

## PAPER

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# Ratiometric fluorescent probe: a sensitive and reliable reporter for the CRISPR/Cas12a-based biosensing platform†

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Due to the excellent activity of *trans*-cleavage and target recognition, the recently discovered CRISPR/Cas12a systems provide a promising opportunity for designing fluorescence biosensing. In the reported CRISPR/Cas12a-based biosensing platform, TaqMan probe is widely used as the reporter. However, the TaqMan probe provides single-wavelength fluorescence changes, which is easily influenced by various analyte-independent confounding factors to produce false-positive signals. In this study, a ratiometric fluorescent probe was designed to act as the reporter of a CRISPR/Cas12a-based system. As a proof-of-concept, fluorescein (FAM) and tetramethylrhodamine (TAMRA) were chosen as the two fluorescence dyes to label one short ssDNA at 5' and 3' ends, respectively, which was designed as one ratiometric fluorescent probe. When the ratiometric probe excites at 480 nm, due to FRET effect, the probe emitted the 580 nm-fluorescence of TAMRA. The activated Cas12a can cleave the dual-labeled ssDNA, resulting in a decrease in the TAMRA's fluorescence and an increase in the FAM's fluorescence. This dual-response fluorescent probe can act as the reporter of the CRISPR/Cas12a-based biosensing platform. Compared with the classic TaqMan, CRISPR/Cas12a-based biosensing with the ratiometric probe as the reporter not only exhibited higher sensitivity but also could distinguish and avoid false positive signals.

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## Introduction

The CRISPR/Cas systems provide RNA-guided adaptive immunity to degrade the nucleic acids of intruding bacteria,<sup>1</sup> and these systems have been used as gene-editing tools.<sup>2,3</sup> Owing to the strict complementary-dependent cleavage principle, the CRISPR/Cas systems have very high accuracy in target nucleic acid recognition. The sequence-specific recognition capability of the CRISPR/Cas systems has been proved to be versatile to develop biosensors for the detection of nucleic acid<sup>4–6</sup> and non-nucleic acid targets.<sup>7,8</sup> The CRISPR/Cas12a exhibits non-specific ssDNase (single-stranded deoxyribonuclease) activity.<sup>5</sup> Based on this principle, many biological recognizing events have been converted to DNA to activate the *trans*-cleavage activity of the CRISPR-Cas12a system. In most of the CRISPR/Cas12a-based fluorescence biosensors, TaqMan probe is utilized as the substrate of Cas12a.<sup>5,8–15</sup> As one short single-

stranded DNA (ssDNA) (*ca.* 5 nt), the classic TaqMan probes are labeled with one quencher and one fluorophore in the 3' and 5' terminals, respectively. In a classic TaqMan probe, the fluorophore is close to the quencher. Due to the fluorescence resonance energy transfer (FRET) between the quencher and the fluorophore, the fluorescence emission of the fluorophore is “turn-off”. After being activated, Cas12a performs a high *trans*-cleavage on ssDNA.<sup>5</sup> Upon binding to the activated Cas12a, the TaqMan probe is cut, and the system emits strong fluorescence. Based on the “turn-on” fluorescence characteristic, TaqMan probe is widely used as a reporter to construct numerous homogenous CRISPR/Cas12a-based biosensors.<sup>5,8–15</sup> The CRISPR/Cas12a-based fluorescence sensing strategy with the TaqMan probe as the reporter is simple and rapid without any washing-steps and has been used to detect various targets, including DNA,<sup>5,10,11</sup> RNA,<sup>12,13</sup> small molecules<sup>8,9</sup> and enzyme activity.<sup>14,15</sup> However, the classic TaqMan probe always provides single-wavelength fluorescence changes, and the absolute-intensity-dependent signal is easily influenced by various analyte-independent confounding factors, which seriously disturb the accuracy of target quantitation.<sup>16</sup> Therefore, it is necessary to explore new reporters for the CRISPR/Cas12a-based biosensors.

