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# Paper/GO/e-Au flexible SERS sensors for *in situ* detection of tricyclazole in orange juice and on cucumber skin at the sub-ppb level: machine learning-assisted data analysis†

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Despite being an excellent surface enhanced Raman scattering (SERS) active material, gold nanoparticles were difficult to be loaded onto the surface of filter paper to fabricate flexible SERS substrates. In this study, electrochemically synthesized gold nanoparticles (e-AuNPs) were deposited on graphene oxide (GO) nanosheets in solution by ultrasonication, resulting in the formation of a GO/Au hybrid material. Thanks to the support of GO, the hybrid material could adhere onto the surface of filter paper, which was immersed into a GO/Au solution for 24 h and dried naturally at room temperature. The paper-based materials were then employed as substrates for a surface enhanced Raman scattering (SERS) sensing platform to detect tricyclazole (TCZ), a widely used pesticide, resulting in better sensitivity compared to the use of paper/Au SERS sensors. With the most optimal GO content of 4%, paper/GO/Au SERS sensors could achieve a limit of detection of  $1.32 \times 10^{-10}$  M in standard solutions. Furthermore, the filter paper-based SERS sensors also exhibited significant advantages in sample collection in real samples. On one hand, the sensors were dipped into orange juice, allowing TCZ molecules in this real sample to be adsorbed onto their SERS active surface. On the other hand, they were pasted onto cucumber skin to collect the analytes. As a result, the paper/GO/Au SERS sensors could sense TCZ in orange juice and on cucumber skin at concentrations as low as  $10^{-9}$  M ( $\sim 2$  ppb). In addition, a machine learning model was designed and developed, allowing the sensing system to discriminate TCZ from nine other organic compounds and predict the presence of TCZ on cucumber skin at concentrations down to  $10^{-9}$  M.

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## 1. Introduction

Raman spectroscopy is a non-destructive analytical tool to study molecular structures. It provides detailed information about the unique vibration modes of samples, allowing the identification of chemical compounds.<sup>1</sup> Since its discovery in 1977, surface-enhanced Raman spectroscopy (SERS) has gained increasing attention as it inherits from Raman the most important

characteristic of chemical fingerprinting while improving its sensitivity. In the SERS phenomenon, Raman scattering of analytes is significantly enhanced when they are adsorbed onto the surface of plasmonic materials, allowing the detection of analytes at trace and even single-molecule levels.<sup>2,3</sup> Noble metal nanoparticles (NPs), such as Au, Ag and Cu, have been employed as active substrates for SERS-based applications for environmental,<sup>4–6</sup> food<sup>7,8</sup> and health<sup>9,10</sup> safety monitoring.

In order to improve the performance of the SERS-based sensing systems, on one hand, many research groups have focused on modification of active materials to improve the sensitivity and reliability of the sensors. Plasmonic NPs with sharp tips and corners, such as nanostars,<sup>11,12</sup> nanoflowers,<sup>13</sup> nanocubes,<sup>14,15</sup> etc., were prepared as SERS substrates to sense various chemical and biological analytes. Besides, SERS active materials have been grown directly on rigid substrates to form nanogaps as well as generate high uniformity of the substrates.<sup>16,17</sup> These strategies aimed to create hot-spots, where the electromagnetic field is particularly intense, allowing the Raman signal to be enhanced in the electromagnetic

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mechanism (EM). Several other groups modified the active substrates to promote substrate-to-analyte charge transfer to enhance the Raman signal in the chemical mechanism (CM). Different semiconductor-metal composites were fabricated and employed as SERS substrates, showing an enhancement in the SERS signal of analytes, compared to using pure plasmonic nanomaterials.<sup>18–20</sup> Besides, with large surface area and excellent adsorption ability, graphene has been considered as the star of the two-dimensional (2D) materials family.<sup>21</sup> Many SERS sensing systems have been designed and developed based on plasmonic materials and graphene or its derivatives.<sup>22,23</sup>

On the other hand, researchers have been more concerned about the practicability of the SERS sensors in real samples. Real applications are challenging for the development of the SERS sensing platform because real samples are much more complicated than standard ones in the laboratory. To be specific, standard samples are usually prepared in solution, which makes it convenient for them to be drop-cast on the surface of widely used rigid substrates, such as silicon wafers and glass slides. However, in real samples, it requires SERS sensors to detect analytes not only in solution but also on different surfaces, either planar or non-planar ones. Thus, flexible substrates, which can be pasted and/or wrapped onto different surfaces, have been designed for both material deposition and sample collection. For example, polydimethylsiloxane (PDMS) films loaded with Au nanostars and nanorods have been fabricated to detect thiabendazole and thiram, respectively, on fruit skin.<sup>24,25</sup> In another approach, commercial adhesive tape was employed to fabricate “sticky” substrates.<sup>26–29</sup> Although the use of adhesive tape is cost- and time-effective, this sticky substrate is only suitable to collect materials and samples on dry surfaces, which would limit the variety of materials and samples. Being hydrophilic but insoluble in water, filter paper is more versatile. SERS active materials have been introduced onto in-paper substrates by drop-casting,<sup>30</sup> dip-coating,<sup>31</sup> spray coating<sup>32</sup> and thermal inkjet printing.<sup>33</sup> Subsequently, the as-prepared substrates can be dipped into aqueous media, or be attached to wet surfaces *via* van der Waals forces to absorb analytes. In a recent study, we fabricated an in-paper SERS sensor by dip-coating filter paper into a silver nanoparticle (AgNP) solution for 24 h and then drying naturally at room temperature. The versatile AgNP-based SERS sensor could detect methylene blue in river water and thiram on apple skin at concentrations down to  $1.0 \times 10^{-10}$  M.<sup>34</sup> In another study, Oliveira *et al.* prepared an Ag-based paper SERS sensor by a drop-casting method to detect rhodamine-6G with detection limit (LOD) as low as  $11.4 \pm 0.2$  pg.<sup>35</sup> These one-step substrate preparation methods were expected to be utilized to fabricate other paper-based substrates using other SERS active nanomaterials. Nevertheless, poor adsorption of gold nanoparticles (AuNPs) on the cellulose fiber network of filter paper, resulting in low SERS sensing performance of the SERS substrate, was reported in several studies.<sup>36,37</sup> However, it did not prevent researchers from developing the idea of combining the excellent SERS performance of AuNPs and the versatility of filter paper. Jang *et al.* suggested performing the spray coating technique to load AuNPs onto filter paper, leading to

enhancements in SERS signals over 2–5 times.<sup>36</sup> However, it required the utilization of a spray gun tool purged with N<sub>2</sub> gas, which should be performed in a laboratory by skilled labor. A simpler approach was proposed by Moram *et al.*, in which aggregation of AuNPs was induced by NaCl before the dip coating step.<sup>38</sup> The addition of NaCl solution (50 mM) led to an intermediate state of aggregation of AuNPs, reducing the distance between them and forming more hot spots. As a result, the SERS signal of MB on an aggregated-AuNP-based substrate was significantly improved. Nevertheless, the aggregation could not be well-controlled, resulting in poor uniformity of the in-paper substrate. Moreover, larger concentrations of NaCl promote the over-aggregation of AuNPs, and in contrast, reduce the number of hot spots, which was reflected by the rapid decrease in the SERS signal of MB on those substrates.

Another issue for the application of SERS sensors is complicated data analysis with the spectral overlapping of various molecules preventing sample detection and identification.<sup>39,40</sup> Recently, machine learning has been developed as a solution to discriminate chemical and biological compounds with similar Raman spectra.<sup>41–43</sup>

In this study, we utilized graphene oxide (GO) nanosheets as the support for grafting electrochemically synthesized AuNPs (e-AuNPs). Subsequently, the composite material was loaded onto filter paper by dip-coating to fabricate a paper/GO/e-Au SERS sensor for detection of tricyclazole (TCZ), a commonly used pesticide. With the most optimal GO content of 4%, it could detect TCZ at concentrations down to  $1.32 \times 10^{-10}$  M in standard solutions and  $10^{-9}$  M in real samples of orange juice and cucumber skin. Thus, we have fabricated a versatile in-paper substrate. Moreover, we also developed a machine learning model, allowing the SERS signal of TCZ to be distinguished from the signals of nine other organic compounds. On real samples of cucumber skin, the artificial neural network could detect and identify TCZ at concentrations down to  $10^{-9}$  M, which was in agreement with the practicability result of the sensing system. Hence, with the machine learning-assisted SERS sensing system, we have targeted all three approaches as discussed above, including: (i) developing a versatile flexible in-paper substrate for detection of an analyte in solution and non-planar surface; (ii) improving the SERS sensing system by material modification; and (iii) discriminating the desired analyte from other organic compounds *via* machine learning.

## 2. Materials and methods

### 2.1. Materials

Sodium citrate (Na<sub>3</sub>C<sub>6</sub>H<sub>5</sub>O<sub>7</sub>, 99.9%), tricyclazole (C<sub>9</sub>H<sub>7</sub>N<sub>3</sub>S, 97.0%), ethanol (C<sub>2</sub>H<sub>5</sub>OH, 98%), graphite flakes, nitric acid (HNO<sub>3</sub>, 90%), potassium permanganate (KMnO<sub>4</sub>, 99%), sodium nitrate (NaNO<sub>3</sub>, 99%) and sulfuric acid (H<sub>2</sub>SO<sub>4</sub>, 98%) were purchased from Shanghai Chemical Reagent and used directly without further purification. Two gold plates (purity: 99.99%) were prepared with dimensions of (100 mm × 5 mm × 0.5 mm). Filter paper was supplied from Hangzhou Special Paper Industry Co., Ltd (NEWSTAR INDUSTRY). Double distilled water was used throughout the experiments.



