their great quenching ability would be significant for enhancing the fluorescence sensing signal through the increment of the signal to background ratio. Here, BQNPs usually refer to those prepared at 400 °C, unless otherwise explained.

### Feasibility of the BQNPs/DNA probe for DNA detection

**Synthesis of the BQNPs/DNA probe.** BQNPs derived from bagasse are good alternatives for developing fluorescent probes used in nucleic acid detection due to their excellent dispersibility, nanometre-sized properties, and high quenching ability. The schematic design of a tDNA detection system based on BQNPs as a fluorescent nanoquencher is shown in Scheme 1. Upon incubation, the FAM-labelled DNA probe specific to the target nucleic sequence would be readily adsorbed onto the surface of BQNPs through  $\pi$ – $\pi$  stacking between the oligonucleotide base ring structures and the honeycomb structures of the BQNPs (see XRD spectra in Fig. 1H) and through the hydrophobic interactions. These noncovalent physisorption interactions guarantee the close proximity of FAM to the BQNP surface, resulting in the quenching of the FAM signal by BQNP (signal off). When there is a tDNA, the DNA probe would specifically hybridize with the tDNA, forming a double-stranded structure (dsDNA), which would release from the surface of the



BQNPs due to the DNA bases being hidden inside the helical structure and no longer available for surface binding, and so only the negatively charged phosphate groups are exposed, causing negative charge repulsion between the phosphate groups and BQNP surface, allowing fluorescence signal recovery (signal on).

The feasibility of BQNPs as nanoquenchers for the fluorescence detection of nucleic acid was thus firstly demonstrated. As indicated in Fig. 3A, a strong fluorescence emission was presented at around 518 nm in the case of the FAM-labelled DNA probe (curve a), and the intensity did not change with the addition of the tDNA (curve b). It was found that when adding BQNPs to the DNA probe, more than 98% of the fluorescence was reduced (curve c), and when the tDNA was incorporated, approximately 63% of the fluorescence signal was recovered (curve d). This suggested that the DNA probe could attach to the surface of BQNPs, which brought the FAM molecules close to BQNPs, causing fluorescence quenching, and with the presence of the tDNA, the double-stranded DNA formed by the hybridization should be detached due to the lower binding force, resulting in the recovery of the fluorescence signal. The incomplete signal recovery may be related to the binding of a certain part of the tDNA to unoccupied surface areas on the BQNPs resulting in a decreased hybridization and recovery efficiency.

The possible mechanism for the binding of DNA probe to the BQNPs was assessed by isoelectric point (pI) analysis (Fig. S3†). By the pH analysis method, the pI of BQNPs was measured to be 7.2, suggesting that the BQNPs should be negatively charged in the detection solution (pH 8), which means that the BQNPs should not be electrostatically attracted to the DNA probe due to its negatively charged backbone. Instead, considering the presence of  $sp^2$  hybridized carbon in the BQNPs (XRD in Fig. 1H), the  $\pi$ - $\pi$  interaction between the aromatic ring in the nucleotide bases of DNA and the BQNPs would be primarily responsible for the attachment of the DNA probe.

In order to more in-depth confirm the binding of the DNA probe to the BQNPs, the structural change of DNA was investigated using circular dichroism analysis. As shown in Fig. 3B, the probe DNA presented characteristic ellipticity positive peaks at 267 and 289 nm, and a negative peak at 246 nm (curve b). Also in case of the BQNPs/DNA probe, the positive peaks were reduced,

while the negative peak disappeared, suggesting a structural alteration of the DNA probe, which occurred due to the attaching of the molecular backbone to the BQNPs (curve c). Moreover, after addition of the tDNA, there was a slight conformational change of the DNA probe caused by the hybridization (curve d).

**Factors affecting the DNA-detection performance.** To achieve optimal assay performance with the BQNPs/DNA probe, several experimental parameters, including concentration of BQNPs and incubation time for maximum quenching efficiency, along with incubation duration and temperature for optimal tDNA detection, were investigated.

The quenching efficiency was determined as  $(F_0 - F)/F_0$ , where  $F_0$  and  $F$  represent the fluorescence intensities of the DNA probe in the absence and presence of BQNPs, respectively. Fig. S4A† shows the fluorescence intensity changes of the DNA probe (30 nM) when reacted with different concentrations of BQNPs. The fluorescence intensity decreased as the concentration of BQNPs increased from 0 to 500  $\mu\text{g mL}^{-1}$ , and the highest quenching efficiency (99%) was achieved at 100  $\mu\text{g mL}^{-1}$  of BQNPs (Fig. S4B†).

