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## Antibody-free colorimetric detection of chlorothalonil in cucumber *via* the inhibition of an enzyme-triggered reaction†

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Currently known rapid determination of fungicides usually relies on antibody-based immunoassay. This paper reports a simple antibody-free colorimetric assay for chlorothalonil *via* the inhibition of an enzyme-triggered reaction. The enzymatic activity of glyceraldehyde-3-phosphate dehydrogenase was significantly inhibited by chlorothalonil, and the color change of NBT-PMS system induced from NADH formation was suppressed, which could be used indirectly to assay chlorothalonil. The limit of detection (LOD) was 0.05  $\mu\text{M}$  with a linear range from 0.5 to 10  $\mu\text{M}$ , and the detection of 1  $\mu\text{M}$  chlorothalonil in solution was achieved with a naked-eye readout. In addition, the colorimetric measurement results of the cucumber samples showed a good recovery rate, although the sensitivity was less effective than the instrumental method. Nevertheless, the results demonstrate that the colorimetric method provides potential opportunities for reliable, cost-effective quantitative detection for chlorothalonil residues in vegetables.

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## Introduction

As a broad-spectrum fungicide widely used on fruits and vegetables,<sup>1–3</sup> chlorothalonil (2,4,5,6-tetrachloroisophthalonitrile, CTL) residue in raw fruits and vegetables and their processed products has drawn wide public attention. Due to its highly toxic to aquatic species and marine organisms, significant cumulative toxicity<sup>3,4</sup> and more toxic and persistent of its main metabolites (4-OH-CTL) in the environment, CTL and its metabolites are considered as a probable human carcinogen by the United States Environmental Protection Agency (USEPA).<sup>12</sup>

The most popular approaches for accurate and precise determination of CTL are based on highly sensitive detectors equipped with chromatographic units such as gas chromatography (GC) coupled with mass spectrometry (MS)<sup>13–15</sup> or electron capture detectors,<sup>16,17</sup> and HPLC with DAD detection<sup>18</sup> or coupled with MS.<sup>6</sup> However, susceptibility to factors such as pH<sup>8</sup> and matrix components<sup>19</sup> makes CTL a nasty pesticide of poor recovery for sample preparation prior to chromatographic analysis. In addition, the need of specialized persons and high cost limit their application for on-site screening. Antibody-based immunoassay systems such as enzyme-linked immunosorbent assay<sup>20–23</sup> (ELISA), immunosensor,<sup>7</sup> immunochip<sup>24</sup> and

immune-strip<sup>25–28</sup> offer potential opportunities for rapid detection of CTL on-site and substantial investigations have been reported and there are already some test kits availability on the market. Despite its high specificity and sensitivity, the preparation process of antibodies is high cost and time-consuming. Moreover, susceptibility to degradation and denaturation of antibodies remains a great challenge during its application. It should be noted that the use of nanoparticles<sup>29</sup> and self-propelled micromotors<sup>30</sup> with inherent enzymatic (artificial enzymes) activity display attractive performance in sensing of phenylenediamines isomers. In spite of its facile synthesis and stable performance, the lack of specific recognition limits their target scope. Unfortunately, there are frequent reports about detection of CTL residue in many types of vegetables and fruits.<sup>31–33</sup> Thus, consumers need a simple, and inexpensive sensor system to monitor the risk of CTL in fruits and vegetables.

It is well known that enzyme-based signal amplification is commonly used in biochemical assays because of their high catalytic efficacy. Long known as a key enzyme in glycolysis, glyceraldehyde-3-phosphate dehydrogenase (EC 1.2.2.12; GAPDH) has recently been assigned numerous other cellular functions.<sup>34</sup> Due to its unique characteristics such as good stability, rapid response, high catalytic efficiency, low cost and biocompatibility, GAPDH is extensively employed in clinical diagnosis and drug design.<sup>35–37</sup> GAPDH catalyze the oxidative phosphorylation of glyceraldehyde-3-phosphate with concomitant reduction of  $\text{NAD}^+$  into  $\text{NADH}$ .<sup>38</sup>

Herein, we report an antibody-free method for visual detection of CTL utilizing a commercially available enzyme GAPDH

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coupled with phenazine methosulphate (PMS)- $\beta$ -nicotinamide adenine dinucleotide (NADH)-nitroblue tetrazolium (NBT) chromogenic system. In addition, we have successfully applied this method for detection of CTL in cucumber with good recovery, demonstrating that quantification of CTL concentration by measuring the color change based on the inhibition of enzyme-triggered reaction has a further potential application for detection of CTL on-site.

