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Colorimetric sensing of calcium carbide over banana peels using 5,5'-dithiobis-(2-nitrobenzoic acid) (DTNB) as a rapid chemoreceptor: a point of care tool for food fraud analysis†

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Point-of-care sensing systems have led to the rapid development of smart and portable devices in the field of analyte detection technology. The high flow of artificially ripened fruits (ARFs) in the market results in severe health risks, and simultaneously, the unavailability of consumer-level 1st screening tool flares up its exigency. Calcium carbide (CaC₂) is a hazardous artificial fruit ripening agent. In this study, a portable one-step colorimetric sensor was developed for the detection of CaC₂ on the surface of ARFs. The presence of CaC₂ was detected using 5,5'-dithiobis-(2-nitrobenzoic acid). The colorimetric reaction of DTNB detected CaC₂ by the free sulfhydryl group detection method which produced a yellow color; the free sulfhydryl group is present as an impurity in CaC₂. The proposed assay displayed an LOD of 50 ppm with a detection time of a few seconds to 1 minute. It exhibited a rapid, highly sensitive and highly selective response, with a linear dynamic range of 0–4000 ppm and R² values of 0.993 and 0.994 for 323 and 412 nm peaks, respectively. The development of this inexpensive, naked eye, portable, and easily accessible CaC₂ sensor reinforces the suggestion that it could bridge the gap between conventional detection techniques and consumer-level detection methods. Our sensor could be useful as a 1st screening tool in various fields, including agriculture, food, and cosmetics industries.

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1. Introduction

Calcium carbide (CaC₂) is a gray carcinogenic synthetic compound manufactured to fulfill the rapidly increasing global demand of the developing sectors, such as iron desulfurization (sulfur removal from iron, including cast iron, pig iron, and steel) and artificial ripening of pre-harvested fruits.^{1,2} The end uses of CaC₂ in different regions globally are expected to exceed US\$ 31 billion by 2031.³ Extended application and global

utilization of CaC₂ are strong evidence of its widespread availability in the market at a low or no cost. There is a significant amount of “CaS” (sulfides) present in CaC₂ organically, as well as in the CaC₂ coming from steel industries after the desulfurization process.^{4–6} CaC₂ is widely used commercially and at home to accelerate the artificial ripening of fruits and crops.^{7,8} India is leading the world in banana, papaya, and mango production. Unfortunately, traders illegally use CaC₂ to artificially ripen these fruits, which violates the Food Safety and Standards Act of 2006 regulations^{9,10} under the Food Safety and Standards Authority of India (FSSAI).¹¹ FSSAI has prohibited the use of CaC₂ due to its serious health risks under the provisions of sub-regulation 2.3.5 of the Food Safety and Standards (Prohibition and Restriction on Sales) Regulations, 2011.^{12–15} In the current scenario, rampant fruit adulteration (artificial ripening of fruits) has been seized, and there is massive loss and waste of agricultural food, particularly in India. The FSSAI and the Food and Drug Administration (FDA) in India are on the lookout for cases of artificially ripened mangoes and have uncovered numerous cases of food fraud,¹⁶ including the Guwahati, India case,¹⁷ Ukkadam city, Coimbatore, India case,¹⁸ and Gudivada and Machilipatnam, India case.¹⁹

CaC₂ produces heat when it comes into contact with moisture, which can cause burns and damage to the mucosa, skin,

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and eyes. Artificial fruit ripening agents (AFRAs) are carcinogenic and highly toxic, and their consumption can cause serious health problems, such as heart disease, as well as lung and kidney failure, in humans and animals.²⁰ Regular consumption of AFRAs can cause dizziness, weakness, skin ulcer, diarrhea, cancer, and central nervous system depression.^{21,22} CaC₂ is particularly used to ripen pre-harvest fruits during storage and transit, as ripened fruits cannot be transported due to their short shelf life.^{23,24} Therefore, fruit quality examination has become necessary to selectively monitor CaC₂ using efficient methodologies. At the consumer end, CaC₂ can be used as a biomarker to monitor fruit quality concerning food safety, allowing consumers to avoid eating CaC₂-ripened fruits for a healthier lifestyle. Thus, detecting the presence of CaC₂ is essential because it is directly linked to health risks. However, there is no simple detection technique available for on-spot monitoring of CaC₂ over fruit peels as a first-screening method, particularly for fruits like mango and banana that are over-ripened with CaC₂ to deceive customers.²⁵ To prevent such life-threatening practices and diseases, we have developed the proposed sensor, which will solve this problem by being the first-screening device for monitoring artificially ripened fruits.

Complex and time-consuming traditional methods,^{26–28} such as liquid chromatography, liquid chromatography–mass spectrometry, data processing techniques, optical techniques, photoacoustic spectroscopy, and electrochemical methods, are currently used for CaC₂ measurement.^{21,29} Although these techniques can accurately quantify CaC₂ at low concentrations, they are not relevant to the consumers in the field.^{30–32} To protect the worldwide environment and human health, analytes (dangerous substances) must be monitored promptly, effectively, and on-site. Colorimetric sensing techniques enable the visual detection of particular chemical substances in real time. Due to its simplicity, reasonable cost, and practicality, as well as the benefit of functioning as a non-label detection method, this methodology has been widely used in a variety of sectors. As a result, the invention of miniaturized analytical procedures with good sensitivity and selectivity for a variety of chemical species (such as biomolecules, organic, and inorganic compounds) has been based on colorimetric detection. Importantly, colorimetric procedures often do not call for specialised equipment or the knowledge of a professional technician.^{33–37} Continuous attempts are being made to make the CaC₂ detection method more ASSURED (affordable, sensitive, specific, user-friendly, rapid, equipment-free, and deliverable) but, due to the restricted chemical functionality, only one colorimetric sensor has been developed so far for CaC₂ detection.^{38,39} However, the development of colorimetric sensors is based on indirect sensing and the reduction of gold nanoparticles (AuNPs). The color of lauryl sulfate (LS)-capped AuNP aggregates changes from red to purple in the presence of arsenic while LS is replaced.⁴⁰ This is a time-consuming assay which demands expertise for analysis. Despite these advancements, incorporating CaC₂ sensors that contain stiff components, such as metals and nanoparticles, and sensitive components, such as enzymes and antibodies, into flexible and polymer-based detection kits remains a technical challenge.^{41–44}