## 2.2. Electrochemically synthesized gold nanoparticles and their characterization

e-AuNPs were synthesized electrochemically in a 200 mL beaker containing 0.1%  $\text{Na}_3\text{C}_6\text{H}_5\text{O}_7$  in distilled water. The electrodes were polished and washed to remove surface oxides before the synthesis. Electrolysis was carried out at room temperature with magnetic stirring at 200 rpm for 2 hours. This resulted in a wine-red colloidal e-AuNP solution. The shapes and sizes of e-AuNPs were recorded by scanning electron microscopy (SEM, Hitachi S-4800) operated at an acceleration voltage of 5 kV, which revealed the spherical shape of the NPs with an average diameter of 19 nm (Fig. S1a†). UV-Vis absorption spectra were recorded using a JENWAY 6850 spectrophotometer. The as-prepared e-AuNPs exhibited a plasmonic band at  $\sim 532$  nm (Fig. S1b†). Full information about NP characterization is reported in our previous study.<sup>44</sup>

## 2.3. Preparation of graphene oxide sheets

GO was synthesized using the Hummers' method as described in a previous study of our group.<sup>45</sup> A mixture of graphite flakes,  $\text{HNO}_3$  and  $\text{KMnO}_4$  was generated with a ratio of 1 : 2 : 1.5 wt. The mixture was transferred to a Teflon autoclave, sealed, and heated in a microwave reactor (UWave-2000 Multifunctional Microwave, MW-2000/SINEO) with a power of 800 W for 1 min to obtain layered graphite. 2.0 g of layered graphite, 8.0 g of  $\text{KMnO}_4$  and 1.0 g of  $\text{NaNO}_3$  were added gradually into 160 mL of  $\text{H}_2\text{SO}_4$  at 5 °C, followed by constant stirring for 30 min. The temperature of the solution was gradually increased up to 45 °C. The solution was stirred constantly for 2 hours. GO was generated when deionized water was dropped into the solution and the temperature was raised to 95 °C for 1 hour. Subsequently,  $\text{H}_2\text{O}_2$  (30%) and HCl (10%) were added into GO solution to eliminate excessive  $\text{KMnO}_4$  and metal ions, and dissolve  $\text{MnO}_2$ . Finally, GO was washed with distilled water using centrifugation.

## 2.4. Fabrication of paper/e-Au and paper/GO/e-Au flexible SERS sensors

A mixture of e-AuNPs and GO in solution underwent 30 minutes of ultrasonication for deposition of e-AuNPs onto GO nano-sheets, resulting in a GO/e-Au solution. Filter paper was cut into 1 cm  $\times$  1 cm pieces and immersed in the solution for 24 hours and dried naturally at room temperature. The structure of cellulose fibers within filter paper before and after the loading of e-AuNPs and GO/e-Au was observed under a DN-117M digital microscope using a 40 $\times$  objective lens. The arrangement of e-AuNPs and GO/e-Au on the cellulose fiber surface in paper/e-Au and paper/GO/e-Au was analyzed by scanning electron microscopy (SEM, Hitachi S-4800) operated at an acceleration voltage of 5 kV.

## 2.5. Standard samples: sample collection and SERS measurements

Distilled water was used to prepare solutions with various concentrations of TCZ ranging from  $10^{-11}$  M to  $10^{-3}$  M. In each

SERS measurement, 5  $\mu\text{L}$  of TCZ solution was drop-cast onto paper/e-Au and paper/GO/e-Au flexible SERS sensors and dried naturally at room temperature. Subsequently, SERS spectra were recorded from the substrate using a MacroRaman™ Raman spectrometer (Horiba) with 785 nm laser excitation. Raman measurements were taken using a 100 $\times$  objective with a numerical aperture of 0.90. The laser power was set at 45 mW, with a 45° contact angle, resulting in a diffraction-limited laser spot diameter of 1.1  $\mu\text{m}$  ( $1.22\lambda/\text{NA}$ ) and a focal length of 115 nm. Each measurement lasted for 10 seconds with three accumulations. Finally, the spectrum was obtained after baseline calibration.

## 2.6. Real samples: sample collection and SERS measurements

Cucumbers were obtained from a local supermarket in Hanoi, Vietnam, and thoroughly washed. For each measurement, 10  $\mu\text{L}$  of TCZ solution at various concentrations (ranging from  $10^{-10}$  M to  $10^{-6}$  M) was directly dropped onto the surface of the cucumber skin and allowed to dry at room temperature. Following this, 10  $\mu\text{L}$  of 98% ethanol was dropped onto the spiked cucumber skin, and a paper/GO/e-Au SERS sensor was immediately attached to the peel. The SERS paper was pressed onto the surface for 3 minutes and then carefully peeled off, following a procedure referred to as the "paste and peel off" method. The SERS paper was subsequently dried at room temperature. SERS measurements were conducted according to the procedure outlined in the previous section.

Orange juice was also bought from a supermarket in Hanoi, Vietnam, and used directly for analysis. To achieve different concentrations (ranging from  $10^{-10}$  M to  $10^{-6}$  M), TCZ was spiked into the samples. The SERS paper was immersed in 10 mL of the aqueous TCZ solution for 30 minutes. Afterward, they were removed and allowed to naturally dry at room temperature, following a procedure referred to as the "dip and dry" method.

The recovery rate of each measurement is calculated using the formula:

$$\text{Recovery (\%)} = \frac{\text{spiked analyte concentration (M)}}{\text{calculated analyte concentration (M)}} \times 100\% \quad (1)$$

## 2.7. Machine learning-assisted data analysis

In order to speed up the process of classifying various pesticides acquired through the SERS sensors we leverage machine learning methodologies. In particular, our dataset consists of 50 spectral samples of 10 different compounds, each having 5 samples. Each data specimen encompasses intensity measurements at 2047 discrete Raman shift values, where only peak values, which retain the most salient information, are selected to feed the ML model. Depending on the compound's nature, the number of retained peaks varies from 2 to 4, resulting in a total of 34 Raman shift values per sample in the ultimate dataset. This dataset is partitioned into two subsets, namely,



a training set and a testing set, comprising 30 and 20 samples, respectively. In order to mitigate the occurrence of data samples exclusively appearing in either the training or test set, we deliberately select the training set and testing set to ensure each compound had three samples in the training set and two samples in the test set.

A wide range of multi-class classification ML models such as Logistic Regression, K-nearest Neighbor, Support Vector Machine, Decision Tree, Random Forest, and Naive Bayes are employed for classifying pesticides. The details of these models can be found in a textbook.<sup>46</sup> We first train these models using the training set and subsequently make predictions on the trained model to measure the accuracy of the substance classification within the testing set. The model achieving the highest classification accuracy will be selected for realistic substance classifications.

All aforementioned data analytics, dimensionality reduction, and model training are executed utilizing the Python programming framework, where Pandas, Scikit-learn, and Matplotlib are the main Python packages used in this contribution. Herein, note that we implement the grid search method in Python to find the best hyperparameters for each model, namely, Logistic Regression ( $C=10$ , solver='liblinear'), K-nearest Neighbor (metric='cosine',  $n\_neighbors=1$ ), Support Vector Machine ( $C=10$ , degree=2, kernel='linear'), Decision Tree (criterion='entropy', max\_depth=6, min\_samples\_split=2), Random Forest (criterion='gini', max\_depth=4, min\_samples\_split=5), and Naive Bayes (alpha=0.01).

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### 3. Results and discussion

#### 3.1. Fabrication of paper/GO/e-Au flexible SERS sensors

Paper/e-Au flexible SERS sensors were fabricated as described in our previous study,<sup>34</sup> in which the e-AgNP solution was replaced by an e-AuNP one. As e-AuNPs were also coated with citrate, it was expected that e-AuNPs could be attached to the filter paper *via* hydrogen bonds between their coating molecules and -OH groups on the cellulose chains of the paper. More importantly, Au-O could also be formed thanks to the presence of -OH groups on the surface of cellulose, leading to direct grafting of e-AuNPs onto the paper surface. However, after 24 h of soaking in the e-AuNP solution the filter paper only showed a slight change in color, which can be observed with the bare eye (Fig. 1a) as well as under a microscope (Fig. 1b and c). The deposition of e-AuNPs on the filter paper was confirmed by a SEM image (Fig. 1e), and is represented by the presence of bright grains on the surface of the paper. However, NP density was much lower than that when using e-AgNP solutions.<sup>34</sup> Thus, despite using the same method of substrate preparation, fewer e-AuNPs were anchored on the cellulose matrix of the filter paper. This was an unexpected result as Au-O bonding was reported to be a strong covalent one,<sup>47</sup> even stronger than Ag-O bonding.<sup>48</sup> However, in a 2015 study, Sun