## Experimental

### Reagents

All chemical used were commercially available. CTL was purchased from Aladdin. Glyceraldehyde 3-phosphate (GAP) and glyceraldehyde-3-phosphate dehydrogenase (GAPDH) were obtained from Sigma (U.S.).  $\text{NAD}^+$ , nitrotetrazolium blue chloride (NBT), sodium pyrophosphate, sodium dihydrogen phosphate, and dibasic sodium phosphate were got from Sinopharm Chemical Reagent Co., Ltd (Shanghai, China). Phenazine methosulfate (PMS) was bought from Sangon Biotech (Shanghai, China). Pentachloronitrobenzene (PCB), pentachloroaniline (PCA), chlorothaldimethyl (CDM), dichlobenil (DCB), phthalide (PTL), quintozene (QTZ) were got from Shanghai Aladdin Bio-chem Technology Co., LTD. Hexachlorobenzene (HCB) and pentachlorophenol (PCP) were purchased from J&K Scientific Ltd.

### Measurements and apparatus

UV-vis absorption spectra were acquired on a TU-1901 spectrophotometer (China). The chromatographic analysis of CTL was performed by a 1260 HPLC system (Agilent) equipped with a VWD detector. The photographs were taken by a IXUS-190 camera (Canon, Japan).

### Colorimetric detection of CTL through enzymatic-triggered reaction

GAPDH (5 U) was preincubated at 37 °C in 50 mM phosphate buffer (pH 5.0) containing noted concentrations of CTL under stirring for 45 min. Following the preincubation, the reaction catalyzed by GAPDH with  $\text{NAD}^+$  as cofactor coupled with NBT-PMS was assay at room temperature in NaPpi buffer (pH 8.5) containing 0.5 mM GAP, 0.35 mM  $\text{NAD}^+$ , 120  $\mu\text{M}$  NBT, 75  $\mu\text{M}$  PMS in a total volume of 200  $\mu\text{L}$ . At that time, the color of the solution was recorded by photograph and determined by following the change in absorbance at 560 nm *via* UV-vis spectrophotometry.

### Pretreatment of cucumber samples

Cucumbers purchased from a market (Haidian, Beijing), were used for recovery examinations. After addition of 10 mL phosphoric acid solution (phosphoric acid/water = 1 : 1 v/v), the sample (100 g) were homogenized by a blender immediately. Then, each sample (100 g) was spiked with some known concentrations of CTL standards and allowed to stand at room temperature for 2 h.

For colorimetric determination, methanol (25 mL) was added to the samples (5 g) in a 50 mL tube and the tube was vigorously shaken for 30 min on an orbital shaker to extract CTL. After centrifuged at 3000 rpm for 5 min, the supernatant (1 mL) was downloaded into SPE cartridge. After washing the cartridge with 1 mL of methanol–water (1 : 9, v/v), the eluate was collected for colorimetric sensor. If needed, the extract solution was further diluted with methanol–water (9 : 1, v/v).

## Results and discussion

### Sensing mechanism of the colorimetric method

Fig. 1 illustrates the sensing mechanism for colorimetric sensor of CTL based on enzymatic reaction catalyzed by GAPDH coupled with PMS-NBT chromogenic system. This method relies on the fact that CTL is an inhibitor of specific nicotinamide adenine dinucleotide (NAD) thiol-dependent glycolytic enzyme-GAPDH,<sup>39,40</sup> leading to a decrease in production of NADH, which fails to induce a color change from yellow to blue of PMS-NBT chromogenic system. According to previous reports, the native GAPDH can catalyze the oxidation of GAP<sup>38</sup> into gluconolactone-6-phosphate (1,3-DPG), accompanied with reduction of  $\text{NAD}^+$  into NADH.<sup>38,41</sup> The simultaneously generated NADH participates in a cycling reaction that exhausts NBT yielding a purple coloured formazan ( $\lambda_{\text{max}}$ : 560 nm) with PMS as an electron carrier.<sup>42–45</sup> The absorbance intensity is proportional to the concentration of NADH. Whereas, previous investigations<sup>39</sup> have confirmed that inhibition of GAPDH by CTL was *via* forming a co-valently bond with amino acid residue Cys 149 in the active pocket (Fig. S1†) that required for NAD binding. The inhibition of activity of GAPDH by CTL leads to quantitative color change depending on the amount of CTL, which can be easily distinguished visually or quantified by the spectrometric measurement. With the absence of CTL, the solution of enzyme-