In general, sensors for direct food quality monitoring must be suitable (ASSURED) for consumers and manufacturing processing units.^{45–48}

This study aimed to use Ellman's reagent [5,5'-dithiobis-(2-nitrobenzoic acid) (DTNB)] to create simple CaC₂ sensors that use a direct sensing method. The DTNB assay is used to assess enzyme activity; it detects sulfide bonds and targets them for cleavage, resulting in ions that are responsible for colorimetric signals.^{49,50} To monitor the presence of CaC₂ in banana peels, we focused on the high sulfide impurity "CaS" present in commercial CaC₂,^{4–6} and a new method was validated, and a solution-based easy colorimetric CaC₂ sensor was developed using the DTNB-based method as 1st screening for artificially ripened bananas. As a proof of concept for real-time detection, bananas were used as the model sample in this study. We developed a simple one-step, sensitive, selective, and prompt colorimetric sensor for CaC₂ detection based on this concept. Since such types of detection kits are commercially unavailable (as the Ministry of Agriculture claims), the presence of CaC₂ can only be tested in laboratories using suitable equipment and additional chemicals. Thus, our sensor will provide a platform that will aid in the resolution of these issues.²⁵ The proposed sensor has the potential to effectively combat artificial ripening practices and agricultural food loss. When consumers have direct access to this type of portable sensor, they will be able to conduct their first-screening test to ensure that they are purchasing good-quality fruits. Accordingly, the sale of artificially ripened fruits and the practice of artificial ripening will become extinct. The study of free sulfhydryl groups is a rapidly developing area in both basic and applied biology. The quantitative measurement of free sulfhydryl groups is a common task in many practical areas where a quick and simple procedure is greatly preferred. Although conventional and traditional assays require time-consuming steps they are exceedingly sensitive and accurate.^{28,47,48} Ellman's spectrophotometric free sulfhydryl group test is quick and easy to use, making it more common for quantifying sulfhydryl. It will also fill the gap between numerous unaware frauds and consumers in various sectors such as medical, industrial, cosmetics and many more fields.

2. Experimental procedure

2.1. Materials and methods

Ellman's reagent (DTNB), sodium phosphate dibasic (Na₂HPO₄·7H₂O), sodium phosphate monobasic monohydrate (NaH₂PO₄·H₂O), CaC₂, ethephon, potassium sulfate, potassium dihydrogen orthophosphate, oxytocin, ethylene glycol, ammonium sulfate, manganese sulfate, magnesium sulfate, potassium sulfate, sodium sulfate, and copper sulfate were purchased from Sigma-Aldrich and directly utilized as received without any further purification. Deionized water, Whatman filter paper, cotton swap, raw bananas, glass beads, and plastic bottle were obtained commercially. Ultraviolet (UV) spectrophotometry, X-ray fluorescence (XRF), scanning electron microscopy (SEM), and X-ray photoelectron spectroscopy (XPS) were used for characterization



in this study. Inductively coupled plasma-optical emission spectroscopy (ICP-OES) was used for the validation of the fabricated sensor.

2.2. DTNB preparation

$\text{Na}_2\text{HPO}_4 \cdot 7\text{H}_2\text{O}$ (10.107 g) and $\text{NaH}_2\text{PO}_4 \cdot \text{H}_2\text{O}$ (1.697 g) were dissolved in 500 mL of distilled water to prepare a 0.1 M sodium phosphate buffer, and the pH was further adjusted to 7.4, 6, 7, 8, and 9. DTNB solutions of various concentrations (0.5 mM, 1 mM, 10 mM, and 15 mM) were prepared using the 0.1 M sodium phosphate buffer. For optimization, different DTNB concentrations were prepared using different pH values of sodium phosphate buffer.

2.3. CaC_2 solution preparation

CaC_2 powder was dissolved in DW, and a 10 000 ppm CaC_2 stock solution was prepared. Using the stock solution, 0, 10, 50, 100, 150, 200, 250, 500, 750, 1000, 2000, 4000, 6000, and 10 000 ppm solutions were prepared and used for the colorimetric assay.

2.4. CaC_2 sensing procedure

CaC_2 solution (from the stock solution) and DTNB solution were mixed to obtain a colorimetric signal for solution-based colorimetric CaC_2 detection and Ellman's reagent reduction. Four different combinations of CaC_2 and DTNB solutions were used for this experiment: (a) 15 mM DTNB (pH 9; 150 μL) and CaC_2 (850 μL), (b) 15 mM DTNB (pH 7; 150 μL) and CaC_2 (850 μL), (c) 0.5 mM DTNB (pH 9; 150 μL) and CaC_2 (850 μL), and (d) 0.5 mM DTNB (pH 7; 150 μL) and CaC_2 (850 μL) [Fig. S1, ESI[†]]. Colorimetric sensing was then conducted using the optimized combination (a).