In recent years, the ratiometric fluorescence method has attracted increasing attention because of its accuracy in detecting the target concentration.<sup>17–21</sup> Compared with the conven-

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tional fluorescence method with single-signal output, the ratiometric fluorescence method can eliminate the negative influence of complex environment through the ratio of fluorescence intensities of two wavelengths.<sup>18–21</sup> Thus, the ratiometric fluorescence methods can effectively overcome the false-positive signal and provide more accurate detection results.<sup>18–21</sup> There are two strategies for achieving ratiometric detection: one is the introduction of a target-insensitive fluorescence signal as the reference.<sup>22,23</sup> Another is to use two target-insensitive signals. In the fluorescence ratiometry with two target-insensitive signals, the addition of the target often leads to specifically decrease one fluorescence signal and increase the other, which makes the ratio of the two fluorescence signals change greatly. Therefore, the ratiometry strategy with two signal changes also possesses a higher sensitivity compared with the other fluorescence method.<sup>16,24</sup>

Enlightened by the aforementioned points, we herein introduced one ratiometric fluorescence probe into the CRISPR/Cas12a-based biosensing. As a proof-of-concept, fluorescein (FAM) and tetramethylrhodamine (TAMRA) were chosen as the two fluorescence dyes to label one short ssDNA at 5' and 3' ends, respectively, to construct one ratiometric fluorescence probe. Due to the FRET effect, the probe emitted the 580 nm-fluorescence of TAMRA. The activated Cas12a can cleave the dual-labeled ssDNA, resulting in the decrease of the TAMRA's fluorescence and increase of the FAM's fluorescence. This dual-response fluorescence probe can act as the reporter of the CRISPR/Cas12a-based biosensing platform. Compared with the classic TaqMan probe, this probe can provide ratio signal for the CRISPR/Cas12a-based biosensing system. This ratiometric CRISPR/Cas12a-based biosensing system exhibited lower detection limit for DNA target than that with the classic TaqMan reporter. In addition, this ratiometric CRISPR/Cas12a-based biosensing system could distinguish and avoid false positive signals, improving the accuracy of the results. Furthermore, we combined this ratiometric CRISPR/Cas12a-based biosensing system with enzyme-mediated strand-displacement amplification (ESDA) to improve the detection sensitivity. Overall, this ratiometric reporter opens a new pattern for signal output of the CRISPR/Cas12a-based biosensing strategy and holds a great potential to expand to other CRISPR/Cas-assisted biosensing.

## Experimental

### Reagents and materials

The HPLC-purified DNA and RNA were synthesized by Sangon Biotechnology Co., Ltd (Shanghai, China). The DNA and RNA sequences are listed in Table S1.† Before use, oligonucleotide solutions were diluted to the desired concentrations with 10 mM Tris-HCl buffer (50 mM NaCl and 10 mM MgCl<sub>2</sub>, pH 7.4). Nicking endonuclease (Nt-BstNBI), Vent (exo-) DNA polymerase and Cas12a were obtained from New England Biolabs, Inc. (Ipswich, US). Millipore Milli-Q water (*ca.* 18 MΩ cm) was used in the whole experiments. Other chemicals were of analytical grade.

### Procedure of the CRISPR/Cas12a-based biosensing of target DNA

First, Cas12a protein (50 nM) was incubated with the gRNA (50 nM) in 1 × 2.1 buffer for 10 min at room temperature (*ca.* 20 °C). Second, 25 μL RNase-free water, 5 μL 10 × 2.1 buffer and 5 μL ratiometric probe (1 μM) were added successively into the Cas12a-gRNA mixture. Then, 5 μL target DNA with different concentrations was added. The resulting mixture was incubated for 50 min at 37 °C. Lastly, the solution was added into 50 μL Tris-HCl buffer for fluorescence test. The fluorescence emission spectrum of the resultant reaction solution was collected on an FS5 Fluorimeter (Edinburgh, UK). The fluorescence spectra were recorded in the 508–650 nm range with an excitation wavelength ( $E_x$ ) of 480 nm. The ratio of the fluorescent intensity at 520 nm ( $F_{520}$ ) to the fluorescent intensity at 580 nm ( $F_{580}$ ) was used to quantify the concentration of target DNA.