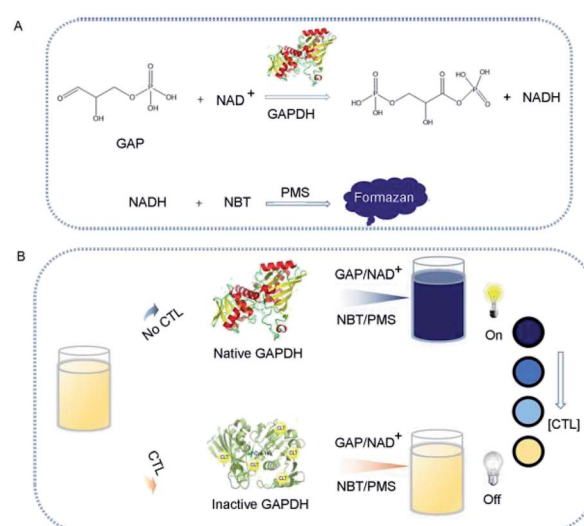


Fig. 1 (A) GAPDH catalyzes the oxidative phosphorylation of GAP to form 1,3-DPG with concomitant reduction of  $\text{NAD}^+$ . With the existence of NADH, NBT and PMS turn into blue formazan. (B) Scheme presentation of colorimetric detection of CTL *via* enzymatic action coupled with PMS/NBT strategy.



triggered reaction of GAP-NBT-PMS system rapidly proceeds and turns blue, and the system is “on”. However, when CTL is pre-mixed with GAPDH, the enzyme-triggered reaction is suppressed and the solution remain yellow, and the system is “off”. Thus, we devise a colorimetric method based on the catalytic capability of GAPDH coupled with PMS-NBT chromogenic system for detection of CTL. This colorimetric sensor is free from utilization of antibody or modification process, and the results can be obtained by either the naked eye or the spectrometer.

### Optimization of experimental conditions

To achieve a highly sensitive response, it's necessary to optimize the experimental conditions. pH is a key factor which can greatly affect the enzyme activity and the chromogenic reaction, thus pH of the reaction system was optimized firstly. As shown in Fig. 2A and S3A,†  $A_{560}$  of the generated formazan products sharply increased with the increasing value of pH and reached to a peak value at about pH 8.5. However, when pH was above 8.5, the absorbance at 560 nm dropped gradually with the increase of pH values, indicating the activity of enzyme was decreased. This was probably because the active-site residues within the catalytic domain center needed appropriate pH value to maintain its protonation state for product formation, and higher pH values led to deprotonation of active-site residues, resulting in the loss of enzyme activity. Therefore, pH 8.5 was selected as the optimal pH value.

GAP is catalyzed to form 1,3-diphosphoglycerate by GAPDH, accompanied by formation of NADH with  $\text{NAD}^+$  as an electron-transferring coenzyme. Therefore, concentrations of  $\text{NAD}^+$  and

GAP would influence the catalytic activity of GAPDH. The results showed that the maximum response was obtained when the concentration of GAP was 0.5 mM and the response slightly decreased with the increase of concentration when concentration of GAP was higher than 0.5 mM (Fig. 2B and S3B†). Similarly, the most remarkable color change was observed when the concentration of  $\text{NAD}^+$  was 0.35 mM, and the color change was moderately decreased when the concentration was above 0.35 mM (Fig. 2C and S3C†). The above results were in accordance with the theory of enzyme kinetics that there is an optimal concentration of the substrates and the reaction is inhibited when the substrate concentration is too high. Thus, 0.5 mM GAP and 0.35 mM  $\text{NAD}^+$  were selected as the optimal conditions for further investigations.