2.5. Color signal characterization

Sensor color change was assessed using digital images of the films before and after exposure to the CaC_2 solution. The images of the enzymatic chemoreceptor-based DTNB sensors were captured using a digital camera. The absorption spectra of all CaC_2 concentrations were recorded using a UV spectrophotometer. Raw bananas were used as the sample model for real-time detection in this study. Our sensor can be used for a variety of other fruits, including those that are subjected to artificial ripening, such as mango, pear, apple, papaya, grapes, and plum. Raw bananas were artificially ripened by applying (spraying)⁵¹ 10,000 ppm (minimum concentration for ripening)²⁴ CaC_2 solution on them and storing them in a closed chamber in the dark at room temperature for two days. After two days, the bananas were taken out and examined for the presence of CaC_2 . CaC_2 was detected by swabbing a wet (water) cotton plug over a banana peel and allowing the analyte to absorb into the cotton plug. After swabbing all the exposed areas, the cotton plug was squeezed into a vial containing the DTNB solution for the colorimetric detection of CaC_2 in a real sample.

2.6. XRF, SEM, EDX, and XPS characterization

X-ray fluorescence (XRF) analysis was used to study the entire CaC_2 composition [Table S1, ESI[†]]. Scanning electron microscopy (SEM)

images of CaC_2 were captured and analyzed to study its morphology (microstructure ameliorates) [Fig. 1(A)]. SEM mapping and X-ray photoelectron spectroscopy (XPS) analysis were used to examine the distribution of elements at the core level and fitted XPS data for carbon, calcium, and sulfur distribution [Fig. 1(B)–(K)–(1–3)]. Data of XPS analysis for the distribution of remaining elements at the core level are mentioned in Fig. S2 (ESI[†]).

2.7. Matrix effect and sulfate test

The banana matrix also contained natural sulfate. We used the proposed sensing technique to cross-check and confirm the selectivity and performance of the developed sensor. An extraction technique was used to prepare the banana matrix sample. By combining 1 g of banana, 3 mL of phosphate buffer, and 5 g of glass beads in a plastic bottle and vigorously shaking it with a vortex, banana pulp was made. For matrix analysis, the pulp extract was separated from the glass beads and combined with the DTNB solution (sulfate test) [Fig. S3, ESI[†]]. Other sulfates (ammonium sulfate, manganese sulfate, magnesium sulfate, potassium sulfate, sodium sulfate, and copper sulfate) were also tested to determine the sensitivity and selectivity of the developed sensor [Fig. S5, ESI[†]].

3. Results and discussion

3.1. Compositional and structural study of CaC_2

The morphology and chemical composition of commercial CaC_2 were examined by the HR-SEM and energy-dispersive X-ray (EDX) elemental mapping studies. CaC_2 is seen to be a large irregular crystal in the SEM image. The surface elemental composition of the crystals was analyzed by elemental mapping studies which evidently showed the presence of calcium (Ca), sulphur (S), silicon (Si), phosphorous (P), magnesium (Mg), aluminum (Al), and iron (Fe) [Fig. 1(A)–(I)]. The trace elements detected like S, Si, P, Mg, Al, Fe, *etc.* are impurities present in the commercial calcium carbide available in the market. XRF data [Table S1, ESI[†]] show the high sulfide content in calcium carbide powder. The molecular states and structure of the compound were elucidated from XPS characterization. In Fig. 1(J)–(1–3), the core level C1s, Ca2p, and S2p spectra of CaC_2 evidently established the presence of C, Ca, and S, respectively. The fitted peaks of the same are shown in Fig. 1(K)–(1–3). The C1s spectrum exhibits three prominent peaks, respectively, at 284.4 eV which corresponds to the $\text{sp}^3\text{-C}/\text{sp}^2\text{-C}$ hybridized carbon, at 286.4 eV which corresponds to the sp-C carbon, and finally, at 290 eV representing C–O–C bonding.^{52,53} The Ca2p core level spectra can be resolved into three components. The peak corresponds to the binding energy 346.3 eV, representing $\text{Ca}2\text{p}_{3/2}$ and another prominent peak at 350.5 eV corresponds to $\text{Ca}2\text{p}_{1/2}$.⁵⁴ The presence of CaC_2 species was further confirmed by the peak at 348 eV S2p core level spectrum which exhibited two prominent peaks at 163.9 eV and 162.5 eV, respectively, representing metallic sulphur and CaS bonding present in the commercial calcium carbide as an impurity.⁵⁵