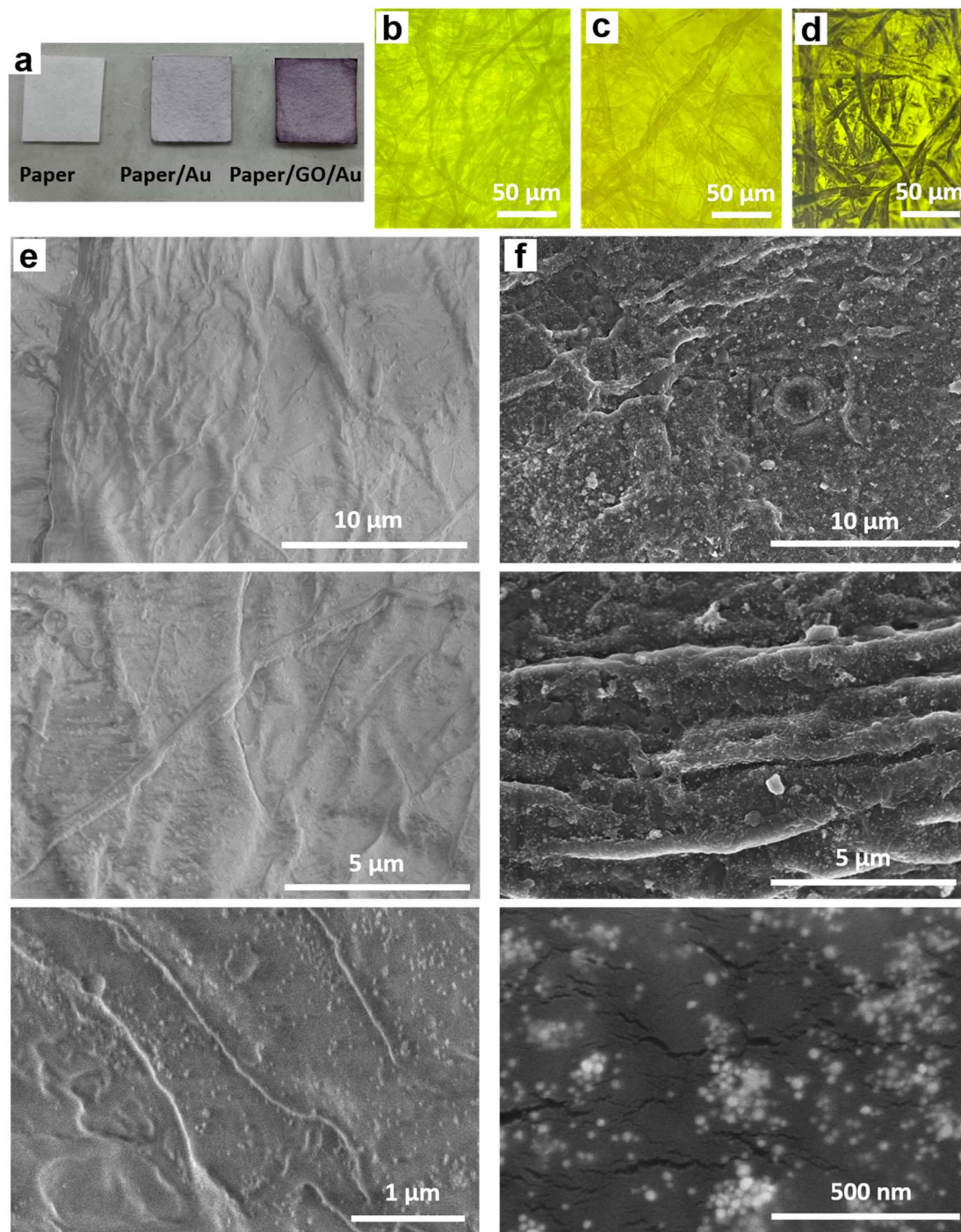
*et al.* calculated the binding energies of O in  $M_x-O$  ( $M = Ag$  or  $Au$ ) models. The authors claimed the binding energy of the Au-O or Ag-O bond could be suppressed by the neighboring Au or Ag atoms. This suppressing effect of the neighboring Au atoms is stronger than that by the Ag ones. Hence, despite the intrinsically stronger Au-O bond, the Au-O bond in  $Au_x-O$  usually appeared to be weaker than the Ag-O bond in  $Ag_x-O$ .<sup>49</sup> In this case, formation of the Au-O bond between the e-AuNPs and cellulose fibers of the filter paper could have been suppressed by many neighboring atoms, and therefore, it was weaker than the Ag-O bond. Hence, there were only a few e-AuNPs attached onto the surface of the filter paper. Furthermore, in our previous study, the size of e-AuNPs was reported to be 19 nm in diameter (Fig. S1a†).<sup>44</sup> The small size and weak Au-O bond would have prevented them from being anchored on the filter paper. Instead, they tended to be washed out through it. Therefore, this simple dip-and-dry method was not suitable to load e-AuNPs onto the paper substrate. Therefore, to combine the advantages of e-AuNPs and filter paper to fabricate a flexible SERS substrate, it required some modifications, which would allow the material to effectively adsorb onto the cellulose fibers. With a large surface, sheet-like GO was selected to combine with e-AuNPs to prepare a SERS active material which would not be easily filtered through the filter paper.

Pristine GO was synthesized using a modified Hummers' method, resulting in a sheet-like structure, which was reported in our 2021 study.<sup>50</sup> GO and e-AuNPs were mixed together at different ratios, including 1, 2, 3, 4 and 5 wt% of GO. Subsequently, the mixture underwent sonication for 30 min, allowing the anchoring of e-AuNPs onto the GO surface. It was reported that the carbon to oxygen ratio in GO could reach approximately three to one,<sup>51</sup> thanks to the presence of oxygenated functional groups such as ketone, hydroxyl, carboxyl and epoxy, exposed on both its edges and basal plane.<sup>52,53</sup> Except for the epoxy group, in which the O atom shares two stable covalent bonds with two C atoms, other functional group can be available for the attachment of e-AuNPs *via* Au-O bonds (Fig. 2 – left side). Although cellulose fibers are also rich in oxygen, the oxygen-containing groups on the surface of cellulose chains need to interact with each other to maintain the stability of the fibers. Therefore, compared to cellulose fibers, oxygen-containing functional groups on the surface of GO sheets could be more available to form Au-O bonds. Moreover, in a 2015 study, Wang and Liu performed a force analysis at the AuNP/GO interface, revealing that the attachment of AuNPs on the GO surface promoted wrinkle formation and the wrinkling in turn strengthened the binding of AuNPs.<sup>54</sup>

Paper/GO/e-Au SERS sensors were fabricated using filter paper and GO/Au solutions with different GO ratios by a dip-and-dry method. Thanks to the presence of GO, we could observe a significant change in the color of the filter paper by the bare eye (Fig. 1a). A microphotograph of the paper/GO/e-Au material with GO content of 4% (Fig. 1d) shows that the cellulose fibers of the filter paper were covered by a black material, which is the characteristic color of GO. This was confirmed by Raman spectroscopy. Fig. 2 shows the Raman spectrum of paper/GO/e-Au, in which the characteristic D- and G-bands of







**Fig. 1** (a) The color of the filter paper before (left) and after being immersed in e-AuNP solution (middle) and GO/e-Au with GO content of 4% (right); microscopic structure of filter paper (b), paper/e-Au (c) and paper/GO/e-Au with GO content of 4% (d); SEM images of paper/e-Au (e) and paper/GO/e-Au with GO content of 4% (f).

GO are obviously detected. In comparison to that of GO, the Raman spectrum of paper/GO/e-Au is noisier, which could be due to the presence of cellulose. However, it does not exhibit the characteristic peaks of cellulose such as those at  $1093\text{ cm}^{-1}$ ,  $1127\text{ cm}^{-1}$  and  $1382\text{ cm}^{-1}$ . In fact, these bands usually show low Raman intensity.<sup>34,55</sup> Moreover, they are overlapped with the D- and G-bands of GO, which exhibit extremely high intensity in

the Raman spectra. In the SEM image (Fig. 1f), the bright particles representing the presence of e-AuNPs were also revealed to be denser on the scaffold of GO on the filter paper. Therefore, with the assistance of GO during the fabrication of the in-paper SERS sensors, we enriched the density of the SERS active materials. As a result, the peak at  $239\text{ cm}^{-1}$  was detected at high intensity in the Raman spectrum of paper/GO/e-Au,



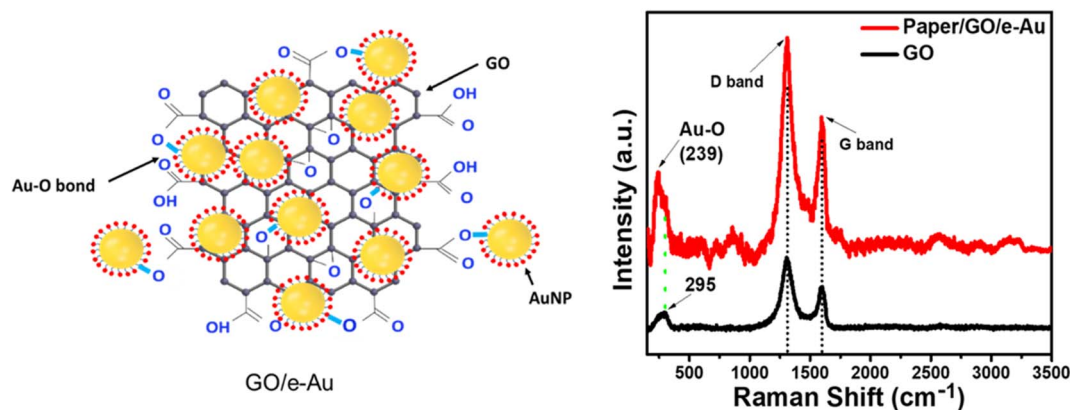


Fig. 2 (Left) Formation of Au–O bonds between e-AuNPs and GO; (right) Raman spectra of GO (black) and paper/GO/e-Au with GO content of 4% (red).

corresponding to the Au–O stretching mode,<sup>56</sup> which was schematically described in Fig. 2. This peak differs from the band in the Raman spectrum of GO (290  $\text{cm}^{-1}$ ), which could be due to excessive chemicals which were not eliminated properly during GO synthesis.

