An azo-coupling reaction-based surface enhanced resonance Raman scattering approach for ultrasensitive detection of salbutamol

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To date, great achievements with GC-MS, HPLC-MS, and fluorescence biosensing techniques have been made to detect illegal additives of salbutamol (SAL) in swine meat. However, these methods are not suitable for rapid on-site screening due to either costly instruments or rather complicated and/or time consuming sample pretreatments. Herein, a simple, rapid and ultrasensitive approach based on an azo-coupling reaction and surface-enhanced resonance Raman scattering (SERRS) is presented. By combining with a magnetic SERS substrate, an indirect detection for SAL, with a LOD of $1.0 \times 10^{-11} \text{ M}$ ($2.39 \text{ pg mL}^{-1}$), was realized. Moreover, a colorimetric method for naked eye detection was successfully carried out for rapid screening of SAL in concentrations higher than $2.09 \times 10^{-5} \text{ M}$ ($5 \text{ pg mL}^{-1}$). In addition, the proposed method was successfully applied for the rapid determination of SAL in real swine meat. The entire process, including pretreatment, coupling reaction and SERRS detection, was performed within 7 min. Moreover, the SERRS fingerprint band being specific to corresponding functional group guarantees the selectivity for the target molecule. Therefore, the proposed strategy in the present study offers a new way to identify trace amounts of analytes, such as SAL as well as other illegal additives in health-related products and food.

Introduction

Salbutamol (SAL) is often used not only as a drug to treat bronchial asthma and smooth muscle but also a type of β-adrenergic agonist to promote protein synthesis in animal body. It can accelerate the transformation and the decomposition of fat, and is therefore often added to animal feed and easily accumulates in animal body as a residue. Upon consuming animal derived food, humans may suffer from intensified breathing, headache, chest tightness, heart palpitations, increased heart rate, muscle tremors, nausea and other symptoms of poisoning. Therefore, according to the Bulletin 235 issued by the Ministry of Agriculture of China (MAC), SAL is a strictly prohibited substance and should not be detected in animal meat.

To date, great efforts have been made to detect SAL with high sensitivities. For this, gas chromatography-mass spectrometry (GC-MS), liquid chromatography mass spectrometry (LC-MS), ultra-high-performance liquid chromatography (UHPLC) coupled to quadrupole-linear ion trap tandem mass spectrometry (MS/MS), surface molecularly imprinted polymers, electro-chemical analysis, capillary electrophoresis, immunochromatographic assay (ICA), fluorescence biosensor, etc. have been reported. However, these methods need either costly instruments or rather complicated, and/or time consuming sample pretreatments. Therefore, it is urgently needed to develop a simple, rapid and ultrasensitive approach for SAL detection.

Surface enhanced Raman scattering (SERS) is a fascinating technique where the Raman signal of the analyte absorbed on a rough metallic surface can be amplified tremendously even for single molecule detection. SERS has gained a lot of attention on account of its non-destructivity, high selectivity, ultrasensitivity, reliability and fast operation in testing chemical and/or biological molecules in trace amounts. For SERS-based SAL detection, a principal challenge is the low binding affinity of these phenolic molecules on silver or gold surfaces, resulting in poor SERS signals when the analytes are mixed with SERS-active nanoparticles directly (Fig. S1†). To date, several indirect methods such as lateral flow immunochromatographic assay, competitive immunochromatographic test and competitive immunoassay, combined with highly sensitive SERS have been developed to detect SAL. However, these
immunochemical methods require complicated preparation of antibody, are costly, and especially, the determination conditions are highly strict. Particularly, when the concentration of SAL is very low, even false results may be obtained, thus greatly restraining their practical application. Therefore, developing a new technique for sample pretreatment that allows cheap, simple, rapid, selective, and sensitive modification on SAL for SERS detection, is pivotal and a great challenge.