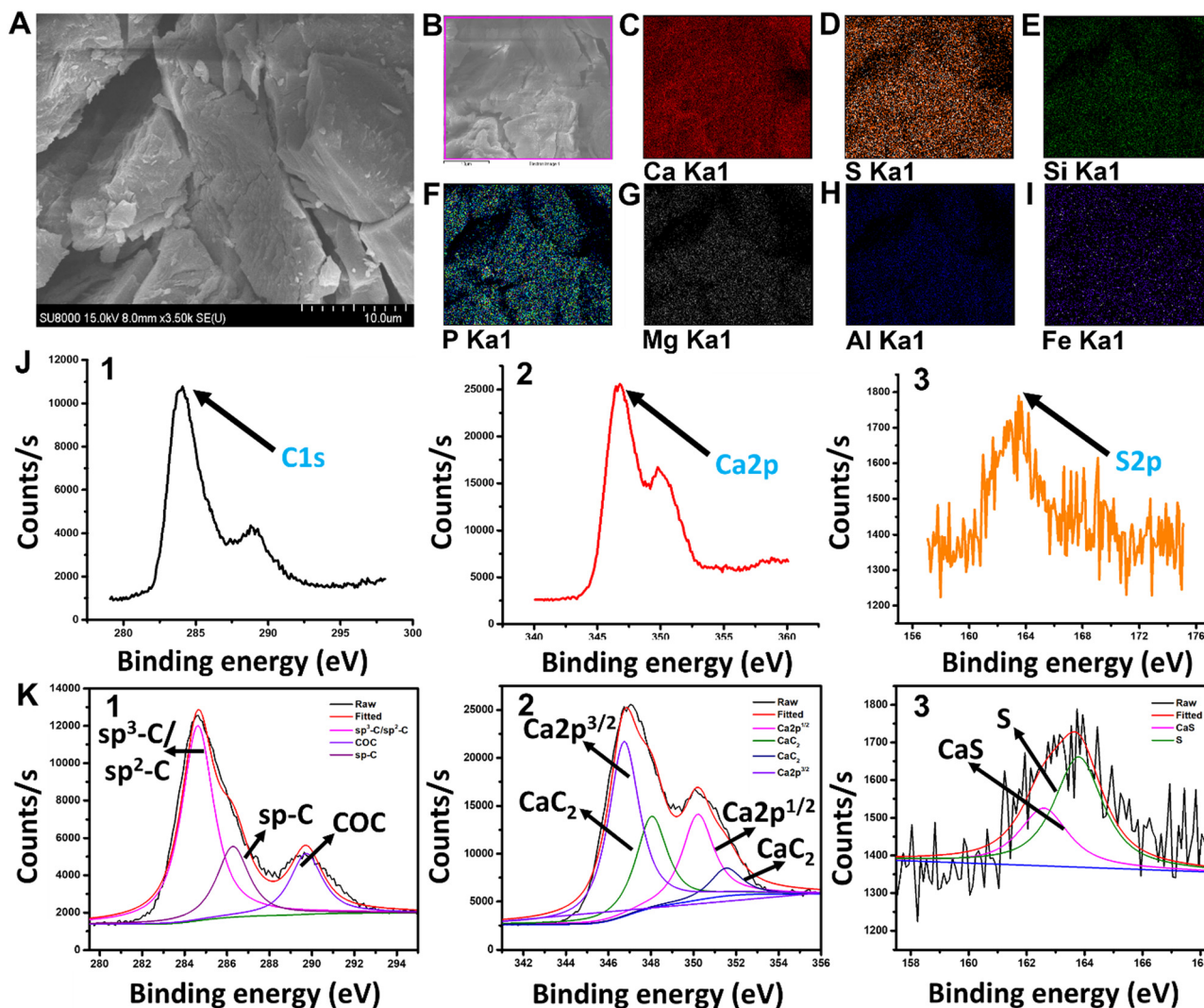


Fig. 1 (A) SEM image of CaC₂ at 10 μm. (B)–(I) Distribution of all elements and their core-level studies based on SEM mapping. (J1–J3) XPS data. (K1–K3) Fitted XPS data for carbon, calcium, and sulfur distribution.

3.2 DTNB solution as a CaC₂ sensor

DTNB is a chemical substance that is used to determine the concentration of free sulfhydryl groups in a sample. George L. Ellman came up with the concept and created this reagent by oxidizing 2-nitro-5-chlorobenzaldehyde to carboxylic acid, adding free sulfhydryl groups *via* sodium sulfide, and linking the monomer *via* iodine oxidation. CaS[−] free sulfhydryl groups of CaC₂ interact with this reagent, slicing the disulfide bond to form the TNB[−] anion,⁵⁶ which ionizes to TNB^{2−} dianion in water at neutral and alkaline pH; TNB^{2−} ions have a yellow color. One mole of free sulfhydryl groups releases one mole of TNB in this rapid and stoichiometric reaction. TNB^{2−} was detected spectrophotometrically by measuring the visible light absorbance at 412 nm with an extinction coefficient of 14,150 M^{−1} cm^{−1} for dilute buffer solutions at pH 9.0 [Fig. 2(A)]. The DTNB buffer solution changed from colorless to yellow and then dark yellow after being exposed to CaC₂ [Fig. 2(E)].⁵⁷ CaC₂ is detected using DTNB because of its impurity form “CaS” contains a significant

amount of sulfide. This contaminant is produced during steel desulfurization in steel plants.^{4–6}

When DTNB reacts with a free sulfhydryl group, mixed disulfide and TNB acid are formed [Fig. 2(F)]. DTNB targets the conjugate base (R-S[−]) “CaS[−]” of CaC₂, the free sulfhydryl group in this reaction, producing TNB, a “colored” species with a high molar extinction coefficient in the visible region [Fig. 2(F)].^{58,59} In the case of CaC₂, the CaS[−] sulfide moiety (present as an impurity source from steel manufacturing industries) interacts with DTNB *via* the above stated mechanism. The standard linear curve fitting can be a good indicator of the assay's strength at 323 and 412 nm peaks [Fig. 2(C) and (D)]. To produce TNB and mixed disulfides, free sulfide groups interact stoichiometrically with TNB. Further at pH 9, TNB ionizes to TNB^{2−} dianion in water. The color of TNB^{2−} ions is yellow. Depending on the CaS concentration in CaC₂, the color of the DTNB buffer solution changed from colorless to yellow and eventually to dark yellow after being exposed to CaC₂ [Fig. 2(F)]. As shown in Fig. 2(C), DTNB steadily reduces to TNB^{2−} ions at low CaC₂ concentrations because there are less