A study of the fluorescence-quenching kinetics was next conducted to determine the time required to completely quench the fluorescence of the DNA probe by the BQNPs. The quenching kinetics was assessed by monitoring the changes of the fluorescence intensities immediately after the addition of the BQNPs. As shown in Fig. S4C,† the fluorescence decrease was swift within the first 1 min (quenching efficiency of 95%), and it gradually reached equilibrium within 5 min (quenching efficiency of 99%). Apparently, the BQNPs were very effective at quenching FAM labelled on the DNA probe, which was attributed to the rapid and strong attachment of the DNA probe. Therefore, to obtain the optimum sensing performance, 5 min was used for quenching the fluorescence of the DNA probe.

The effect of the incubation time for the hybridization of tDNA in the detection process was evaluated, and 60 min was found to be optimal to achieve the highest signal recovery (Fig. S4D†). The effect of the incubation temperature during tDNA detection was also investigated, since temperature changes may affect the buffer conditions and the stability of DNA. As shown in Fig. S4E,† the fluorescence sensing signal was extremely weak when the incubation temperature was 4 °C. In contrast, the response signals increased significantly to nearly the same levels in the cases of 25 °C and 40 °C, showing that the proposed probe has a wide range of working temperatures. In the subsequent investigation, the incubation temperature was set at 25 °C.

**Sensitivity, stability, and specificity of the BQNPs/DNA probe.** The quantitative detection of single-stranded tDNA was performed using the BQNPs/DNA probe under optimal conditions. As shown in Fig. 4A, the fluorescence intensities increased gradually with the increment in tDNA concentrations. Also, the change in fluorescence intensity could be expressed as  $(F - F_0)/F_0 \times 100\%$ , where  $F_0$  and  $F$  represent the fluorescence intensities in the absence and presence of tDNA, which was linearly correlated with the concentration of tDNA in the range of 2 to 50 nM, with a correlation equation of  $y = 0.6344x -$

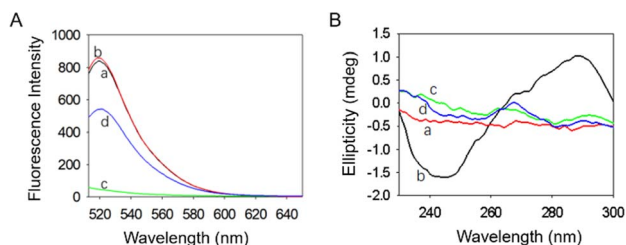
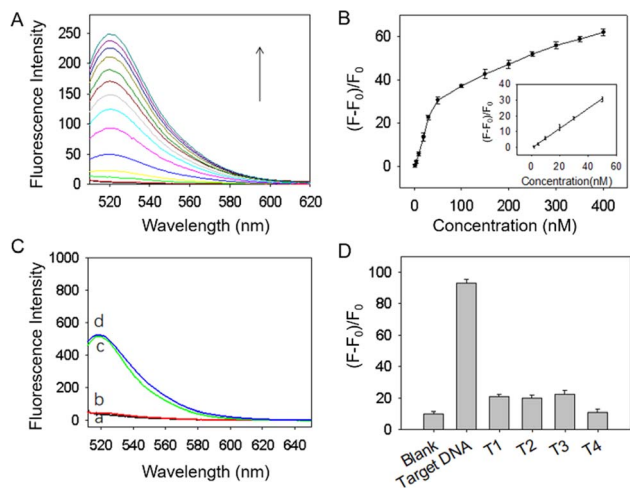


Fig. 3 (A) Fluorescence emission spectra of the DNA probe under various conditions: (a) DNA probe, (b) DNA probe + tDNA, (c) DNA probe + BQNPs, and (d) DNA probe + BQNPs + tDNA. The concentrations of BQNPs, DNA probe, and tDNA were 20  $\mu\text{g mL}^{-1}$ , 2 nM, and 2 nM, respectively. (B) Circular dichroism spectra of (a) BQNP, (b) DNA probe, (c) BQNPs/DNA probe, and (d) BQNPs/DNA probe + tDNA.