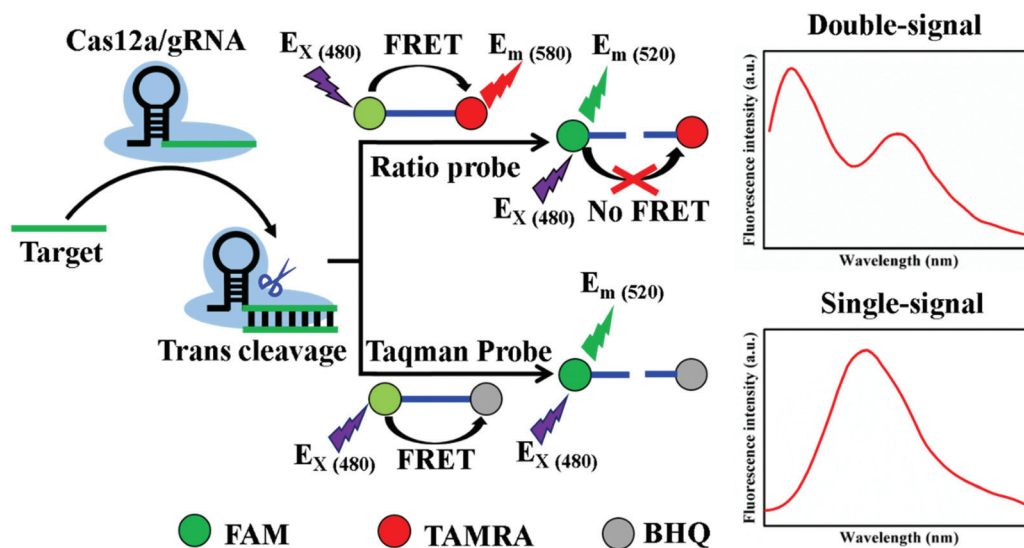
### Procedure of ESDA with the CRISPR/Cas12a-based biosensing for DNA detection

10 μL target DNA with different concentrations and 1 μL template DNA (0.1 nM) were added into 39 μL Nt-BstNBI nicking endonuclease buffer (pH 7.9 Tris-HCl (25 mM), containing 1.25 mM dithiothreitol, 125 mM NaCl, and 12.5 mM MgCl<sub>2</sub>). Then, the above mixture was heated at 90 °C for 5 min, and was naturally cooled to room temperature (*ca.* 20 °C). Subsequently, 39 μL ThermoPol buffer, 1 μL dNTP (25 mM), 5 μL Nt-BstNBI (1 U μL<sup>-1</sup>) and 5 μL vent (exo-) polymerase (1 U μL<sup>-1</sup>) were added into the above mixture. After reaction for 1.5 h at 55 °C, 30 μL ESDA product was mixed with 5 μL 10× Neb buffer 2.1, 5 μL ratiometric probe (1 μM), 5 μL gRNA (50 nM) and 5 μL Cas12a protein (50 nM) were added into the mentioned solution. Then, the solution was incubated at 37 °C for 50 min. The fluorescence spectra of the reaction solution were recorded. The ratio of  $F_{520}$  to  $F_{580}$  was used to quantify the target DNA concentration.

## Results and discussion

### Principle of ratiometric CRISPR/Cas12a-based biosensing strategy for target DNA detection

The principle of this ratiometric CRISPR/Cas12a-based biosensing is illustrated in Scheme 1. The guide RNA (gRNA) consisted of a 20 nt-long repeated sequence (black color) and a programmable target-recognizing sequence (green color), where the repeated sequence generally promotes gRNA to form the hairpin structure. The gRNA can interact with Cas12a protein to form a Cas12a/gRNA complex *via* sequence- and shape-specific interactions.<sup>25</sup> The Cas12a/gRNA complex can specific bind target DNA (ssDNA) to produce gRNA-DNA heteroduplex, and then Cas12a is activated to show the ssDNase activity. The activated Cas12a can effectively cleave the ratiometric probe (reporter). The ratiometric probe is dual-labeled ssDNA with FAM and TAMRA as the fluorescence donor and acceptor, respectively. The maximum excitation wavelength ( $E_x$ ) of FAM is 480 nm, and the maximum emission wave-