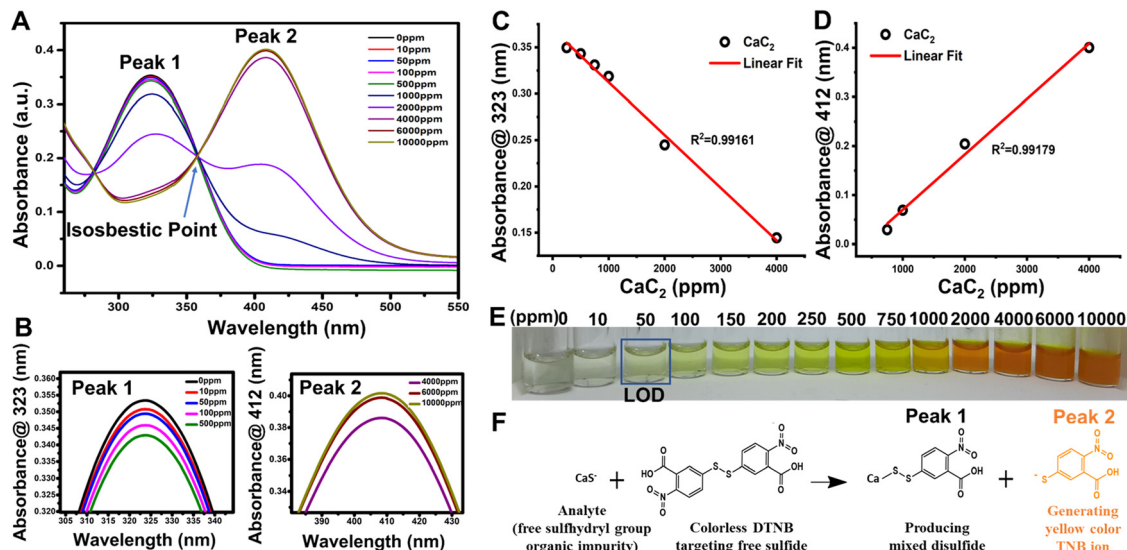


Fig. 2 (A) CaC_2 (0–10 000 ppm) detection absorbance spectra with a redshift in peaks from 323 to 412 nm, indicating the formation of TNB ions, which are responsible for the yellow color development. (B) Magnified representation of plot (A) for a clear understanding of absorption values at 323 and 412 nm peaks. (C) and (D) Linear curve fittings of the peaks at 323 and 412 nm, respectively, for standard CaC_2 concentrations. (E) Solution based CaC_2 colorimetric sensing, and (F) reduction of DTNB ions responsible for the redshift and color development.

sulfide ions available to interact with it. DTNB exhibits a spectroscopic absorbance peak at 323 nm, whereas TNB^{2-} exhibits a peak at 412 nm. All the DTNB undergoes a rapid stoichiometric reaction in the presence of high sulfide content and produces yellow color [Fig. 2(E)], as seen by the shift in the absorbance spectra of TNB^{2-} , which showed absorbance at a wavelength of 412 nm, instead of 323 nm as DTNB quickly converted to TNB^{2-} ions [Fig. 2(D)]. Ellman's assay has an isosbestic point of around 356 nm [Fig. 2(A)]. The isosbestic point can be determined either tabularly or graphically by measuring the full spectrum of the absorbance values for each solution to confirm that the molar ratio between Ellman's reagent and the test sample is equivalent across each test solution. A smooth peak at 412 nm also indicates that our solution is within the assay's working range. Fig. 2(C) and (D) depict the linear curve fittings of peaks at 323 and 412 nm, respectively, for standard CaC_2 concentrations. Ellman's reagent can also be used to measure free sulfhydryl groups such as glutathione in pure solutions⁶⁰ as well as biological specimens such as blood.⁶¹ It can also be used to determine the number of free sulfhydryl groups in proteins.⁶² Despite the widespread use of Ellman's reagent in various sensing applications, no study has been published on using Ellman reagent-based sensors to detect CaC_2 on fruit skin after CaC_2 artificial ripening.

3.3. Quantification of sulfhydryl concentrations in an unknown sample

The absorbance spectra of unknown solutions can be compared to those of the standard curve to ensure that the concentrations of the solutions are within the standard curve's range [Fig. 3]. The absorbance values of unknown CaC_2 samples 1, 2, 3, and 4 were plotted along the standard curve [Fig. 3(C) and (D)], and they corresponded to those of the standard curve.

The linear trend of the sensor for the unknown sample can be used to determine the amount of CaC_2 present in banana peels [Fig. 3(C) and (D)]. The number of sulfhydryl groups in the unknown sample was quantified based on molar absorptivity. An 850 μL aliquot of four different unknown samples mixed with 150 μL of Ellman's reagent solution produced absorbances of (1) 0.294, (2) 0.463, (3) 0.326, and (4) 0.201 on using a 1 cm spectrophotometric cuvette, which was plotted linearly [Fig. 3(C) and (D)]. The sulfhydryl concentrations in ppm were calculated for the four different unknown samples a, b, c, and d, and the results corresponded to 5.72 ppm, 7.53 ppm, 3.6 ppm, and 4.73 ppm, respectively [Table S2, ESI[†]].