As reactants in the second step, the concentrations of NBT and PMS were closely connected with the generation of blue products, thus, their concentrations were further optimized. As depicted in Fig. 2D, E, S3D and S3E,† the absorbance at 560 nm reached to a plateau when the NBT concentration was 100  $\mu\text{M}$ . In the concentration range of 0–75  $\mu\text{M}$ , the color change was enhanced with the increase of concentration of PMS, and the change was declined when PMS was above than 75  $\mu\text{M}$ . As a result, concentrations of NBT and PMS were selected 100 and 75  $\mu\text{M}$ , respectively.

Finally, the incubation time of CTL with GAPDH was investigated in the same way as the above. Compared with control,  $A_{560}$  increased with the incubation time, and it reached to a minimum value at 45 min, but it remained unchanged with a further increase of the time (Fig. 2F). In consideration of the requirement of rapid determination, we selected 45 min for further investigations.

### Sensitivity of the colorimetric method

To examine the sensitivity of the enzymatic inhibition strategy-based colorimetric assay for the determining CTL, a series of different concentration of CTL standard solution were added to the GAPDH/GAP-NBT/PMS chromogenic system under the optimized condition. As shown in Fig. 3A, the absorbance band at 560 nm ( $A_{560}$ ) decreased progressively with the concentration of CTL ranging from 0 to 25  $\mu\text{M}$ . Meanwhile, an obvious color change from blue to yellow was also be found (the inset portion of Fig. 3A) and a sensitivity of about 1  $\mu\text{M}$  CTL was achieved with naked eyes. In addition, the logarithm of  $A_{560}$  ( $\log(A_{560})$ ) decreased correspondingly over the CTL concentration. With concentration ranging from 0.5 to 10  $\mu\text{M}$ , the  $\log(A_{560})$  displayed a linear correlation with  $R^2 = 0.993$  (Fig. 3B). The limit to detection (LOD) was estimated to be 0.05  $\mu\text{M}$  ( $3\sigma$ ). The high sensitivity and relative broad linear range is attributed to the high catalytic property of GAPDH.

As summarized in Table 1, we draw a comparison between the proposed colorimetric detection and other previously reported instrumental chromatographic analysis or antibody-based immunoassay. Compared to instrumental chromatographic analysis, limit of detection for CTL in this work was much higher than that of GC-MS or LC-MS. Also, the sensitivity of the proposed colorimetric assay was not remarkably

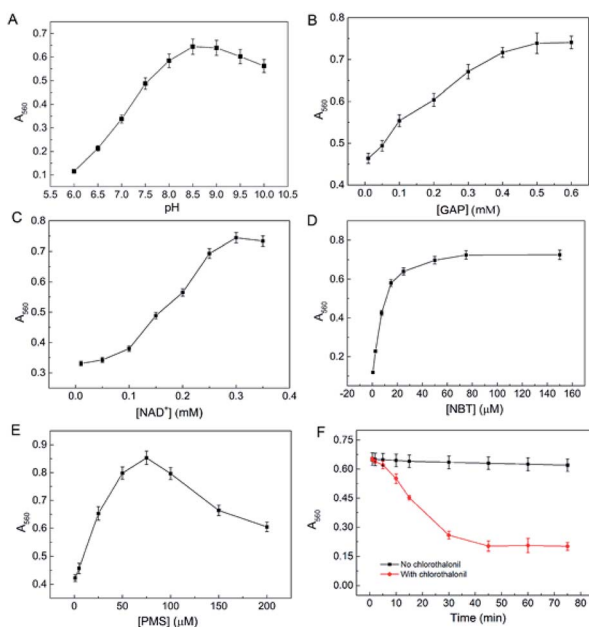


Fig. 2 Optimization of experimental conditions. Absorbance at 560 nm ( $A_{560}$ ) obtained by spectrometer with different pH values (A), various concentrations substrates (GAP (B),  $\text{NAD}^+$  (C), NBT (D) and PMS (E)), and varied incubation time intervals (F) in the presence of 5  $\mu\text{M}$  CTL.