**Scheme 1** Principle of ratiometric probe or TaqMan probe as reporter in CRISPR/Cas12a-mediated biosensing platform.

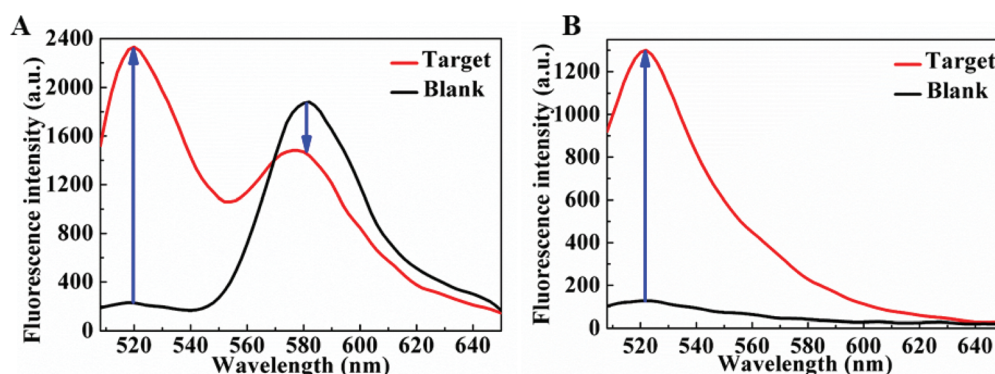
length ( $E_m$ ) of FAM is 520 nm. The  $E_m$  of TAMRA is 580 nm. When the ratiometric probe excites at 480 nm, the TAMRA's fluorescence at 580 ( $F_{580}$ ) is observed due to FRET. After the cleavage of the ratiometric probe, FRET is eliminated, resulting in appearance of a FAM's fluorescence at 520 nm ( $F_{520}$ ). Thus, the change of two fluorescence intensities can be monitored, and the ratio of  $F_{520}$  to  $F_{580}$  can be used to quantify the amount of target DNA. When the classic TaqMan probe is acted as the reporter of the CRISPR/Cas12a system, only one signal ( $F_{520}$ ) appears in the biosensing system.

In order to verify the feasibility of the ratiometric reporter, we added 0.5 nM target DNA to the CRISPR/Cas12a-based biosensing system. The fluorescence emission spectra of the system were recorded in the absence and presence of the target DNA. As illustrated in Fig. 1A, without the target DNA, FAM emission ( $F_{520}$ ) was rather weak (ca. 228), whereas TAMRA fluorescence ( $F_{580}$ ) was very strong (ca. 1870), indicating that FRET is quite efficient for the FAM-TAMRA pair. With target

DNA,  $F_{520}$  intensity increased from 228 to 2330, whereas the  $F_{580}$  intensity decreased from 1870 to 1470. Therefore, the signal-to-noise ratio of the ratiometric reporter was estimated to be ca. 13. In addition, we measured the fluorescence response of the biosensing system with the classic TaqMan as the reporter under the same experimental conditions. As shown in Fig. 1B, the presence of target DNA resulted in the increase of  $F_{520}$  from 127 to 1290. The signal-to-noise ratio of the TaqMan reporter was ca. 10. So, the sensitivity of the ratiometric probe was higher than that of the TaqMan probe.

### Optimization of experimental conditions

In order to improve the performance of the CRISPR/Cas12a-based biosensing, the experimental conditions were optimized. First, we explored the effects of the length of the ratiometric probe on the response. The three DNA probes with different bases numbers (5, 8 and 11) were designed to act as the reporter of the CRISPR/Cas12a-based biosensor. The



**Fig. 1** Fluorescence emission spectra of the CRISPR/Cas12a-based biosensing system in the absence and presence of target DNA with the ratiometric probe (A) or TaqMan probe (B). Experimental conditions: 5  $\mu$ L 50 nM gRNA, 5  $\mu$ L 50 nM Cas12a, 5  $\mu$ L 1  $\mu$ M probe, 5  $\mu$ L 0.5 nM target DNA, and 50 min reaction time.

results (Fig. S1†) showed that the 8-nt probe had the highest signal-to-noise, and was used as the ratiometric probe. We also optimized the other conditions of the CRISPR/Cas12a system, including incubation time, ratiometric probe concentration, Cas12a concentration and gRNA concentration (Fig. S2†). The optimized experimental conditions were 50 min of incubation time, 1  $\mu$ M of ratiometric probe, 50 nM of gRNA, and 50 nM of Cas12a.

### Performance of the CRISPR/Cas12a-based biosensing for target DNA

Under the optimal experiment condition, the fluorescence emission spectra of the CRISPR/Cas12a system with the ratiometric probe were obtained in the presence of target DNA with different concentrations. As shown in (Fig. 2A), with increasing target DNA concentration,  $F_{520}$  gradually increased, whereas  $F_{580}$  steadily decreased. The ratio ( $F_{520}/F_{580}$ ) increases linearly with target concentration in 0.01–0.75 nM range (Fig. 2B). The limit of detection (LOD) was calculated to be 3 pM ( $3\sigma$ ). For a comparison, we investigated the fluorescence response of the CRISPR/Cas12a-based biosensing with the TaqMan probe as the reporter under the same conditions (Fig. S3A†). The fluorescence intensity was linearly dependent on the target DNA

concentration in the range of 0.025 nM–0.25 nM (Fig. S3B†). The LOD of the TaqMan probe was estimated to be 10 pM ( $3\sigma$ ), which was above threefold higher than that of the ratiometric probe. We also checked the precision of the two reporters. Fig. 3 shows the results of 11 parallel measurements of the CRISPR/Cas12a-based biosensing at three different concentration levels (0.025 nM, 0.1 nM, and 0.25 nM). The relative standard deviation (RSD) of the ratiometric probe was estimated to be 2.1%, 2.4% and 1.7%, respectively (Fig. 3A). The RSD of the TaqMan probe was 4.8%, 4.2%, and 4.1%, respectively (Fig. 3B). It is obvious that the precision of the ratiometric probe is better than that of the classic TaqMan probe. The ratiometric probe is more resistant to fluctuations caused by the experimental environment.