3.4. Selectivity and sensitivity of the DTNB solution as a CaC_2 sensor

The visual observation of color changes in association with UV absorbance spectra [Fig. 2(D)] supported the colorimetric change of DTNB to TNB^{2-} ions in the system in the presence of CaC_2 . It was necessary to determine DTNB selectivity in response to other AFRAs. Thus, common ripening agents, such as ethephon, potassium sulfate, potassium dihydrogen orthophosphate, oxytocin, and ethylene glycol, were added to various vials containing the DTNB sensor solution and spectral analysis was carried out [Fig. 4(B)], and the results revealed that the DTNB sensor was only sensitive to CaC_2 , with its color changing ability from colorless to yellow [Fig. 4(A)]. Additionally, by doing the assay five more times, we were able to test the reproducibility and sulfide content in five distinct batches of the same 500 ppm CaC_2 concentration, as shown in [Fig. 4(C)]. Spectral analysis of all five batches confirmed that the sulfide content was similar. The response time of the DTNB sensor was measured by exposing the sensor to CaC_2 solution and taking images at 1 min intervals. The sensor quickly changed color



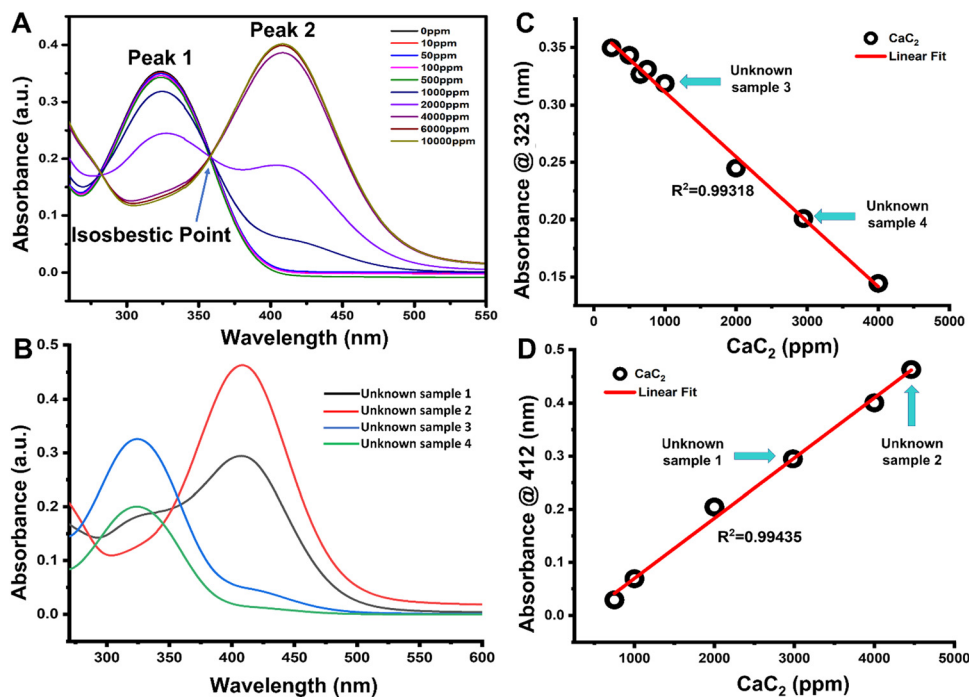


Fig. 3 The performance of the DTNB sensor in CaC_2 detection based on broad range concentration (0–10 000 ppm). (A) Absorbance spectra and ion formation, with a redshift in peaks from 323 to 412 nm for standard CaC_2 concentrations, indicating the formation of TNB ions responsible for the yellow color development. (B) Absorption spectra of unknown samples 1, 2, 3, and 4, and (C) and (D) fitting of unknown sample concentrations in standard linear curves at peaks at 323 and 412 nm, respectively.

from colorless to yellow, and no further significant color change was observed after 10 min of exposure (negligible change) [Fig. 4(D)]. With the help of spectral analysis, it was confirmed that after 10 min, the formation of TNB ions got

saturated and no change in peak intensity was observed at concentrations of 50 and 100 ppm CaC_2 [Fig. 4(E) and (F)]. In this study, time-dependent degradation of the colorimetric signal was performed, and it was discovered that the sensor's