### 3.2. SERS sensing performance: a comparison of paper/e-Au and paper/GO/e-Au flexible SERS sensors

For standard samples, TCZ solution prepared in distilled water was drop-cast onto in-paper substrates as demonstrated in

Fig. 3a. On filter paper, in the absence of the active SERS substrate, there was no peak of TCZ detected (Fig. S2†). Fig. 3c compares the SERS spectra of TCZ ( $10^{-5}$  M) on different e-AuNP-based in-paper substrates without and with GO at different GO contents (*i.e.*, 1, 2, 3, 4 and 5% wt). Without GO, the SERS intensity of TCZ on the paper/e-Au substrate was low with characteristic peaks at 430  $\text{cm}^{-1}$ , 592  $\text{cm}^{-1}$ , 985  $\text{cm}^{-1}$ , 1312  $\text{cm}^{-1}$  and 1372  $\text{cm}^{-1}$ . However, the intensities of the peaks at 592  $\text{cm}^{-1}$  and 985  $\text{cm}^{-1}$  are relatively low, so they could not be clearly detected as Raman peaks (Fig. S2†). In the presence of

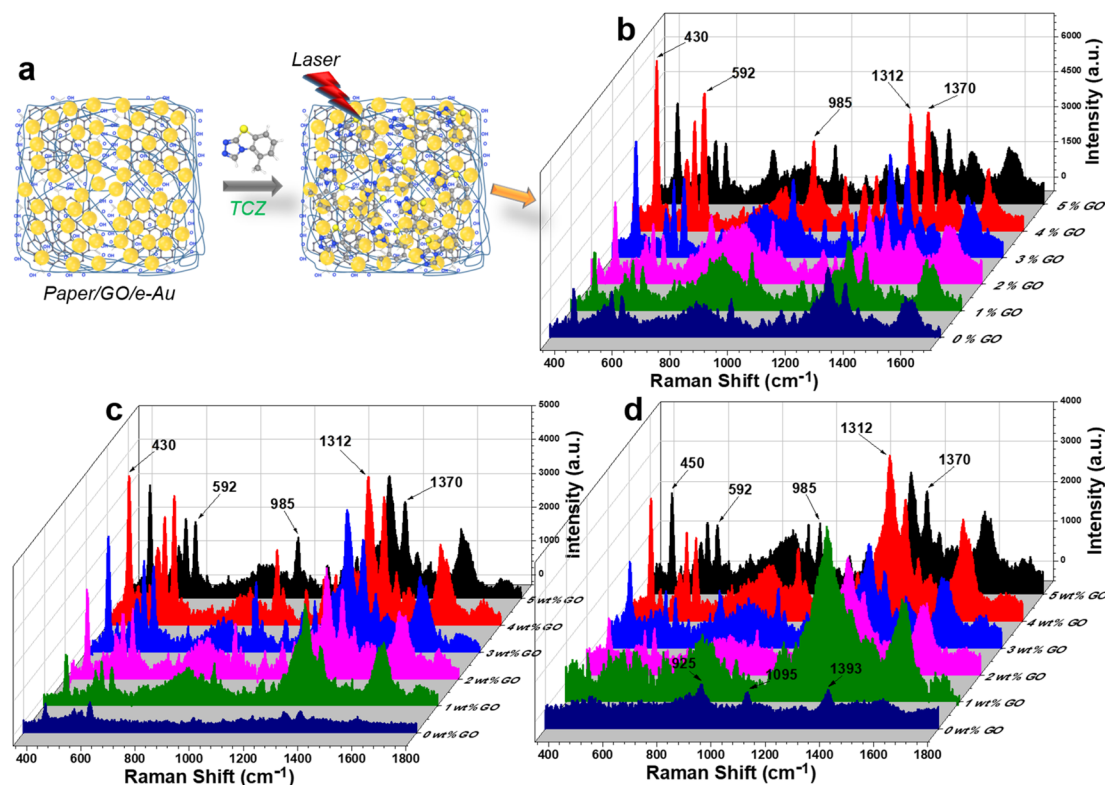


Fig. 3 (a) Schematic method to detect TCZ in standard samples; SERS spectra of TCZ at  $10^{-4}$  M (b),  $10^{-5}$  M (c) and  $10^{-6}$  M (d) using a paper/e-Au substrate and paper/GO/e-Au substrate with different GO contents.



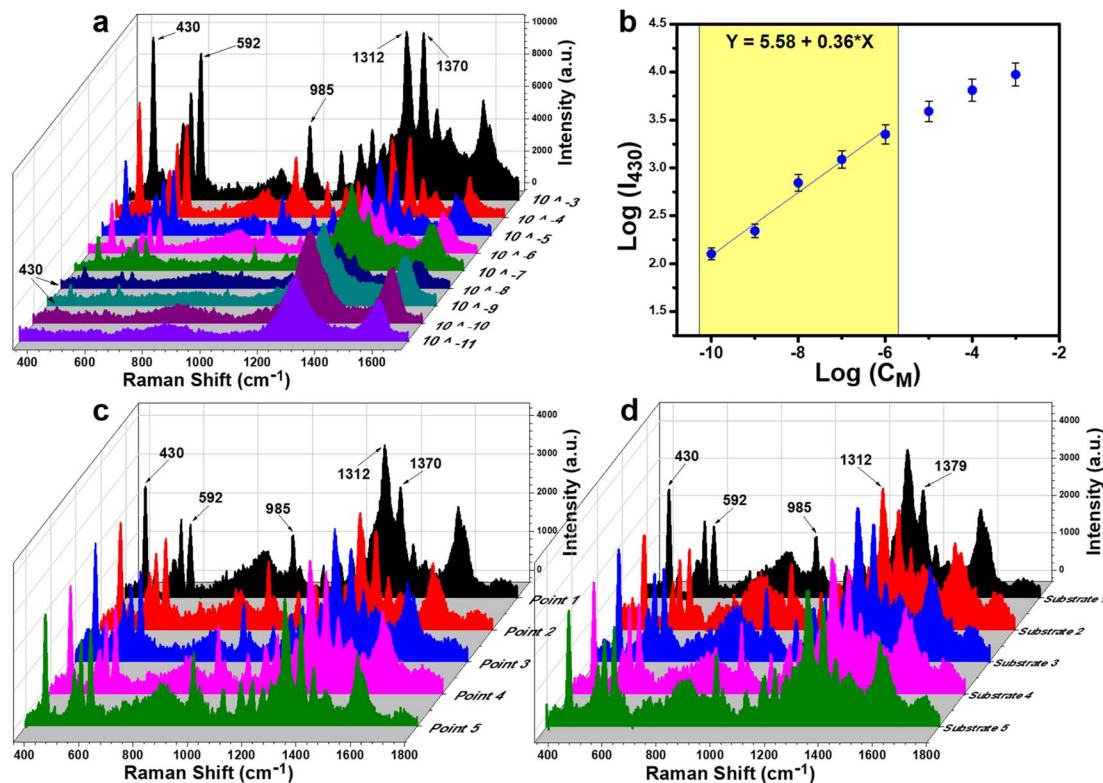


Fig. 4 (a) SERS spectra of TCZ ( $10^{-11}$  M to  $10^{-3}$  M) on the paper/GO/e-Au substrate; (b) plot of log of SERS intensity at  $430\text{ cm}^{-1}$  (slope  $0.36 \pm 0.02$ , intercept  $5.58 \pm 0.02$ ); uniformity (c) and reproducibility (d) of the SERS sensor for TCZ ( $10^{-5}$  M) detection based on the paper/GO/e-Au substrate.