Compared to the conventional single-signal probe, the ratiometric probe is expected to possess the capability of distinguish false positive signals.<sup>18–20</sup> As illustrated in Fig. 2A, the addition of the target DNA could lead to the increase of  $F_{520}$ , accompanying the decrease of  $F_{580}$ . If the increase of  $F_{520}$  and decrease of  $F_{580}$  in this biosensing system do not occur simultaneously, the signal will be false-positive. To prove this, we took bovine serum albumin (BSA) as an example to study the effects of some proteins on the response of the reporter. As

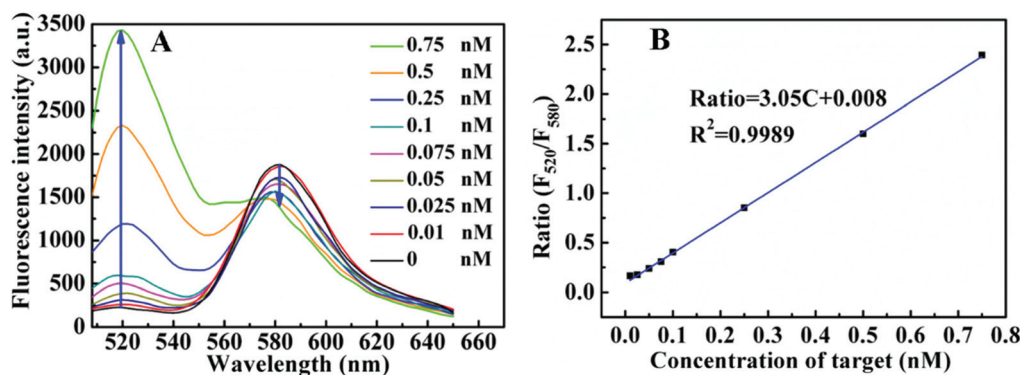


Fig. 2 (A) Fluorescence emission spectra of the CRISPR-Cas12a biosensing system in the presence of target DNA with increased concentrations, and (B) linear relationship between the fluorescence intensity ratio ( $F_{520}/F_{580}$ ) and target DNA concentration. Experimental conditions: 5  $\mu$ L 50 nM gRNA, 5  $\mu$ L 50 nM Cas12a, 5  $\mu$ L 1  $\mu$ M probe, 5  $\mu$ L different concentrations of target DNA.

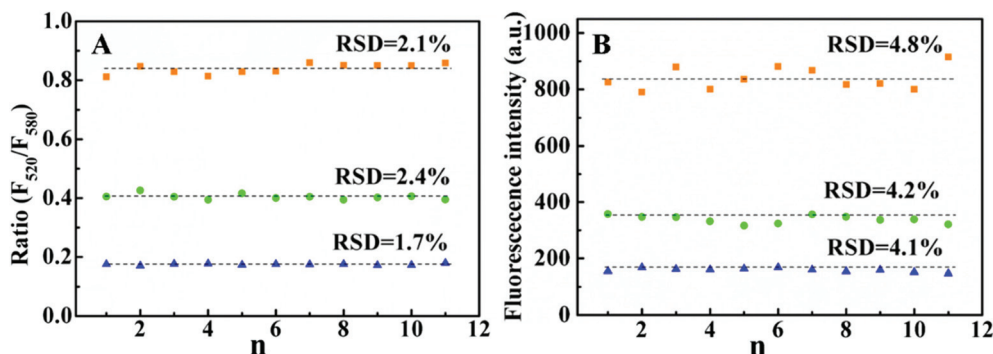


Fig. 3 11 parallel measurements of the CRISPR/Cas12a-based biosensing system in the presence of target DNA with different concentrations (0.025, 0.1, 0.25 nM from down to top) using the ratiometric probe (A) and TaqMan probe (B) as the reporter.