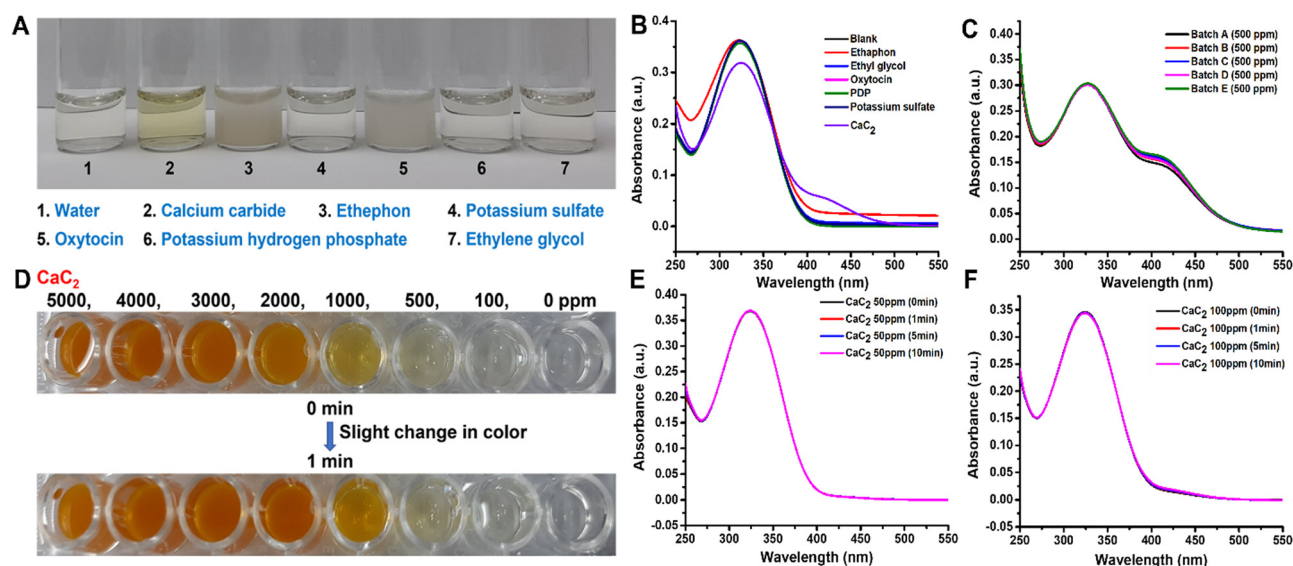


Fig. 4 DTNB sensor performance in the presence of other ripening agents as well as the time-dependent stability. (A) The DTNB sensor was exclusively sensitive to CaC_2 , exhibiting a color change from colorless to yellow, and was insensitive to other less common ripening agents (ethephon, potassium sulfate, potassium dihydrogen orthophosphate, oxytocin, and ethylene glycol) involved in the fruit-ripening process. (B) Spectral analysis of all interfering compounds. (C) Reproducibility and spectral analysis of CaC_2 in different batches. (D) The sensor changed color quickly, and no further significant color change was observed after 1 min of exposure (negligible change). (E and F) Stability and time-dependent spectral analysis of 50 and 100 ppm CaC_2 .



color remained stable for more than a month [Fig. S4, ESI†]. This rapid color change indicates that the free sulfide group from CaC_2 reacted with the DTNB sensor to form TNB ions. The detection limit was calculated as the average value of absorbance at zero concentration with three standard deviations.^{63,64} The DTNB sensors were exposed to different CaC_2 concentrations ranging from 0 to 10 000 ppm, and their images were taken immediately to determine their limit of detection (LOD) for CaC_2 [Fig. 2(E)]. When the CaC_2 concentration was less than 50 ppm, the color change was quite visible to the naked eye. However, at concentrations of 100 ppm and above, the color change of the sensors was clearly visible. The LOD of the DTNB sensors corresponded to that of CaC_2 .

3.5. Artificially ripened fruit monitoring

The developed DTNB-based sensor was used for real-time monitoring of CaC_2 in banana fruit peels as a model sample for our work in order to demonstrate its practical application for CaC_2 monitoring. Our sensor is applicable to other fruits as well, such as mango, pear, apple, papaya, grapes, plum, *etc.*, which are considered for the artificial ripening treatment. The color transitioning pattern of the banana-exposed sensor [Fig. 5] was found to be similar to that of the pure CaC_2 -exposed sensor.

3.6. Monitoring of the matrix effect and other sulfates

We used the proposed sensor to test the interference of the natural sulfate content and the additional sulfate contents in the banana matrix. It was discovered that there was no interference effect of matrix and other sulfates (ammonium sulfate, manganese sulfate, magnesium sulfate, potassium sulfate, sodium sulfate, and copper sulfate) [Fig. S2 and S5, ESI† respectively]. According to the one-step analysis, our proposed sensor demonstrated the best performance for CaC_2 detection with high sensitivity and selectivity immediately after being exposed to samples.

3.7. Assessment of sensor performance

Currently, colorimetric sensors are very attractive due to their inherent advantages of direct understanding of detection results, portability, low-cost efficiency, user-friendliness, and high sensitivity. Only one colorimetric assay has been published for CaC_2 detection; however, it does not meet the basic requirements as it is complicated and non-portable, and requires a trained person for handling. To combat such challenges, we proposed a simple sensor and the performance of our CaC_2 sensor is compared with other prominent work in Table 1. For all existing techniques complicated methods, difficulty in understanding at the consumer level, and non-

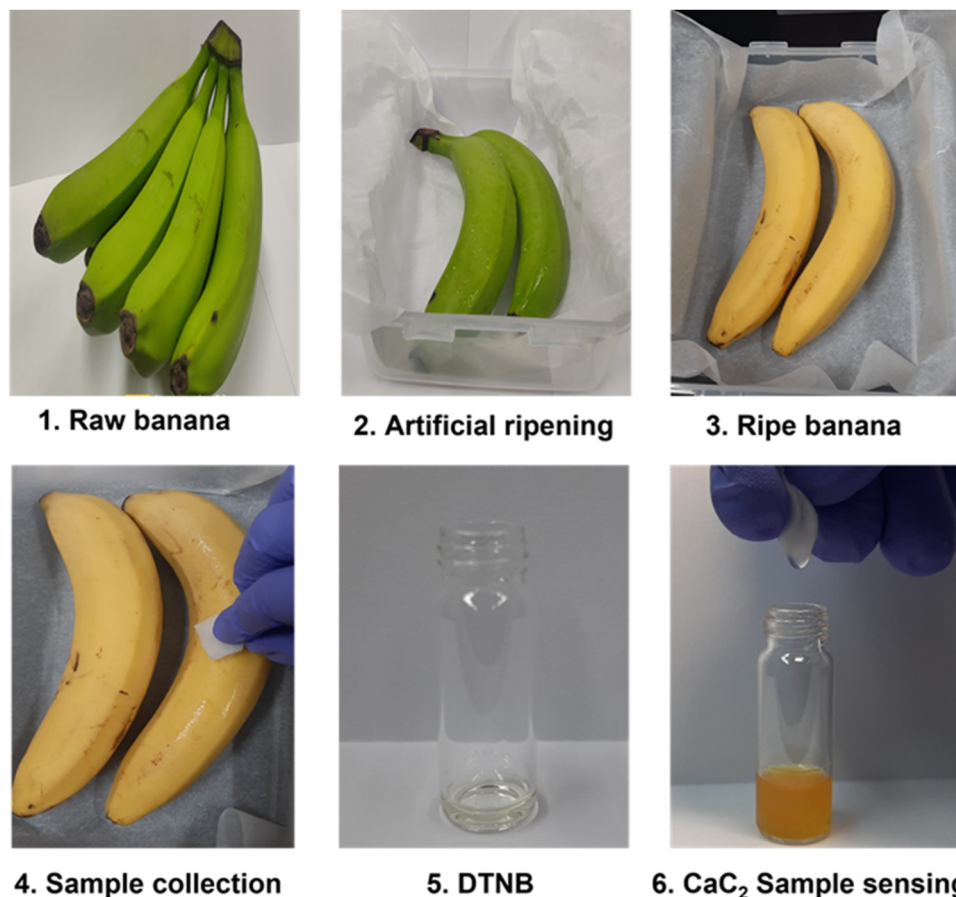


Fig. 5 (1–6) CaC_2 monitoring using the DTNB sensor on the skin of artificially ripened bananas, with a step-by-step illustration of real-time sensing.