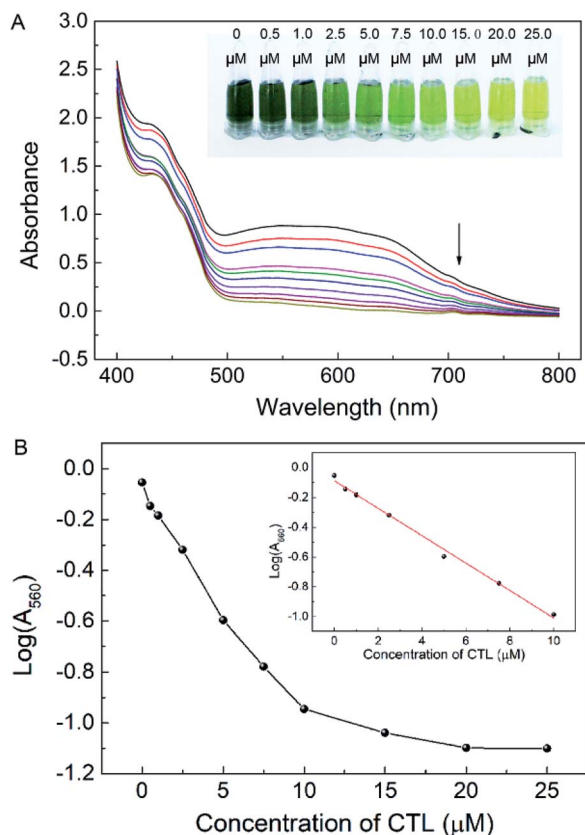


Fig. 3 (A) UV-vis spectra of the GAPDH/GAP-NBT/PMS chromogenic system in the presence of different concentrations of CTL. Inset portion was the corresponding photographic images. (B)  $\log(A_{560})$  of the reaction system versus the concentration of CTL. Inset portion showed the linear dependence of  $A_{560}$  at low CTL concentration (error bars represented the standard deviation of three measurements).

more effective than ELISA. Since the MRLs of CTL in vegetables was set from  $5 \text{ mg kg}^{-1}$  in China ( $5 \text{ mg kg}^{-1}$  in USA and  $1 \text{ mg kg}^{-1}$  in Europe), the sensitivity of the proposed analysis was effective enough to detect CTL residues in vegetables. Moreover, the approach avoided the use of expensive and complex instruments apparatus required in instrumental analysis and utilization of expensive antibody of the immunoassay, which undoubtedly improves their application in real samples.

### Selectivity of the colorimetric method

To determine the selectivity of the colorimetric assay for CTL, CTL ( $5 \text{ } \mu\text{M}$ ) and several analogues of closely related molecular structure at the level of  $50 \text{ } \mu\text{M}$  were carried out to undergo the chromogenic reaction. As shown in Fig. 4, it was observed that only in the presence of CTL, the solutions turned yellow and showed a decrease in  $A_{560}$ . However, the other pesticides with the level 10 times higher than CTL stay blue, similarly to the blank sample, and color and  $A_{560}$  of the solution had no notable change, revealing that the inhibition of the reaction was not occurred. The other pesticides yielded only negligible response comparing to that of CTL. CTL inhibits the activity of GAPDH by covalent interaction with cysteine-149 located at the active center responsible for the binding of GAP. Due to difference in the molecular size and chemical nature of the reactive toxiphores, pesticides or chemical compounds with similar chemical structure do not inhibit the enzyme. It should be noted that certain trichloromethyl sulfonyl fungicides suppress activity of GAPDH. However, CTL differed from those fungicides in that it did not react with non-thiol groups of either GAPDH and had a different reaction rate with the GAPDH. This indicated that other pesticides didn't inhibit the catalytic activity of GAPDH and didn't interrupt the chromogenic system of GAPDH coupled with NBT/PMS. The colorimetric method has excellent specificity to CTL and the colorimetric system is suitable for selective detection of CTL.

### Application of cucumber samples

As a proof of its applicability, cucumber was chosen as a representative specie to perform the spiked test. Different concentrations of CTL standard solutions were added into the cucumber sample (final concentration  $50.0$ ,  $100.0$ ,  $150.0 \text{ } \mu\text{g kg}^{-1}$ , respectively). As listed in Table 2, the recovery values were  $82.0\%$ ,  $90.2\%$  and  $84.9\%$ , respectively, which was consistent with that obtained by HPLC (Table S1†). Therefore, this facile, antibody-free assay can be used for sample with simple matrixes directly with satisfactory results. However, better sample clean-up procedures are necessary to eliminate interference for sample with more complicated matrixes such as vegetables and fruits. The reason is that substances such as pigment or sulphur-containing compounds in the real samples can interfere the chromogenic reaction, which has distinct impact on the quantification of analyte. We will make efforts to apply the