Pauly’s reaction is also known as the azo-coupling reaction, where the diazonium ion couples with either a phenol, amine or hetero-cyclic compound, finally producing azo dyes.26 Pauly’s reagent has been reported to be used for the determination of imidazole or phenol-containing molecules,27 based on the optical absorption of the azo products. Their propensity toward binding to silver nanoparticles (AgNPs) as substrates makes it possible to supply a strong Raman signal for the analytes.28 Therefore, by combining with the surface-enhanced resonance Raman scattering (SERRS) technique,29,30 we will propose an azo-coupling reaction-based SERRS approach for SAL determination. In comparison to the antibody-based immunoprobe, this method will avoid the use of expensive biological reagents and a tedious and time-consuming process for sample pretreatment. By using just a simple chemical reaction with color change, SAL measurement in high sensitivity and selectivity can be made possible. Moreover, in order to improve the test efficiency, magnetic nanoparticles (Fe3O4@Ag, Fig. S2†) were adopted to enrich the sample and reagent C were rapidly added into with manual vortex reaction for 1 min at room temperature. A mixture of 20 mM 2,2’-benzidinedisulfonic acid and 12 M HCl (0.1 mL) in 10 mL solution was used for reagent A; reagent B was 0.8% NaNO2, and reagent C was 4% Na2CO3 in water. The coupling reaction was performed by mixing reagents A, B, C, and SAL together. For that, reagent B was first added into reagent A, and then the sample and reagent C were rapidly added into with manual vortex reaction for 1 min at room temperature. After that, the obtained solution was mixed with a solution containing Fe3O4@Ag (50 mg mL−1, 20 μL) for Raman spectral measurement.

Azo-coupling reaction
Reagents A, B and C were prepared as follows, respectively, and then stored at 4 °C ready for use. A mixture of 20 mM 2,2’-benzidinesulfonic acid and 12 M HCl (0.1 mL) in 10 mL solution was used for reagent A; reagent B was 0.8% NaNO2, and reagent C was 4% Na2CO3 in water. The coupling reaction was performed by mixing reagents A, B, C, and SAL together. For that, reagent B was first added into reagent A, and then the sample and reagent C were rapidly added into with manual vortex reaction for 1 min at room temperature. After that, the obtained solution was mixed with a solution containing Fe3O4@Ag (50 mg mL−1, 20 μL) for Raman spectral measurement.

SERS detection for real samples
The process of detection of SAL from homogenized swine meat was composed of the following three steps: SAL extraction, coupling reaction and SERS detection. (1) SAL extraction: 1.0 g SAL and 5 mL PBS were mixed in a centrifuge tube and scrolled in a vortex for 2 min, then it was filtered by a microporous aperture membrane (0.45 μm) to obtain the filtrate for further analysis. (2) Coupling reaction: the diazonium ion of aromatic amine was synthesized by adding 0.8% NaN02 (200 μL) into 20 mM aromatic amines (200 μL), then the extract of SAL (200 μL) and 4% Na2CO3 (200 μL) were added to the solution containing
diazonium ions in a centrifuge tube (1.5 mL), finally followed by a manual vortex reaction for 1 min at room temperature. (3) SERRS detection: the particles of Fe₃O₄@Ag (50 mg mL⁻¹, 20 µL) were dispersed in the above solution and scrolled 1 min for full adsorption of azo dyes, and were then separated and enriched by a 16 hole magnetic rack. SERRS measurement of the SAL-derived azo dye was performed by using a BWTEK Raman spectrometer with a portable optical fiber probe and 532 nm excitation.

Characterizations
The morphology of magnetic particles was characterized using a JEOL JSM-6700F field emission scanning electron microscope (FE-SEM) operating at 3.0 kV. High performance liquid chromatography (HPLC) analysis was carried out on a Shimadzu LC-20AB system (Kyoto, Japan) equipped with a CTO-10AS column oven and an RF-10Axl fluorescence detector. The chromatographic separation of analytes was performed on a VP-ODS C18 column (5 µm, 4.6 mm x 150 mm i.d.). The mobile phase used in HPLC was acetonitrile/water (0.1% H₃PO₄, pH 2.5) = 25/75. Its flow rate was fixed at 1.0 mL min⁻¹, and SAL the excitation wavelength was set at 276 nm and the emission wavelength was 310 nm. The injection volume was 20 µL for each performance. UV-Vis absorption spectra were taken at 200–800 nm using a Shimadzu UV-2550 spectrometer. The SERS spectrum of the analyte was recorded using a BWTEK Raman spectrometer equipped with different excitation wavelengths, finally fixed at 532 nm after optimization. All the Raman spectra were collected with a laser power of 5.00 mW, while an integration time of 5 s and an accumulation of 5 times was selected for each spectrum.

Results and discussions
Coupling reaction on SAL
SAL has a low cross section for Raman scattering and also is not easily adsorbed on the substrate, so direct SERS detection on SAL is a critical challenge for samples dissolved in solution. Owing to the structural characteristic of SAL as a phenolic compound, an indirect strategy was designed for detection to convert it into an azo dye, which fortunately supports high SERS activity.26,29 The procedure follows the Pauly reaction (Fig. 1), which can be summarized briefly as: the −NH₂ group of the agent interacted with the freshly prepared HNO₂ (originated from NaNO₂ and HCl) to generate the diazonium ions (−N≡N⁺) under acidic conditions; which was then reacted with SAL in basic solution and adjusted by adding Na₂CO₃. According to the mechanism of the coupling reaction, SAL generates azo dye at the ortho-position of phenolic hydroxyl by electrophilic substitution. This procedure is fast and can be completed within 1 min.