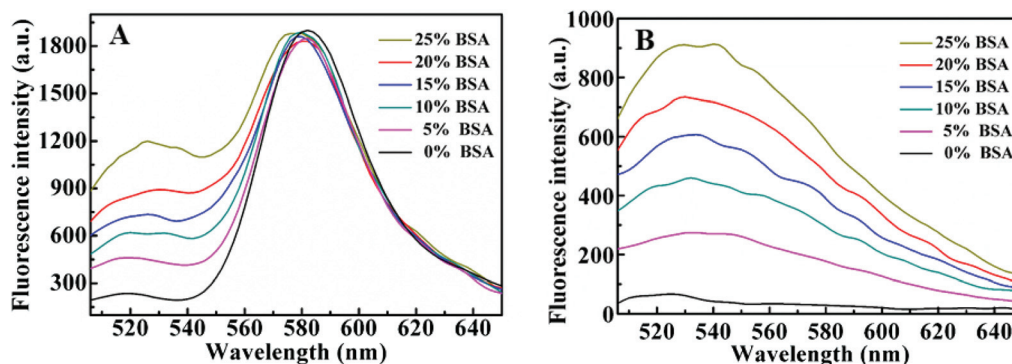


Fig. 4 Fluorescence spectra of the ratiometric probe (A) and TaqMan probe (B) in the presence of different concentrations of BSA.

shown in Fig. 4A, by increasing the BSA amount from 0 to 25%, the  $F_{520}$  intensity of the ratiometric probe increased evidently, whereas the  $F_{580}$  intensity did not almost change. From the response of this ratiometric probe, it is easy to judge that the BSA-induced change is a false positive signal. However, for classic TaqMan probe, a gradually increasing false-positive signal was observed (Fig. 4B), which is undistinguishable from a true target response. In addition, the similar result was observed when  $\text{Ca}^{2+}$  was added into the reporter probe.  $\text{Ca}^{2+}$  could decrease the intensities of  $F_{520}$  and  $F_{580}$  in this ratiometric probe response (Fig. S4†). This may be attributed to that  $\text{Ca}^{2+}$  can bind to the phosphate backbone of the ratiometric DNA fluorescent probe,<sup>26</sup> reducing the solubility of the DNA probe. However, as shown in Fig. 5A, the ratio of  $F_{520}$  to  $F_{580}$  remained unchanged when increasing  $\text{Ca}^{2+}$  from 0 to 10 mM. This is to say, the ratiometric probe can effectively avoid the false positive signal from  $\text{Ca}^{2+}$ .  $\text{Ca}^{2+}$  also decreased the fluorescence of the classic TaqMan probe (Fig. 5B), but the classic TaqMan probe cannot distinguish whether the decrease of the fluorescence signal is due to the change of the  $\text{Ca}^{2+}$  concentration or the change of target concentration. Therefore, the ratiometric probe as a reporter of the CRISPR/Cas12a-based biosensing can distinguish and even avoid false positive signals. The ratiometric probe is suitable for an *in vivo* assay.

Currently, there is a lack of an appropriate carrier to deliver the CRISPR/Cas12a system into living cells.<sup>27</sup> Therefore, we did not apply the CRISPR/Cas12a ratiometric fluorescent probe in practical applications.

#### Combining enzyme mediated strand-displacement amplification with CRISPR/Cas12a-based biosensing

Enzyme mediated strand-displacement amplification (ESDA) is an efficient tool for amplified assay.<sup>28–30</sup> To further enhance the sensitivity of the ratiometric CRISPR/Cas12a-based biosensing, we combined ESDA with this CRISPR/Cas12a-based biosensing. As shown in Fig. 6A, the ESDA template contained three domains: binding region of the target DNA (red), nicking site (purple) and region for generating activator (green). The hybridization of the target DNA and the template triggers the ESDA process. In the presence of dNTPs and DNA polymerase, the partial duplex (template/target) is extended along the template to form a complete dsDNA with one nicking site. Nicking endonuclease cleaves the extended DNA strand to release the activator, and simultaneously the cleavage reaction can trigger a new extension—cleavage—release cycle. As a result, the ESDA reaction continuously occurs, and many activators can be produced. The binding of the activator onto gRNA activates the *trans*-cleavage activity of Cas12a to cleave the ratiometric