**Fig. 4** (A) Fluorescence emission spectra of DNA probe after incubation with different concentrations of tDNA (0, 2, 5, 10, 20, 30, 50, 100, 150, 200, 250, 300, 350, and 400 nM, from bottom to top). (B) Calibration plot for the quantitative detection of tDNA at different concentrations. Inset: Calibration curve for the low-concentration detection of tDNA. (C) Fluorescence emission profiles of the BQNP/DNA probe (formed by the DNA probe (2 nM) and BQNP (20  $\mu\text{g mL}^{-1}$ )) (a) without and (c) with tDNA (2 nM), and the fluorescence emission profiles of the BQNP/DNA probe after treating with DNase I (b) without and (d) with tDNA (2 nM). (D) Selectivity of the BQNP/DNA probe for DNA analysis. The concentrations of tDNA, T1, T2, T3, and T4 were all 30 nM and the concentration of BQNP/DNA probe was 200  $\mu\text{g mL}^{-1}$ . The error bars illustrate the standard deviations from triplicate measurements.

0.6696 and  $R^2 = 0.9988$ . The detection limit was 1.04 nM as calculated by the signal to noise ratio of three, which is comparable or superior to the previously reported methods (see Table S1†). Moreover, the relative standard deviation for tDNA detection was less than 7%, suggesting a reasonable reproducibility of the method.

The stability of the BQNP/DNA probe was evaluated by treating it with DNase I. According to Fig. 4C, the BQNP/DNA probe showed negligible changes in fluorescence signal after treatment with DNase I (see curves a and b), while after adding tDNA, the fluorescence signal recovered to almost its original level (see curves c and d). Clearly, the DNA probe on the BQNP was resistant to DNase I cleavage and even remained normally functional with its adsorption to the BQNP.

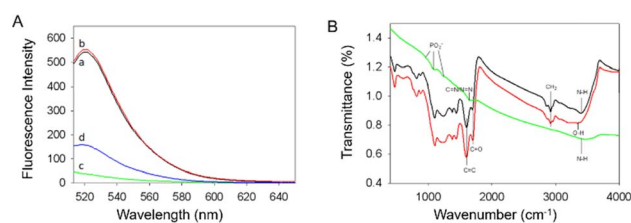
This sensing platform is based on the high selectivity of the single nucleotide chain, which contributes to its specificity. Therefore, we also evaluated the selectivity of the designed BQNP/DNA probe in response to irrelevant DNA, such as mismatched single-base DNA (T1, T2, and T3) and non-complementary DNA (T4). As shown in Fig. 4D, the fluorescence signal increased significantly after incubation with tDNA, whereas the responses to the non-complementary DNA and one-base mismatched DNA were negligible. Based on these results, the proposed BQNP/DNA probe demonstrated sufficient selectivity towards tDNA and potentially could be used for single nucleotide polymorphism analysis.

## Feasibility of the BQNP/aptamer probe for E2 detection

**Preparation of the BQNP/aptamer probe.** To achieve specific E2 detection, a BQNP/aptamer probe was additionally proposed by conjugating the FAM-labelled E2 aptamer (also called E2 aptamer probe) attached with an amino group ( $-\text{NH}_2$ ) at the 3' termini to the carboxyl groups ( $-\text{COOH}$ ) of BQNP, thus forming a BQNP/aptamer probe. The feasibility of the BQNP/aptamer probe for E2 detection is shown in Scheme 1. In the absence of E2, covalent conjugation ensures the close proximity of FAM labelled on the E2 aptamer to the BQNP, resulting in a quenching of the FAM signal by the BQNP. The presence of the target E2, on the other hand, causes the aptamer probe to change its secondary structure and form an aptamer/E2 complex after selectively capturing E2, which has a relatively weak affinity for BQNP and separates from the BQNP, thereby restoring FAM fluorescence.

To investigate whether the E2 aptamer probe was covalently bonded to BQNP, FTIR spectral profiles of the original BQNP, E2 aptamer probe, and BQNP/aptamer probe after removing the unbound aptamers were analyzed. As shown in Fig. 5B, the characteristic peaks of BQNP were observed at  $1600\text{ cm}^{-1}$  ( $\text{C}=\text{C}$ ),  $1700\text{ cm}^{-1}$  ( $\text{C}=\text{O}$ ), and  $3370\text{ cm}^{-1}$  ( $-\text{OH}$ ).<sup>17,18</sup> The characteristic peaks of the E2 aptamer were observed at 965, 1084, and  $1229\text{ cm}^{-1}$ , corresponding to phosphate asymmetric and symmetric vibrations, while the vibration at  $1649\text{ cm}^{-1}$  was due to the presence of both  $\text{C}=\text{N}$  double bonds and  $\text{N}=\text{N}$  double bonds.<sup>31</sup> In the BQNP/aptamer probe, there was another characteristic peak of the E2 aptamer at  $3416\text{ cm}^{-1}$ , which was ascribed to the nucleic acid bases of  $\text{NH}$  stretching,<sup>17,18</sup> indicating that the E2 aptamer was successfully combined with BQNP.