Selection of Pauly reagent for SAL
Then, to investigate the resonance absorption of the target molecules at excitation light, SAL-derived azo dyes were obtained by adding different aromatic amines as (1) sulfanilic acid and (2) 2,2'-benzidineisulfonic acid (Fig. 2). As exhibited in the photographs of Fig. 2, both processes showed obvious color reaction supplying orange solution finally, although the intermediate diazonium salts were colorless (inset of Fig. 2). In addition, neither product of diazonium salt showed a UV-Vis absorption peak at 450 nm (green line), indicating that there was no interference of it on the subsequent Raman detection. However, when using (2) 2,2'-benzidineisulfonic acid as a derivative agent, the absorbance strength of the azo dyes was relatively higher than that of (1) sulfanilic acid (pink line) under identical concentrations. Therefore, finally (2) 2,2'-benzidineisulfonic acid is selected as the derivative agent for SAL to provide a more sensitive SERRS response in the following process.

The UV-Vis absorption spectra of SAL, diazonium salt (blank), SAL-derived azo dyes and Fe₃O₄@Ag are illustrated respectively in Fig. 3A, which indicate neither SAL or diazonium salt has an absorption band in the visible range except the SAL-derivative shows a new band at about 450 nm. So, neither of three will interfere with the detection of SAL-derived azo dyes even when the SAL, diazonium salt and/or derivative reagent are largely in excess. Fortunately, the product of the Pauly reaction, azo compound exhibits both a high SERS cross-section and an absorption band centered at 450 nm. As it is very close to the localized surface plasmon resonance (LSPR) absorption of Fe₃O₄@Ag nanoparticles (415 nm), it is possible to be resonated by excitation to the Raman signal. Therefore, the Fe₃O₄@Ag-based SERRS probe for the ultrasensitive and selective indirect detection of SAL is established by detecting SAL-derived azo dyes. Consequently, the SERRS spectra of diazonium salt (black) and SAL-derived azo dyes (red) are shown and compared in
azo dyes are located at 1117, 1160, 1375, 1420, 1591 cm\(^{-1}\) shown in Fig. 5A, the main characteristic peaks of SAL-derived azo dye, based on an azo-coupling reaction. As shown, the best enhanced SERRS signal is obtained at the excitation of 514 nm because of the extensive localized surface plasmon resonance (LSPR) absorption. However, in considering the stronger fluorescence interference induced by 514 nm, finally 532 nm is chosen as the excitation wavelength in the following detection.

**Selection of excitation wavelength**

It has been demonstrated that the SERRS enhancement factor is closely related to the plasmon resonance of the nanoparticle, laser excitation wavelength, and the molecular resonance.\(^{35}\) If the target molecule contains an absorption band overlapping with the laser excitation, an additional order of magnitude in Raman signal will be achieved.\(^{36,37}\) The SERRS spectra of the azo compound under different incident laser wavelengths of 514, 532, 633, 785 nm are shown in Fig. 4A. By comparing the SERRS intensities, which are close to each other under the excitation wavelengths of 514 and 532 nm, we see that both are relatively higher than the other two wavelengths. Fig. 4B records the histogram of the SERRS peaks intensity at 1591 cm\(^{-1}\) of SAL-derivative under different excitation. As shown, the best enhanced SERRS signal is obtained at the excitation of 514 nm because of the extensive localized surface plasmon resonance (LSPR) absorption. However, in considering the stronger fluorescence interference induced by 514 nm, finally 532 nm is chosen as the excitation wavelength in the following detection.

**Sensitivity of SAL detection**

The indirect SERS analysis is performed after target SAL is changed to azo dye, based on an azo-coupling reaction. As shown in Fig. 5A, the main characteristic peaks of SAL-derived azo dyes are located at 1117, 1160, 1375, 1420, 1591 cm\(^{-1}\), being attributed to \(v(C-N)\), \(v(C-C)\) from phenol groups, \(\delta(C-H)\), \(\nu(N=N)\), and \(\nu(C=C)\) from phenyl rings, respectively.\(^{28,34}\) By comparing all curves, the existence of Raman peaks for SAL even at a concentration as low as 1.0 \(\times\) \(10^{-11}\) M is speculated, which confirms that this method can be used to achieve trace detection of SAL. Fig. 5B shows the calibration curves by plotting the SERRS intensities of SAL at 1591 cm\(^{-1}\) versus the log concentration of SAL. Additionally, the color of SAL-derived azo dyes in solution is diminished and became colorless (or light yellow) with the concentration decrease of SAL, as shown in Fig. S3.\(^{†}\) Therefore, we can determine the concentrations of SAL by naked eyes based upon the color changes of the solution when it is higher than 2.09 \(\times\) \(10^{-5}\) M (5 \(\mu\)g mL\(^{-1}\)). However, when the concentration of SAL is less than 5 \(\mu\)g mL\(^{-1}\), there is no obvious color difference among the different concentrations of SAL. To improve the sensitivity of this azo-coupling reaction, Raman spectrometer is required.