GO in the in-paper substrate, we could observe those characteristic bands of TCZ with obvious improvement in intensity (Fig. 3b and S2†). The band at  $430\text{ cm}^{-1}$  represents C–N–C deformation vibration. The band at  $592\text{ cm}^{-1}$  is assigned to C–S–C deformation vibration. The band at  $985\text{ cm}^{-1}$  is associated with C–C symmetric stretching vibration. The bands at  $1312\text{ cm}^{-1}$  and  $1372\text{ cm}^{-1}$  represent C–N stretching

vibration.<sup>8,57,58</sup> In a recent study concerning the use of AgNPs to detect TCZ, Su *et al.* proposed that Ag–S bonds would have formed, allowing TCZ to replace citrate and fluorescein on the surface of AgNPs. In this case, Au–S bond formation would have also occurred. The sulfur atom in the TCZ molecule, which already shares two covalent bonds with two carbon atoms, could have been linked to Au *via* a dative bond with two shared

Table 1 Several reported SERS-based TCZ sensors using noble metal NPs as active substrates

Material	Measuring support	LOD	Linear range	Real sample	Sample collection method	Ref.
Au@AgNPs	Capillary tube	$2.75 \times 10^{-7}$ M	$2.75 \times 10^{-5}$ to $2.75 \times 10^{-7}$ M	Pear	Homogenizing and centrifuging samples for extraction; dipping SERS substrates into sample solutions	8
rGO–Ag nanocomposite	Glass slide	$10^{-6}$ M	$10^{-3}$ to $10^{-6}$ M	—	—	63
AgNPs	Si substrate	$5.28 \times 10^{-9}$ M	$5.28 \times 10^{-4}$ to $5.28 \times 10^{-9}$ M	Lettuce	Homogenizing and centrifuging samples for extraction; drop-casting the sample solution onto the SERS substrate	64
Multi-stacked Au–Ag bimetallic nanowires	Microwell (active substrates fixed within the wells)	$5.5 \times 10^{-8}$ M	$5.5 \times 10^{-4}$ to $5.5 \times 10^{-8}$ M	Whole milk	Directly drop-casting milk samples onto SERS substrates/ dipping SERS substrates into milk samples	65
GO/e-Au	Filter paper	$1.32 \times 10^{-10}$ M	$10^{-6}$ to $10^{-11}$ M	Cucumber Orange juice	“Paste and peel off” for cucumber skin “Dip and dry” for orange juice	This work





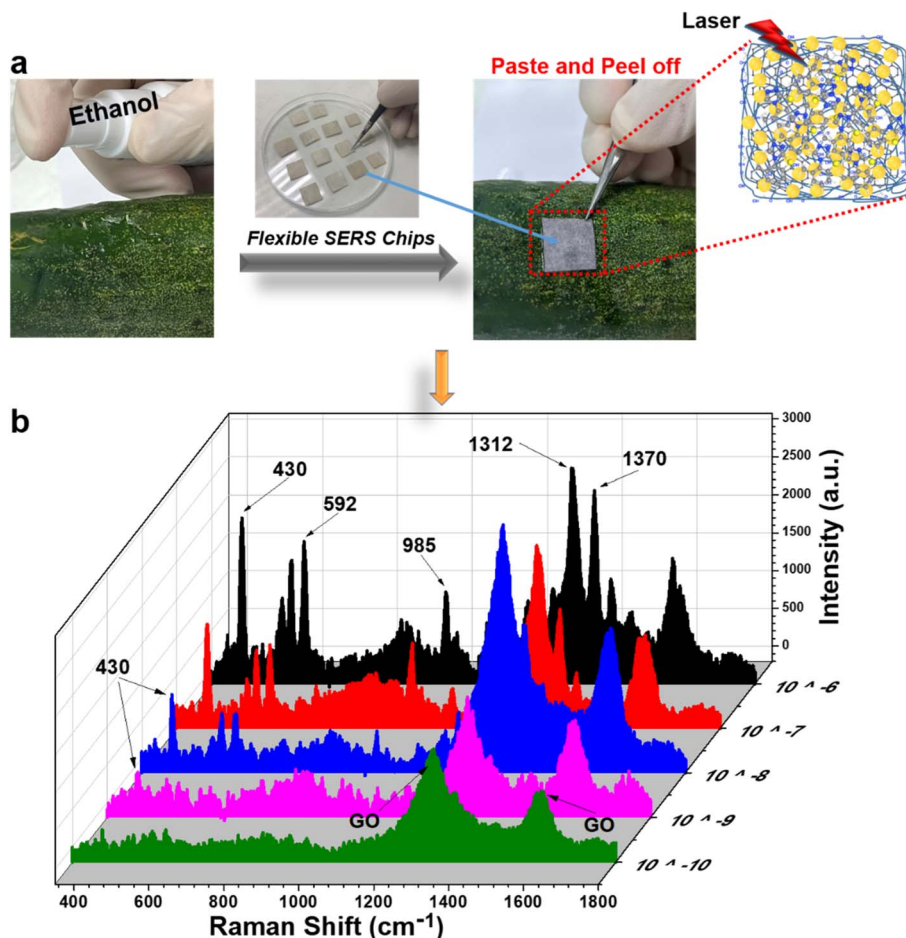


Fig. 5 (a) "Paste and peel off" method to collect TCZ on the surface of a cucumber; (b) SERS spectra of thiram (10<sup>-10</sup> M to 10<sup>-4</sup> M) on the surface of a cucumber using the paper/GO/e-Au SERS substrate.

electrons (Fig. S4†).<sup>59,60</sup> Therefore, the band of the C-S-C bond remained even when TCZ was linked to AuNPs. Besides, the formation of the dative bond also reduced the distance between TCZ and AuNPs, allowing TCZ to better experience the SERS effect *via* EM enhancement. In addition, this chemical linkage was convenient for charge transfer.

With the addition of GO, SERS enhancement of TCZ has been improved. It is clear that at low content of GO (1–4%), the SERS intensity of TCZ increases with the increase of GO content. Higher content of GO could have improved the adsorption of

TCZ onto paper/GO/e-Au. In addition, the increase in GO content might have provided more carboxyl groups for the attachment of AuNPs, leading to the enrichment of AuNPs on the cellulose fibers. However, when GO content reaches 5%, the TCZ intensity decreases compared to the use of paper/4% GO/e-Au. It is worth mentioning that the decrease is the most significant at the band of 592 cm<sup>-1</sup> (63%) while the decreases in the levels of TCZ intensity are 45%, 50%, 47%, and 45% at 430 cm<sup>-1</sup>, 985 cm<sup>-1</sup>, 1312 cm<sup>-1</sup> and 1372 cm<sup>-1</sup>, respectively. Fig. S3† shows the intensity of the band at 592 cm<sup>-1</sup> of TCZ

Table 2 Practicability of paper/GO/e-Au SERS sensors for TCZ detection on cucumber skin and in orange juice

Real sample	Spiked concentration (M)	Detected concentration (M)	Recovery (%)
Cucumber skin	10 <sup>-6</sup>	9.32 × 10 <sup>-7</sup>	93
	10 <sup>-7</sup>	8.21 × 10 <sup>-8</sup>	82
	10 <sup>-8</sup>	8.54 × 10 <sup>-9</sup>	85
	10 <sup>-9</sup>	7.82 × 10 <sup>-10</sup>	78
	10 <sup>-10</sup>	8.72 × 10 <sup>-7</sup>	87
Orange juice	10 <sup>-6</sup>	8.72 × 10 <sup>-7</sup>	87
	10 <sup>-7</sup>	9.11 × 10 <sup>-8</sup>	91
	10 <sup>-8</sup>	7.58 × 10 <sup>-9</sup>	76
	10 <sup>-9</sup>	8.59 × 10 <sup>-10</sup>	86





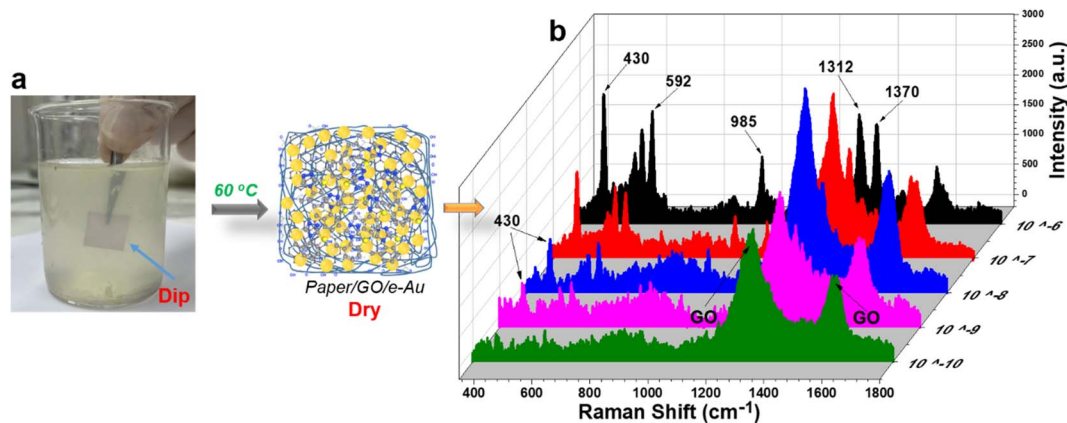


Fig. 6 (a) "Dip and dry" method to collect TCZ in orange juice; (b) SERS spectra of TCZ ( $10^{-10}$  M to  $10^{-4}$  M) in orange juice using paper/GO/e-Au SERS substrates.

( $10^{-5}$  M) on paper/GO/e-Au with GO content ranging from 0% to 5%. A similar trend could be observed with the concentrations of  $10^{-4}$  M and  $10^{-6}$  M TCZ (Fig. 3b and d). As the band at  $592\text{ cm}^{-1}$  was assigned to C-S-C deformation vibration, which is closest to the Au surface due to Au-S bonding, the decrease at this band might be related to the decrease of Au surface area on the substrate. Hence, a high GO content of 5% could have allowed GO nanosheets to envelop AuNPs, preventing TCZ to bind directly to AuNPs *via* Au-S linkage. Moreover, it was reported that the encapsulation of AuNPs by GO could reduce the SERS effect of the plasmonic material because it could cover AuNPs and block the nanogaps between them, preventing analytes from experiencing EM enhancements.<sup>21,61</sup> Hence, with a GO content of 5%, we also observed the decrease in SERS intensity at other characteristic bands of TCZ due to partial encapsulation of AuNPs by GO. To overcome this, Xu *et al.* had to perform thermal annealing at  $400\text{ }^{\circ}\text{C}$  for 2 h on their graphene-encapsulated SERS substrate to activate its SERS effect.<sup>61</sup> Nevertheless, in the case of depositing GO/e-Au composites on paper substrates, that thermal treatment was not a suitable approach for our system. Therefore, we optimized the performance of the paper/GO/e-Au substrate by selecting the most appropriate content of GO sheets to prevent them from covering the e-AuNPs. Therefore, the GO content of 4% was selected to further develop TCZ sensors.