The viability of a fluorescent probe-constructed using BQNP and E2 aptamers was first established for E2 detection. As shown in Fig. 5A (curve a), the free FAM-labelled E2 aptamers exhibited a strong fluorescence emission at 518 nm, and after adding the target E2, the fluorescence signal was not changed (curve b), demonstrating that the fluorescence intensity of free aptamers was not affected by E2. In contrast, the BQNP effectively quenched FAM fluorescence by more than 98% in the case of the BQNP/aptamer probe (curve c). Furthermore, the quenching efficiency was higher than that of most reported nanoquenchers.<sup>32,33</sup> In the presence of the target E2, the aptamer changes its secondary structure upon the selective



**Fig. 5** (A) Fluorescence emission spectra of the E2 aptamer probe under different conditions: (a) E2 aptamer probe; (b) E2 aptamer probe + E2; (c) BQNP/aptamer probe; (d) BQNP/aptamer probe + E2. The concentrations of BQNP/aptamer probe, E2, and E2 aptamer were 10  $\mu\text{g mL}^{-1}$ , 500  $\text{ng mL}^{-1}$ , and 10 nM, respectively. (B) FTIR spectra of BQNP (red colour), E2 aptamer probe (green colour), and BQNP/aptamer probe (black colour).



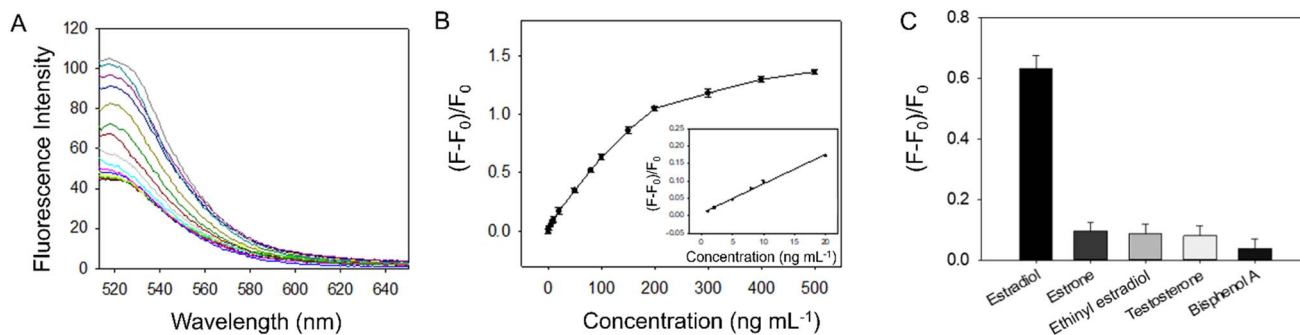


Fig. 6 (A) Fluorescence emission spectra of the BQNP/aptamer probe without or with E2 added at varying concentrations (1, 2, 5, 8, 10, 20, 50, 80, 100, 150, 200, 300, 400, and 500 ng mL<sup>-1</sup>, from bottom to top). (B) E2 concentration calibration plot for the proposed fluorescence probe. Inset: Calibration curve for E2 detection. (C) Specificity of the BQNP/aptamer probe for E2 (100 ng mL<sup>-1</sup>) detection. As much as 200 ng mL<sup>-1</sup> of irrelevant chemicals were detected, including estrone, ethinyl estradiol, testosterone, and bisphenol A.  $F$  and  $F_0$  are the fluorescence intensities in the presence and absence of E2, respectively. The error bars illustrate the standard deviations of three replicate measurements.