To confirm the reproducibility of the proposed SERRS method, the standard solutions of SAL from 10 batches under the identical concentrations of 1.0 \(\times\) \(10^{-10}\) M, were tested. For each sample, the SERRS spectrum and its histogram are shown in Fig. S4.\(^{†}\) which are obtained on randomly selected spots on the substrate, respectively. By comparing to the sample-to-sample variation at 1591 cm\(^{-1}\) in the SERRS spectra, remarkable reproducibility is exhibited.

**Detection of SAL in real samples**

To further demonstrate the practical application of the azo-coupling reaction based SERRS method for SAL detection in real samples, we purchased 5 packages meat randomly from 5 different shops in farm-product markets, in numbering as #1–5. These samples need pretreatments including liquid–solid extraction, coupling reaction and SERRS detection in turn. The spike-and-recovery results are listed in Table 1. None of the original sample is evaluated to contain SAL as tested by SERRS and HPLC (chromatograms and standard curves are shown in Fig. S5.\(^{†}\)). Then, the original samples are spiked SAL to 1.0 \(\times\) \(10^{-6}\) M by using its standard solutions for subsequent extraction and detection, and the results show recoveries in the range of 78–117%, demonstrating the practical application of the azo-coupling reaction based SERRS method and confirming it would be applicable to the rapid detection of SAL in homogenized swine meat. The detection limit in real sample is displayed in Fig. S6.\(^{†}\) which can be estimated as 1.0 \(\times\) \(10^{-10}\) M. In comparison to other reported methods for SAL detection (Table S1.\(^{†}\)), the present study exhibits significant advantages to realize rapid and ultra-sensitive detection.
also other illegal additives involved in health products and food, when choosing the corresponding Raman band as specific.

Interference assay for SAL detection

In order to evaluate anti-interference of the assay to SAL detection, different possible interferences such as clenbuterol, lysine, glycine, glycerol, KCl and NaCl at a concentration of 1.0 × 10⁻³ M, is mixed with SAL (1.0 × 10⁻⁶ M) under the optimal condition. After mixing with magnetic particles, each of them is tested based on the proposed azo-coupling SERRS method. The results show achromatic colour only for the control, while orange for all other solutions containing SAL (Fig. 6A), exhibiting no effect of either on the detection of SAL. Similarly, Fig. 6B gives the SERRS intensity of the SAL derivative and different interference after adding into the magnetic particles as above. It shows a stable intensity at 1591 cm⁻¹, indicating that the mixture of SAL with different interference does not affect the detection of SAL.

Conclusions

In summary, aiming to rapidly and ultrasensitively detect illegal SAL in swine meat, we propose a new approach in the present study based on an azo-coupling reaction and surface enhanced resonance Raman scattering (SERRS). By combining with a magnetic SERS substrate, as well for separation at the same time, an indirect detection on SAL in solution with a LOD of 1.0 × 10⁻¹¹ M by SERRS and 2.09 × 10⁻⁵ M by naked eye is realized firstly. In addition, with improvement the proposed method is successfully applied to the determination of SAL in real swine meat and offers a simple, practicable, high sensitive and high anti-interference approach. Particularly, the proposed strategy offers a new opportunity for trace amounts of not only SAL but

Table 1: Application of azo-coupling reaction based SERRS method for the determination of SAL in homogenized swine meat

<table>
<thead>
<tr>
<th>Sample no.</th>
<th>Detection by SERRS and HPLC before spiking</th>
<th>Spike value (x 10⁻⁶ M)</th>
<th>Detection by SERRS (x 10⁻⁶ M)</th>
<th>Recovery (%)</th>
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<td>105</td>
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</tbody>
</table>

Fig. 6 (A) The photographic images and (B) the SERRS intensity of SAL (1.0 × 10⁻⁴ M) and in mixing with the magnetic particles and different interference (1.0 × 10⁻³ M) as A: clenbuterol, B: lysine, C: glycine, D: glycerol, E: KCl and F: NaCl, respectively.

Conflicts of interest

There are no conflicts between the authors to declare.

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Notes and references