Table 1 Comparison of the fabricated CaC₂ sensor performance with those of the existing ones

Sensing technology	Fruit	Analyte	LOD	Time	Handling	Portable	Ref.
Infrared thermal emission sensor	Banana	Ethylene	5 ppm	—	Complicated	No	65
LS-capped AuNP sensor	Mango	CaC ₂	0.01 ppm	5 min	Complicated	No	40
HPLC/MS/MS	Banana	Ethephon	0.92 ppm	—	Complicated	No	66
Image processing/multispectral imaging sensor	Banana	—	—	—	Complicated	No	67
DTNB colorimetric sensor	Banana	CaC ₂	50 ppm	Prompt	Simple and user friendly	Yes	Present work

portability are the main concerns to be solved. Due to complex methodologies, these methods are expensive and instrument dependent as well as require expensive chemicals. Our sensor has simplified the assay method and results are understandable by applying one step detection. As summarized in Table S3 (ESI[†]), the presented sensor enhanced the sensitivity and selectivity, reduced sensing time and sample volume, also it is very convenient and portable when compared to other existing techniques as it is just a small bottle containing our active sensing solution and consumers just need to add few drops of sample to the test bottle. Overcoming all concerns and fulfilling all needs, creating our sensor could be a potentially ASSURED tool for CaC₂ detection in ARFs and filling the gap between laboratory-based analysis and on-site detection of analytes for a variety of purposes. The developed sensor must meet a few important specifications for its validation; it is all about sensor resolution, accuracy, and precision. The relative standard deviation (RSD) of the developed sensing in response to 50 ppm CaC₂ for ten measurements was observed as less than 5.0%, indicating good reproducibility of the developed sensor. The shelf life of the developed sensing solutions and their sensing responses were checked every two days for two weeks with 50 ppm CaC₂. The sensing solutions were stored at refrigerator temperature to monitor the storage stability. Each experiment was performed in triplicate. After two weeks of storage, the developed sensing solution retained about 92.5% of its initial response, indicating good storage stability and precision preferences.

3.8. Comparative study and validation of the DTNB-based sensor

A comparative study is required for the validation and assessment of the developed method against other credible traditional methods for ARF monitoring assurance. We performed and compared the CaC₂ analysis using our developed method and ICP-OES to confirm the presence of sulfide content and the linearity, sensitivity, and accuracy of our developed method for its validation. Fig. S6 (ESI[†]) depicts the ICP-OES's and our sensor's comparative absorbance linear plot of sulfide content in standard CaC₂ solutions with *R*² of 0.9895 and 0.9732, respectively. Data comparison demonstrates our sensor's linearity and sensitivity. Table S3 (ESI[†]) compares the detection patterns of sulfide in CaC₂ using the developed sensor (visual/colorimetric data for naked eye readability in the CaC₂ sample) and the detection analyzed using the traditional technique (ICP-ES data for sulfide quantification in the CaC₂ sample). It has been observed that the developed method has the ability to show instant color change-based detection at the CaC₂ concentration of 50 ppm; however, in ICP-MS, the detection

is based on total detectable sulfide content *via* an analytical technique which may also show recovery losses in trace sulfide content analysis. In addition to this, the complete analytical procedure of ICP-MS requires several typical steps, such as sample extraction (2–3 h), and a single sample run of 20–30 min for sulfide content detection in CaC₂, which could be eliminated *via* the currently developed detection method.

4. Conclusions

A colorimetric CaC₂ sensor was developed using an optimized enzymatic receptor DTNB solution. This paper describes the sensing and characterization of CaC₂ using an enzymatic chemoreceptor. The sensor was discovered to significantly detect the presence of CaC₂ in banana fruit peels. Our results revealed that the DTNB reagent emits a colorimetric (yellow) signal when it comes into contact with CaC₂. This signal indicates the presence of sulfide in CaC₂. The DTNB sensor is simple, and it responds to artificially ripened fruits, such as bananas, by exhibiting a strong color change from colorless to yellow to dark yellow. The sensor only responds to CaC₂ when exposed to another ripening agent. Furthermore, this work was optimized to prevent the consumption of artificially ripened fruits. The color transition of the DTNB enzymatic chemoreceptor sensors was discovered to have a linear relationship with increasing CaC₂ levels, and a redshift was observed after reaching the isosbestic point. Based on this linear trend, the LOD using a UV spectrophotometer was 10 ppm, while the LOD with the naked eye was 50 ppm.

Author contributions

Sonam Sonwal: writing – original draft, investigation, methodology, validation, and data curation. Shruti Shukla: data curation and writing – review and editing. Munirah Alhammadi: writing – review and editing. Reddicherla Umapathi: writing – review and editing. Hemanth P. K. Sudhani: writing – review and editing. Youngjin Cho and Yun Suk Huh: conceptualization, funding acquisition, writing – review and editing, and supervision.

Conflicts of interest

The authors declare no conflict of interest.

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