Table 1 Detection of E2 in water samples

Sample sources	Spiked E2 concentration (ng mL <sup>-1</sup> )	Mean found (ng mL <sup>-1</sup> )	Recovery (%)	RSD (%)
Tap water	5	4.86	97.2	2.8
	8	8.51	106.4	6.4
	10	10.38	103.8	3.8
	20	19.36	96.8	3.2
Lake water	5	4.76	95.3	4.7
	8	8.5	106.1	6.1
	10	10.37	103.7	3.7
	20	19.3	96.4	3.6
Underground water	5	4.76	95.2	4.8
	8	8.46	105.7	5.7
	10	10.4	104.1	4.1
	20	19.4	96.9	3.1

capture of the target E2, leading to the formation of aptamer/E2 complexes with weak affinities to BQNP, which allows the E2 aptamers to move away from the BQNP surface, hence allowing the FAM fluorescence to be recovered (curve d). The achievement of a partial signal recovery was possibly due to the binding of some E2 to unoccupied surface areas of BQNP, thereby the amount of E2 captured by the aptamer was reduced. In addition, it may possibly be related to incomplete detachment of the aptamers from the BQNP following E2 capture.

**Sensitivity and specificity of the BQNP/aptamer probe.** To explore the sensitivity of the sensing probe, varying concentrations of E2 were reacted with the BQNP/aptamer probe, and the fluorescence recoveries were monitored. Fig. 6A shows that the fluorescence intensity increased gradually with increasing the E2 concentration. With a linear equation of  $y = 0.0084x + 0.0074$  ( $R^2 = 0.9933$ ), the fluorescence intensity change ratio (where  $F_0$  and  $F$  represent the fluorescence intensities without and with E2, respectively) was linearly correlated with the concentration of E2 (1–20 ng mL<sup>-1</sup>) (Fig. 6B). Furthermore, this assay is portable, cost-effective, easy to operate, and has a limit of detection of 0.4 ng mL<sup>-1</sup> based on a signal to noise ratio of three, which is comparable to other methods (Table S2†).

Several irrelevant endocrine disrupting chemicals, including estrone (E1), ethinyl estradiol (EE), testosterone, and bisphenol

(BPA), were considered in order to test the specificity and selectivity of the proposed fluorescence probe. In Fig. 6C, it can be seen that the irrelevant chemicals exhibited negligible fluorescence responses despite their concentrations being 2-fold higher than the target molecules (E2). Consequently, the BQNP/aptamer probe demonstrated excellent specificity for detecting the target molecules due to their high affinity for the target molecules (E2).

**Practical applications of the BQNP/aptamer probe.** A practical evaluation of the proposed BQNP/aptamer probe was carried out by monitoring E2 in tap water, lake water, and underground water samples. For the three environmental water samples spiked with E2 at four different concentrations (5, 8, 10, 20 ng mL<sup>-1</sup>), high recoveries were achieved between 95.2% and 106.4% with RSDs of 2.8% and 6.4%, as shown in Table 1. Consequently, the BQNP/aptamer probe designed for water pollution monitoring was validated and is a reliable alternative to conventional methods to detect E2 in practical water samples.

## Conclusions

Bagasse-derived quencher nanoparticles (BQNP) obtained *via* low-temperature (400 °C) pyrolysis were demonstrated to possess a remarkable quenching ability towards the



fluorescence of 6-carboxyfluorescein. These nanoparticles exhibited a significantly higher quenching rate constant compared to those prepared at high temperatures and also compared to other typical quenchers. Taking advantages of the distinctive characteristics of BQNPs, including their high quenching efficiency, rich functional groups, and excellent dispersibility, as a proof of concept, BQNP/DNA and BQNP/apptamer probes were successfully constructed by adsorption and covalent interactions of single-stranded oligonucleotides. Both probes were capable of the selective and sensitive detection of single-stranded DNA and 17 $\beta$ -estradiol, with satisfactory detection limits of 1.04 nM and 0.4 ng mL<sup>-1</sup>, respectively. Because of the outstanding quenching performance, extremely good cost-effectiveness, great dispersibility, and environmental friendliness, the BQNPs have significant potential for the development of low-cost and sensitive fluorescence sensors.

## Data availability

The authors confirm that the data supporting the findings of this study are available within the article and its ESI.†

## Author contributions

Xiaoli Qi was responsible for conceptualization, data curation, formal analysis, investigation, methodology, validation, writing – original draft, writing – review & editing. Hui Hu was responsible for investigation, data curation, formal analysis, methodology, validation, visualization, writing – original draft. Yunxian Piao was responsible for funding acquisition, conceptualization, project administration, supervision, writing – original draft, writing – review & editing. Lina Liang and Yuqing Lin, were responsible for conceptualization, resources, writing – original draft. Yudan Liu and Haifeng Sun were responsible for methodology, formal analysis.

## Conflicts of interest

There are no conflicts to declare.

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