Table 1 Comparison of different detection method of CTL

Methods	Linear range ( $\mu\text{M}$ )	Limit of detection ( $\mu\text{M}$ )	Use of large instrument	Use of antibody	Ref.
GC-MS	0.19–37.6	0.19	Yes	No	5
HPLC-MS	0.001–1.12	0.004	Yes	No	6
Immunosensor	0.03–0.16	0.09	No	Yes	7
ELISA	0.0007–0.005	0.0002	No	Yes	8
IC-ELISA	0.0003–0.004	0.001	No	Yes	9
IC-ELISA	0.004–0.022	0.001	No	Yes	10
Colloidal gold-based strip test	0.005–0.01	0.35	No	Yes	11
Colorimetric assay	0.5–10	0.05	No	No	This study



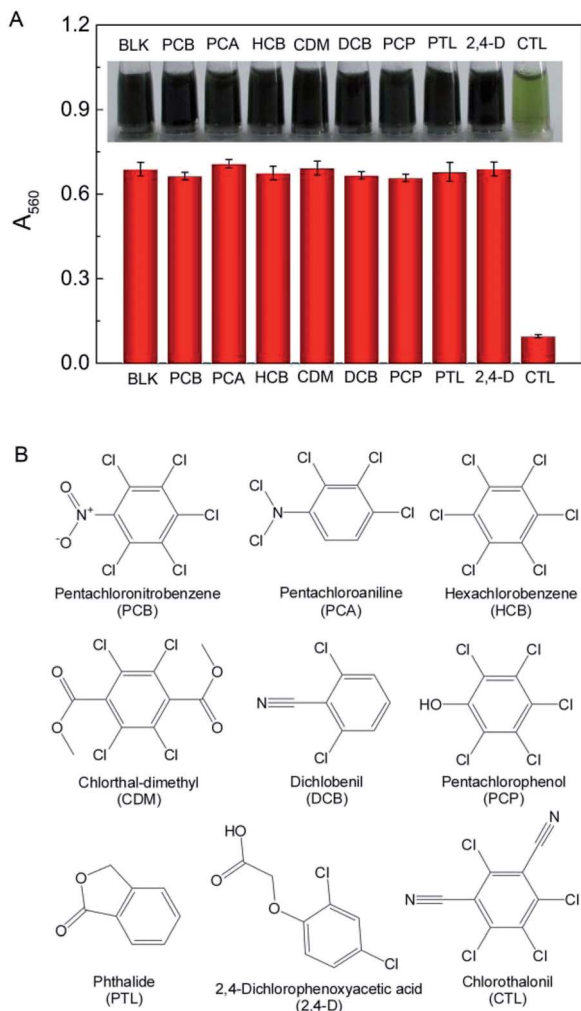


Fig. 4 (A)  $A_{560}$  of mixture spiked with CTL and other compounds (5  $\mu$ m). Inset portion showed the corresponding picture. (B) Chemical structures of the selected compounds.

Table 2 The recovery under different concentration of CTL in cucumber samples

Sample	Added ( $\mu$ g kg <sup>-1</sup> )	Visual assay		
		Found ( $\mu$ g kg <sup>-1</sup> )	Recovery (%)	RSD (%)
Cucumber 1	50.0	41.0 $\pm$ 1.6	82.0	3.9
Cucumber 2	100.0	90.2 $\pm$ 4.3	90.2	4.8
Cucumber 3	150.0	127.4 $\pm$ 6.9	84.9	5.4

colorimetric procedures to determine CTL in other agro-products in future work.

## Conclusions

In summary, we developed a facile, sensitive colorimetric biosensor for CTL detection with straightforward readout and implementation. This strategy was relied on the inhibition of GAPDH activity by CTL, resulting in color change of NBT/PMS

system, which could be visualized by naked eyes or quantification *via* the UV-vis spectrophotometer. The linear range of the method was from 0.5 to 10  $\mu$ M, and 1  $\mu$ M CTL could be distinguished with naked eyes. Compared with the existing techniques for CTL detection, the established method was large instrument-free and antibody-free. In addition, we successfully applied the method to the determination of CTL in cucumber. Therefore, it has great potential application as a selective and sensitive platform for detection of CTL in other agro-products.

## Conflicts of interest

There are no conflicts to declare.

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