The presence of GO nanosheets not only enriched e-AuNPs on the in-paper SERS sensors, but also improved the adsorption of analytes onto the substrate. Kim *et al.* stated that molecules containing a large number of C=C bonds were more adhesive to the GO surface.<sup>62</sup> Meanwhile, TCZ contains an aromatic ring with 3 C=C bonds. It allowed TCZ to be adsorbed onto the GO surface. As a result, in addition to the TCZ molecules that bind directly to the Au surface, in the presence of the GO scaffold, more TCZ molecules would locate near AuNPs to experience the resonance effect around the SERS active materials. This might also explain the appearance of the band at  $985\text{ cm}^{-1}$ , which represents the C-C symmetric stretching vibration, in the SERS spectra of TCZ on all GO-containing in-paper substrates.

Paper/4% GO/e-Au was selected to be the substrate for the TCZ sensors. Nine TCZ solutions at different concentrations ( $10^{-3}$  M to  $10^{-11}$  M) were prepared in distilled water. Subsequently, these samples were drop-cast on the prepared substrate. Fig. 4a demonstrates the SERS spectra of TCZ at these concentrations on the paper/4% GO/e-Au substrate. It is obvious that the SERS intensity of TCZ increases with the increase of its concentration. Down to the concentration of  $10^{-10}$  M, the characteristic bands of TCZ are still able to be detected, and then disappear at the concentration of  $10^{-11}$  M. The plot of logarithmic SERS intensity at  $430\text{ cm}^{-1}$  against the logarithmic concentration of TCZ shows a good linear relationship in the region from  $10^{-6}$  M to  $10^{-10}$  M with a linear regression of  $R^2 = 0.98$  (Fig. 4b). For each point of the calibration curve, SERS signals were recorded three times. Based on the linear equation in Fig. 4b, the LOD was calculated to be  $1.32 \times 10^{-10}$  M, which is lower than that of many established SERS-based sensors for TCZ detection using noble metal NPs as shown in Table 1. Moreover, thanks to the assistance of GO nanosheets and cellulose fibers, e-AuNPs were spread quite evenly on the SERS paper. Uniformity of the SERS sensors was evaluated by measuring five random points on one substrate (Fig. 4c), revealing an RSD of 9.5%. Besides, five different paper/4% GO/e-Au substrates were prepared independently using the method as described in Section 2.4. The SERS signals of TCZ ( $10^{-5}$  M) on these substrates were recorded, resulting in five spectra as shown in Fig. 4d. The RSD value for the reproducibility of the method was calculated to be 12.1%. LOD and RSD values were calculated as described in the ESI.†

### 3.3. Practicability of paper/GO/e-Au sensors to detect TCZ in real samples

As discussed in our previous study,<sup>34</sup> in-paper SERS sensors were convenient to detect analytes using two sample collection methods: "paste and peel off" and "dip and dry". Using both methods, the practicability of paper/GO/e-Au sensors to detect TCZ was evaluated on cucumber skin and orange juice, respectively.



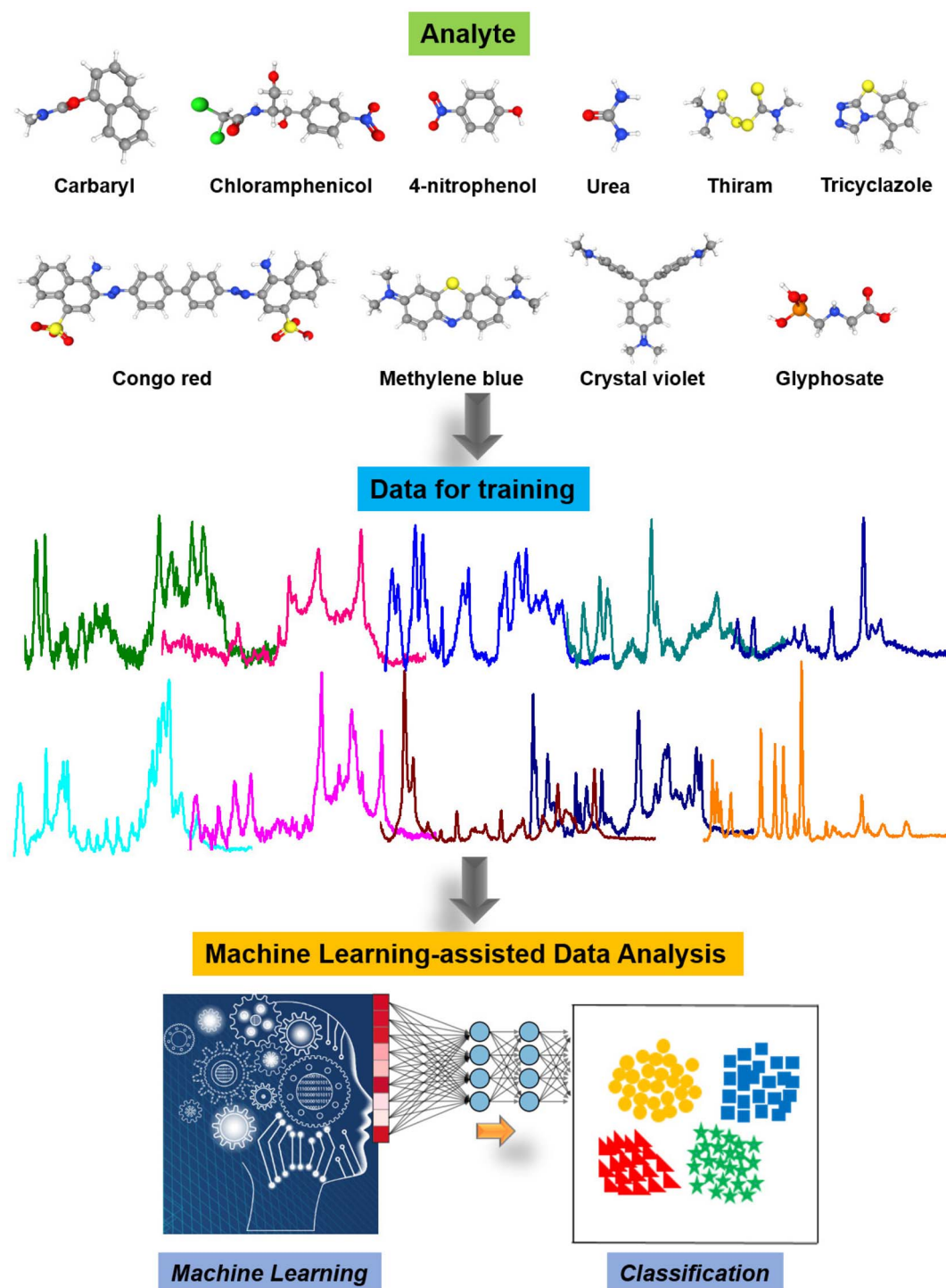


Fig. 7 Designing data for machine-learning-assisted identification and quantification of TCZ on cucumber skin. Our SERS data design comprises ten molecules including carbaryl, chloramphenicol, 4-nitrophenol, urea, thiram, TCZ, Congo red, methylene blue, crystal violet, and glyphosate.

The “paste and peel off” method was employed to collect TCZ on cucumber peel as described in Section 2.5 and Fig. 5a. To disassociate TCZ from the peel, alcohol solution was sprayed on the cucumber skin. This simple extraction allowed TCZ to be exposed on the cucumber peel. Moreover, water in the alcohol solution slowed the evaporation of the solution, and therefore, the in-paper sensors could adhere on it to adsorb the analyte

extracted from the cucumber skin. Fig. 5b shows the SERS spectra of TCZ at different concentrations ( $10^{-6}$  M to  $10^{-10}$  M) on cucumber peel collected on the SERS sensors. The characteristic peaks of TCZ can be detected in the spectra at concentrations down to  $10^{-9}$  M. The recovery rates range from 78 to 93% (Table 2). A part of spiked TCZ might have been absorbed into deeper layers of the peel matrix, which could not be extracted by alcohol



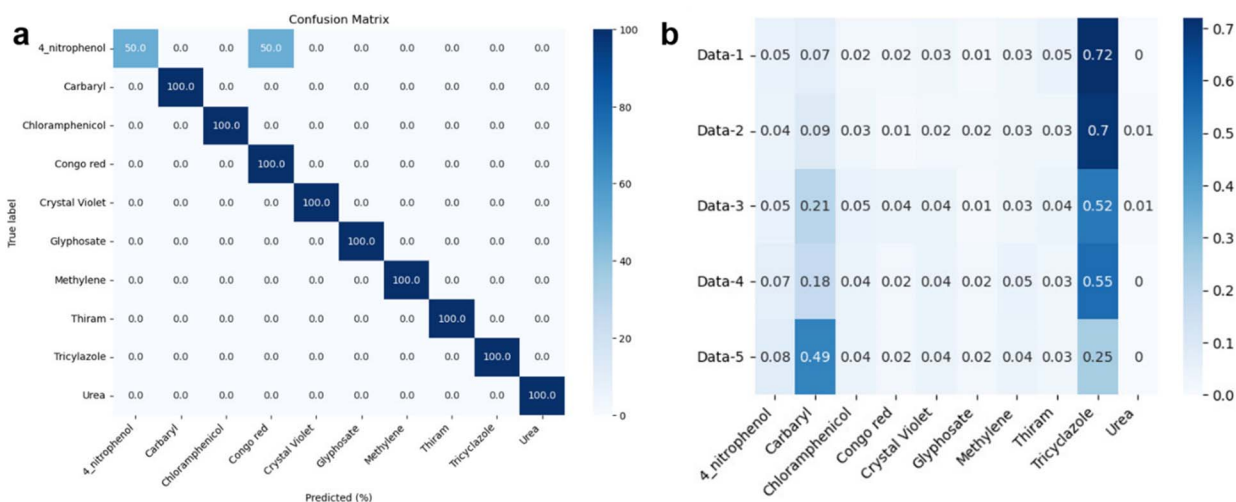


Fig. 8 (a) Confusion matrix and (b) prediction of TCZ on cucumber skin (data-1: TCZ ( $10^{-6}$  M); data-2: TCZ ( $10^{-7}$  M); data-3: TCZ ( $10^{-8}$  M); data-4: TCZ ( $10^{-9}$  M); data-5: TCZ ( $10^{-10}$  M)).