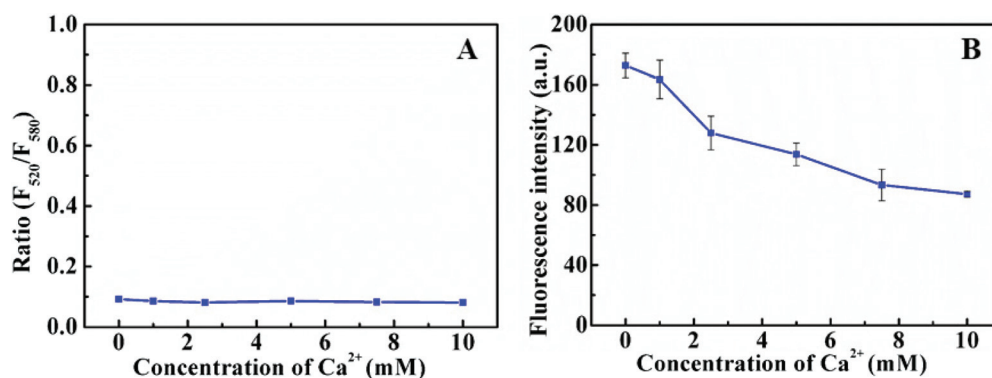
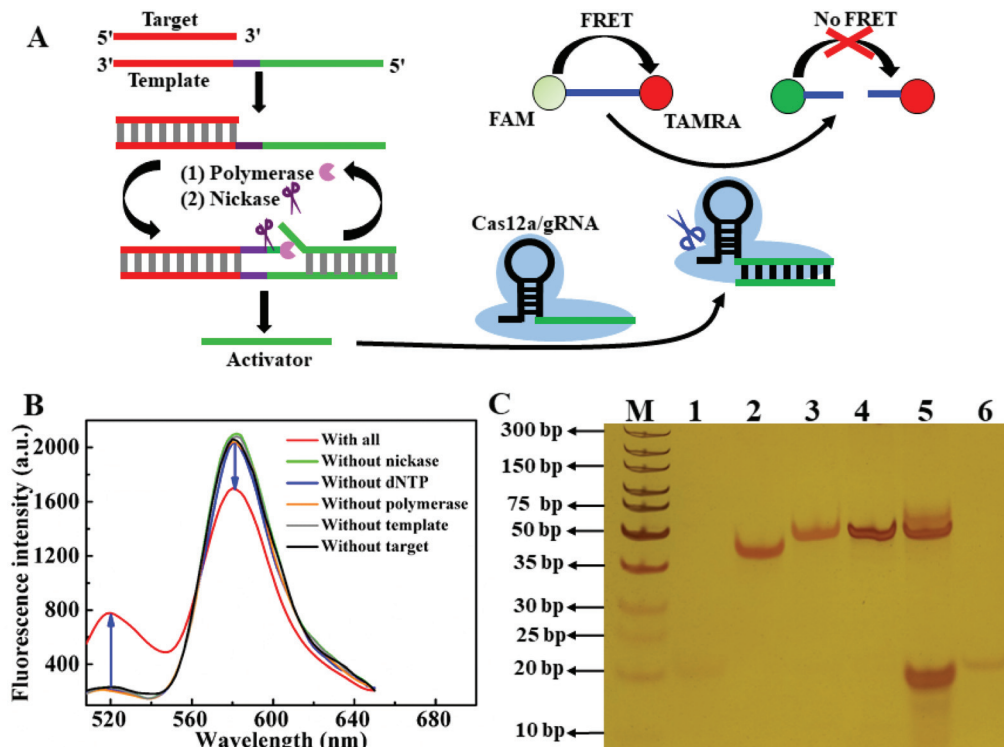


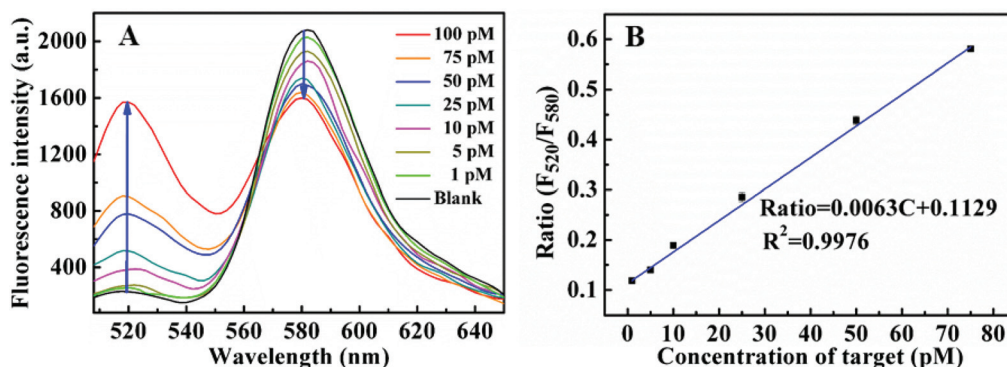
Fig. 5 Effect of  $\text{Ca}^{2+}$  concentration on the fluorescence response of the ratiometric probe (A) and TaqMan probe (B) in 20 mM Tris-HCl (pH 7.4) buffer.