solution, and therefore, the recovery rates were lower than 100%. Moreover, the paper/GO/Ag SERS sensors allowed the detection of TCZ at concentrations as low as  $10^{-9}$  M ( $\sim 2$  ppb), which is much lower than the maximal residue limit (MRL) of 2 ppm in vegetables in China and the MRL of 0.01 ppm in vegetables in the European Union (EU) and Japan. In addition, with the use of this in-paper SERS sensor, we could avoid a step of sample preparation including homogenizing and multiple rounds of centrifuging, which was usually performed for sample collection in fruit and vegetable samples (Table 1).

The “dip and dry” method was utilized to collect TCZ spiked in orange juice at different concentrations from  $10^{-6}$  M to  $10^{-10}$  M as described in Section 2.5 and Fig. 6a. Fig. 6b shows the SERS spectra of TCZ at these concentrations on paper/GO/e-Au SERS sensors. Similar to the sample on cucumber skin, characteristic peaks can be detected in the spectra of TCZ at concentrations down to  $10^{-9}$  M. The recovery rates range from 76 to 91% (Table 2). Once again, the recovery rates are all lower than 100%. It is worth mentioning that for real samples, both cucumber skin and orange juice, sample collection methods were different from that used to collect TCZ in standard samples, for which, we drop-cast TCZ solution directly onto the flexible SERS substrate. As a result, the in-paper substrate could adsorb more TCZ molecules, leading to a higher SERS signal. Therefore, the recovery rates of the real sample are all lower than 100%. However, in this study, we aim to develop a flexible SERS substrate that can collect TCZ directly from the real samples without complicated pre-preparation, and the concentration of TCZ collected from different samples with different natures using different sample collection methods can be calculated using the same calibration curve with acceptable recovery rates. Paper/GO/e-Au SERS sensors were still able to detect TCZ at concentrations as low as  $10^{-9}$  M in orange juice and on cucumber skin using a fast and simple procedure; therefore, the SERS papers are promising to be employed in other real applications.

### 3.4. Machine learning-assisted data analysis

Fig. 7 illustrates the machine learning-assisted system we designed for discriminating TCZ from nine other organic compounds, including carbaryl, chloramphenicol, 4-nitrophenol, urea, thiram, Congo red, methylene blue, crystal violet and glyphosate. In addition to those of TCZ, the SERS spectra of these compounds were recorded and employed as training data, enriching the neural network. For each compound, we selected several characteristic peaks as shown in Table S1.† Training and testing were performed as described in Section 2.6. The mean normalized number of spectra sorted in the validation set is presented as a confusion matrix (Fig. 8A), showing excellent capability for discriminating TCZ from other compounds with an accuracy of 100%. The network could predict the presence of TCZ not only in standard samples, but also in real samples at different concentrations. Fig. 8B demonstrates the prediction of the network using the SERS spectra of TCZ collected on cucumber skin as described in Section 3.4 at the concentration range of  $10^{-6}$  M to  $10^{-10}$  M. From  $10^{-6}$  M to  $10^{-9}$  M, the network could predict the presence of TCZ with accuracies higher than 50%. Nevertheless, at a concentration of  $10^{-10}$  M, the prediction was incorrect. This result was in agreement with the SERS spectrum of TCZ ( $10^{-10}$  M) in Fig. 5b, in which we hardly detected the characteristic bands of TCZ. This preliminary result is encouraging as the system could predict the presence of TCZ on real samples at concentrations down to  $10^{-9}$  M ( $\sim 2$  ppb), which is lower than the MRL of TCZ established in various countries and regions around the world.

## 4. Conclusions

In this study, we designed and developed a flexible sensing platform based on paper/GO/e-Au SERS sensors and an artificial neural network to detect and identify TCZ in real samples. With the most optimal content of 4%, GO not only played an important role in material fabrication as a platform for the





anchoring of e-AuNPs, but also promoted the adsorption of the analyte, resulting in an impressive LOD of  $1.32 \times 10^{-10}$  M in standard samples. Moreover, the flexible substrate could detect TCZ in real samples of cucumber skin and orange juice at concentrations down to  $10^{-9}$  M ( $\sim 2$  ppb). Furthermore, we developed a machine learning model, which could discriminate TCZ from nine other organic compounds. Also, it could predict the presence of TCZ on cucumber skin at concentrations as low as  $10^{-9}$  M. The study confirms the potential of integrating machine learning and flexible substrates for fast determination of chemicals in real samples.

## Author contributions

H. A. Nguyen: conceptualization, methodology, formal analysis, writing – original draft; Q. D. Mai: conceptualization, validation, investigation, writing – review & editing; D. T. N. Nga: conceptualization, validation, investigation; M. K. Pham: validation, investigation; Q. K. Nguyen: methodology, validation, AI-data analysis; T. H. Do: methodology, validation, AI-data analysis; V. T. Luong: methodology, validation, AI-data analysis, writing – review & editing; V. D. Lam: methodology, validation, formal analysis; A. T. Le: conceptualization, methodology, supervision, project administration, writing – review & editing.

## Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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## References

- 1 D. R. Walt, I. Biran and T. K. Mandal, in *Encyclopedia of Physical Science and Technology*, ed. R. A. Meyers, Academic Press, New York, 3rd edn, 2003, pp. 803–829, DOI: [10.1016/B0-12-227410-5/00954-6](#).
- 2 X. X. Han, R. S. Rodriguez, C. L. Haynes, Y. Ozaki and B. Zhao, *Nat. Rev. Methods Primers*, 2022, **1**, 87.
- 3 K. Chen, Y. H. Ong, C. Yuen and Q. Liu, in *Imaging in Dermatology*, ed. M. R. Hamblin, P. Avci and G. K. Gupta, Academic Press, Boston, 2016, pp. 141–154, DOI: [10.1016/B978-0-12-802838-4.00013-3](#).
- 4 J. F. Li, Y. F. Huang, Y. Ding, Z. L. Yang, S. B. Li, X. S. Zhou, F. R. Fan, W. Zhang, Z. Y. Zhou, D. Y. Wu, B. Ren, Z. L. Wang and Z. Q. Tian, *Nature*, 2010, **464**, 392–395.
- 5 R. A. Halvorson and P. J. Vikesland, *Environ. Sci. Technol.*, 2010, **44**, 7749–7755.
- 6 W. Liao, Q. Lin, S. Xie, Y. He, Y. Tian and Y. Duan, *Anal. Chim. Acta*, 2018, **1043**, 64–71.
- 7 L. Wu, W. Zhang, C. Liu, M. F. Foda and Y. Zhu, *Food Chem.*, 2020, **328**, 127106.
- 8 N. Hussain, H. Pu and D.-W. Sun, *Food Chem.*, 2021, **350**, 129025.
- 9 Y. Peng, C. Lin, Y. Li, Y. Gao, J. Wang, J. He, Z. Huang, J. Liu, X. Luo and Y. Yang, *Matter*, 2022, **5**, 694–709.
- 10 R. Haldavnekar, K. Venkatakrishnan and B. Tan, *Nat. Commun.*, 2018, **9**, 3065.
- 11 A. S. D. S. Indrasekara, S. Meyers, S. Shubeita, L. C. Feldman, T. Gustafsson and L. Fabris, *Nanoscale*, 2014, **6**, 8891–8899.
- 12 X. Qi, X. Wang, Y. Dong, J. Xie, X. Gui, J. Bai, J. Duan, J. Liu and H. Yao, *Spectrochim. Acta, Part A*, 2022, **272**, 120955.
- 13 Y. Xu, F. Y. H. Kutsanedzie, M. M. Hassan, J. Zhu, H. Li and Q. Chen, *Sens. Actuators, B*, 2020, **324**, 128718.
- 14 H.-C. Hu, S.-H. Wu, L.-X. Jin and J.-J. Sun, *Biosens. Bioelectron.*, 2022, **210**, 114283.
- 15 J. M. McLellan, Z.-Y. Li, A. R. Siekkinen and Y. Xia, *Nano Lett.*, 2007, **7**, 1013–1017.
- 16 Y. Shan, Y. Yang, Y. Cao, C. Fu and Z. Huang, *Nanotechnology*, 2016, **27**, 145502.
- 17 T. Xu, J. Fu, X. Wang, G. Lu and B. Liu, *Front. Chem.*, 2022, **10**, 887900.
- 18 S. Cong, X. Liu, Y. Jiang, W. Zhang and Z. Zhao, *Innovation*, 2020, **1**, 100051.
- 19 J. Zhou, J. Zhang, H. Yang, Z. Wang, J.-a. Shi, W. Zhou, N. Jiang, G. Xian, Q. Qi, Y. Weng, C. Shen, Z. Cheng and S. He, *Nanoscale*, 2019, **11**, 11782–11788.
- 20 L.-J. Gu, C.-L. Ma, X.-H. Zhang, W. Zhang, S. Cong and Z.-G. Zhao, *Chem. Commun.*, 2018, **54**, 6332–6335.
- 21 N. Zhang, L. Tong and J. Zhang, *Chem. Mater.*, 2016, **28**, 6426–6435.
- 22 Y. Long, W. Wang, W. Xiong and H. Li, *J. Alloys Compd.*, 2022, **901**, 163660.
- 23 C. Zhu, Q. Zhao, X. Wang, Z. Li and X. Hu, *Microchem. J.*, 2021, **165**, 106090.
- 24 B. Fortuni, Y. Fujita, M. Ricci, T. Inose, R. Aubert, G. Lu, J. A. Hutchison, J. Hofkens, L. Latterini and H. Uji-i, *Chem. Commun.*, 2017, **53**, 5121–5124.
- 25 S. Kumar, P. Goel and J. P. Singh, *Sens. Actuators, B*, 2017, **241**, 577–583.
- 26 J. Jiang, S. Zou, Y. Li, F. Zhao, J. Chen, S. Wang, H. Wu, J. Xu, M. Chu, J. Liao and Z. Zhang, *Microchim. Acta*, 2019, **186**, 603.
- 27 J. Chen, Y. Huang, P. Kannan, L. Zhang, Z. Lin, J. Zhang, T. Chen and L. Guo, *Anal. Chem.*, 2016, **88**, 2149–2155.
- 28 J. Jiang, S. Zou, L. Ma, S. Wang, J. Liao and Z. Zhang, *ACS Appl. Mater. Interfaces*, 2018, **10**, 9129–9135.
- 29 J. Sitjar, J.-D. Liao, H. Lee, L. P. Pan, B. H. Liu, W.-e. Fu and G. D. Chen, *Nanomaterials*, 2019, **9**, 1750.
- 30 Y. Hong, Y. Li, L. Huang, W. He, S. Wang, C. Wang, G. Zhou, Y. Chen, X. Zhou, Y. Huang, W. Huang, T. Gong and Z. Zhou, *J. Biophotonics*, 2020, **13**, e201960176.
- 31 C. Wang, B. Liu and X. Dou, *Sens. Actuators, B*, 2016, **231**, 357–364.
- 32 H. S. Siebe, Q. Chen, X. Li, Y. Xu, W. R. Browne and S. E. J. Bell, *Analyst*, 2021, **146**, 1281–1288.



- 33 P. M. Fierro-Mercado and S. P. Hernández-Rivera, *Int. J. Spectrosc.*, 2012, **2012**, 716527.
- 34 Q. D. Mai, H. A. Nguyen, N. X. Dinh, N. T. Thu Thuy, Q. H. Tran, P. C. Thanh, A.-T. Pham and A.-T. Le, *Talanta*, 2023, **253**, 124114.
- 35 M. J. Oliveira, P. Quaresma, M. Peixoto de Almeida, A. Araújo, E. Pereira, E. Fortunato, R. Martins, R. Franco and H. Águas, *Sci. Rep.*, 2017, **7**, 2480.
- 36 W. Jang, H. Byun and J.-H. Kim, *Mater. Chem. Phys.*, 2020, **240**, 122124.
- 37 S. S. B. Moram, C. Byram, S. N. Shibu, B. M. Chilukamarri and V. R. Soma, *ACS Omega*, 2018, **3**, 8190–8201.
- 38 S. S. B. Moram, C. Byram and V. R. Soma, *Bull. Mater. Sci.*, 2020, **43**, 53.
- 39 N. Kim, M. R. Thomas, M. S. Bergholt, I. J. Pence, H. Seong, P. Charchar, N. Todorova, A. Nagelkerke, A. Belessiotis-Richards, D. J. Payne, A. Gelmi, I. Yarovsky and M. M. Stevens, *Nat. Commun.*, 2020, **11**, 207.
- 40 H. Zhou, L. Xu, Z. Ren, J. Zhu and C. Lee, *Nanoscale Adv.*, 2023, **5**, 538–570.
- 41 F. U. Ciloglu, M. Hora, A. Gundogdu, M. Kahraman, M. Tokmakci and O. Aydin, *Anal. Chim. Acta*, 2022, **1221**, 340094.
- 42 J. Q. Li, P. V. Dukes, W. Lee, M. Sarkis and T. Vo-Dinh, *J. Raman Spectrosc.*, 2022, **53**, 2044–2057.
- 43 Y. X. Leong, Y. H. Lee, C. S. L. Koh, G. C. Phan-Quang, X. Han, I. Y. Phang and X. Y. Ling, *Nano Lett.*, 2021, **21**, 2642–2649.
- 44 M. Q. Doan, N. H. Anh, N. X. Quang, N. X. Dinh, D. Q. Tri, T. Q. Huy and A.-T. Le, *J. Electron. Mater.*, 2022, **51**, 150–162.
- 45 P. N. Minh, V.-T. Hoang, N. X. Dinh, O. Van Hoang, N. Van Cuong, D. Thi Bich Hop, T. Q. Tuan, N. T. Khi, T. Q. Huy and A.-T. Le, *New J. Chem.*, 2020, **44**, 7611–7620.
- 46 S. Ben-David and S. Shalev-Shwartz, *Understanding Machine Learning: from Theory to Algorithms*, Cambridge University Press, Cambridge, 2014.
- 47 H.-J. Zhai, C. Bürgel, V. Bonacic-Koutecky and L.-S. Wang, *J. Am. Chem. Soc.*, 2008, **130**, 9156–9167.
- 48 K. A. Moltved and K. P. Kepp, *J. Phys. Chem. C*, 2019, **123**, 18432–18444.
- 49 K. Sun, M. Kohyama, S. Tanaka and S. Takeda, *J. Energy Chem.*, 2015, **24**, 485–489.
- 50 M. Q. Doan, N. H. Anh, H. Van Tuan, N. C. Tu, N. H. Lam, N. T. Khi, V. N. Phan, P. D. Thang and A.-T. Le, *Adsorpt. Sci. Technol.*, 2021, **2021**, 1169599.
- 51 I. Tiginyanu, V. Ursaki and V. Popa, in *Nanocoatings and Ultra-Thin Films*, ed. A. S. H. Makhlof and I. Tiginyanu, Woodhead Publishing, 2011, pp. 330–354, DOI: [10.1533/9780857094902.2.330](https://doi.org/10.1533/9780857094902.2.330).
- 52 S. Azizighannad and S. Mitra, *Sci. Rep.*, 2018, **8**, 10083.
- 53 A. Abdelkhalek, M. A. El-Latif, H. Ibrahim, H. Hamad and M. Showman, *Sci. Rep.*, 2022, **12**, 7060.
- 54 F. Wang and J. Liu, *Nanoscale*, 2015, **7**, 919–923.
- 55 J. H. Wiley and R. H. Atalla, *Carbohydr. Res.*, 1987, **160**, 113–129.
- 56 A. A. Bessonov, T. V. Basova, V. G. Kiselev, L. A. Sheludyakova, N. B. Morozova and I. K. Igumenov, *Vib. Spectrosc.*, 2009, **51**, 283–288.
- 57 H. Tang, D. Fang, Q. Li, P. Cao, J. Geng, T. Sui, X. Wang, J. Iqbal and Y. Du, *J. Food Sci.*, 2012, **77**, T105–T109.
- 58 Q.-Q. Li, Y.-P. Du, Y. Xu, X. Wang, S.-Q. Ma, J.-P. Geng, P. Cao and T. Sui, *Chin. Chem. Lett.*, 2013, **24**, 332–334.
- 59 H. Häkkinen, *Nat. Chem.*, 2012, **4**, 443–455.
- 60 D. Santamaría-Pérez, D. Daisenberger, J. Ruiz-Fuertes, T. Marqueño, R. Chulia-Jordan, C. Muehle, M. Jansen, P. Rodriguez-Hernandez, A. Muñoz, E. R. Johnson and A. Otero-de-la-Roza, *Chem. Sci.*, 2019, **10**, 6467–6475.
- 61 W. Xu, J. Xiao, Y. Chen, Y. Chen, X. Ling and J. Zhang, *Adv. Mater.*, 2013, **25**, 928–933.
- 62 T.-H. Kim, K.-B. Lee and J.-W. Choi, *Biomaterials*, 2013, **34**, 8660–8670.
- 63 V. Van Cat, N. X. Dinh, L. T. Tam, N. V. Quy, V. N. Phan and A.-T. Le, *Mater. Today Commun.*, 2019, **21**, 100639.
- 64 W. Chen, F. Long, G. Song, J. Chen, S. Peng and P. Li, *J. Raman Spectrosc.*, 2020, **51**, 611–618.
- 65 S. Kim, W. Choi, D. J. Kim, H. S. Jung, D.-H. Kim, S.-H. Kim and S.-G. Park, *Nanoscale*, 2020, **12**, 12942–12949.