**Fig. 6** (A) Scheme of combining ESDA and CRISPR/Cas12a-based ratiometric biosensing platform for DNA detection, (B) fluorescence spectrum of the sensing system under different conditions, and (C) the gel electrophoresis under different conditions, lane M: DNA marker; lane 1: target; lane 2: template; lane 3: target + template; lane 4: target + template + vent (exo-) polymerase + dNTPs; lane 5: target + template + Vent (exo-) polymerase + dNTPs + nickase; lane 6: activator.

probe, generating an obvious fluorescence response. As shown in Fig. 6B, when template, polymerase, nicking endonuclease and dNTP co-existed in the reaction system, the addition of 50 pM target DNA resulted in the signal change ( $F_{520}$  increase and  $F_{580}$  decrease). However, when any of four components (dNTPs, template, nicking endonuclease and polymerase) was absent, target DNA did not induce a significant change of fluorescence. To certify whether the signal amplification would work as designed, we performed the gel electrophoresis experi-

ments. As shown in Fig. 6C, the clear band on lane 1, 2 and 6 corresponded to target (22 nt), template (51 nt) and activator (20 nt). When the target was mixed with the template, a new upper band appeared in lane 3, which indicated the formation of a partial duplex between the target and the template. Once polymerase and dNTP were added into the system, a dark band in lane 4 emerged (51 bp), suggesting that the partial duplex was extended along the template DNA to produce the complete duplex. When nickase was unceasingly added into



**Fig. 7** Fluorescence spectra (A) and calibration curve (B) of the ESDA-CRISPR/Cas12a sensing system in the presence of increasing target DNA concentrations using the ratiometric probe as the signal reporter. Experimental conditions: 30  $\mu\text{L}$  ESDA product was mixed with 5  $\mu\text{L}$  10 $\times$  NEB buffer 2.1, 5  $\mu\text{L}$  ratiometric probe (1  $\mu\text{M}$ ), 5  $\mu\text{L}$  gRNA (50 nM) and 5  $\mu\text{L}$  Cas12a protein (50 nM).

the system, the band at 51 bp became weak (lane 5), and in the meantime a dark band appeared at the same position as the activator (lanes 5 and 6). The gel electrophoresis results showed that a small number of targets were transformed into a large number of activators through ESDA, thereby enhancing the sensitivity of the CRISPR/Cas12a-based method.

Under the optimized experiment condition (Fig. S5†), we investigated the fluorescent response of the ESDA-CRISPR/Cas12a-based biosensing system to different concentrations of the target DNA in the 1–100 pM range. As shown in Fig. 7A, by increasing the target DNA concentration,  $F_{520}$  gradually increased, whereas  $F_{580}$  steadily decreased. The ratio ( $F_{520}/F_{580}$ ) increases linearly with target DNA concentration in the range from 1 pM to 75 pM (Fig. 7B). According to the  $3\sigma$ , the LOD was estimated to be 0.3 pM, which was about 10-fold lower than that of the ratiometric CRISPR/Cas12a sensing system without ESDA. For comparison, we also investigated the fluorescence response of this biosensing system using the TaqMan probe as the reporter under the same conditions (Fig. S6†), and the LOD was calculated to be 2 pM, which was much higher than that of the ratiometric probe. These results further suggested that the ratiometric probe had higher sensitivity than the classic TaqMan probe.

## Conclusions

In summary, we used two fluorophores as the donor and acceptor to design a ratiometric probe for expanding the output pattern of a single signal readout. The ratiometric probe was used as the reporter of the CRISPR/Cas12a-based biosensing system. Compared with the classic TaqMan as the reporter, the CRISPR/Cas12a-based biosensing with the ratiometric probe as the reporter not only exhibited higher sensitivity but also could distinguish and avoid false-positive signal. This study shows that the ratiometric fluorescence probe is an effective reporter for the CRISPR/Cas12a-based biosensor system, which will promote the development of the ratio fluorescence method. Furthermore, the ratiometric fluorescence probe may provide great potential for other CRISPR/Cas enzyme-assisted biosensors.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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