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# Magnetic solid-phase extraction of pyrethroid insecticides from tea infusions using ionic liquid-modified magnetic zeolitic imidazolate framework-8 as an adsorbent

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A simple, sensitive, and reliable magnetic solid-phase extraction (SPE) method coupled with GC-MS/MS for the effective analysis of four pyrethroids from tea infusions was developed. A magnetic adsorbent, named ionic liquid-modified magnetic zeolitic imidazolate framework-8 ( $\text{Fe}_3\text{O}_4/\text{ZIF-8}/\text{IL}$ ), was prepared by immobilizing an ionic liquid (IL) on the surface of  $\text{Fe}_3\text{O}_4/\text{ZIF-8}$ . The textures of  $\text{Fe}_3\text{O}_4/\text{ZIF-8}/\text{IL}$  were confirmed by material characterization, and the results suggested that the adsorbent possessed high magnetism ( $59.0 \text{ emu g}^{-1}$ ), an adequate Brunauer–Emmett–Teller (BET) surface area ( $104 \text{ m}^2 \text{ g}^{-1}$ ), and a large pore volume ( $0.68 \text{ cm}^3 \text{ g}^{-1}$ ). To confirm the extraction performance of the prepared  $\text{Fe}_3\text{O}_4/\text{ZIF-8}/\text{IL}$ , several experimental conditions affecting the extraction efficiency were investigated. Under the optimum conditions, the limits of determination (LODs) for the four pyrethroids were in the range of  $0.0065\text{--}0.1017 \mu\text{g L}^{-1}$  ( $S/N = 3 : 1$ ) with an intra-day relative standard deviation (RSD) of  $\leq 9.70\%$  and inter-day RSD of  $\leq 11.95\%$ . The linear ranges were  $0.5\text{--}50 \mu\text{g L}^{-1}$  for bifenthrin and  $0.5\text{--}500 \mu\text{g L}^{-1}$  for permethrin, cypermethrin, and flucythrinate, with determination coefficients higher than 0.999. Finally, the proposed technique was successfully applied for the determination of pyrethroids in real tea infusions. This work could be extended to other IL-modified metal–organic frameworks (MOFs) and to the development of different sample pretreatment techniques.

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## 1. Introduction

Tea is one of the most widely consumed drinks in the world since it has excellent refreshing and mild stimulant effects on and even medicinal benefits for humans.<sup>1</sup> Tea can be classified as green, white, oolong, black, or Pu-erh and is usually brewed before consumption.<sup>2,3</sup> Because of the high humidity and warm temperatures in the environments in which tea is cultivated, tea plantations suffer from pest damage and consequent economic loss. Therefore, insecticides are commonly used to control pests on this crop. Pyrethroids, as the third most commonly applied group of insecticides, are widely used for pest control in agricultural and non-agricultural fields because of their high activities and low mammalian toxicities.<sup>4</sup> However, because of their excessive application and accumulation over time,

pyrethroids can enter into the human body through the food chain, creating a substantial health risk.<sup>5</sup> Accordingly, several international organizations have set maximum residue levels for pyrethroids in tea, such as  $0.2\text{--}30 \text{ mg kg}^{-1}$  in the United States and  $0.01\text{--}3 \text{ mg kg}^{-1}$  in the European Union.<sup>6</sup> Therefore, the development of a sensitive and reliable analytical technique for monitoring pyrethroid insecticides in tea is of prime importance.

Sample pretreatment is a crucial component in the determination of pesticide residues in complicated matrices. Numerous sample preparation methods have been established for the analysis of pyrethroids, including liquid–liquid extraction, liquid–liquid microextraction, solid-phase extraction (SPE), solid-phase microextraction, and QuEChERS.<sup>7–11</sup> Magnetic SPE (MSPE), as an alternative to SPE, has gained extensive recognition since its introduction in 1999.<sup>12</sup> In this technique, a magnetic adsorbent is added to a sample solution and dispersed with the aid of a vortex, shaker, or sonicator. Then, an external magnetic field is used to facilitate efficient retrieval of the magnetic adsorbent.<sup>13</sup> This extraction process is rapid and simple because of the magnetism of the magnetic adsorbent, which means that phase separation can be readily performed without tedious or time-consuming filtration or

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centrifugation steps.<sup>14</sup> Recently, many magnetic adsorbents have been prepared for MSPE, such as magnetic molecularly imprinted polymers, magnetic carbon nanomaterials, and magnetic metal–organic frameworks (MOFs).<sup>15</sup>

Zeolitic imidazolate frameworks (ZIFs), a new subclass of MOFs with chemically stable zeolite-like frameworks, are usually formed from metal ions (*e.g.*, Zn and Co) and imidazolate organic linkers.<sup>16</sup> ZIFs are microporous with high surface areas and uniform structured cavities, which endow them with remarkable physical, chemical, and thermal stabilities and excellent adsorption abilities.<sup>17,18</sup> Among the different types of ZIFs, ZIF-8 has attracted widespread attention because of its high surface area, hierarchical porosity, easy preparation and cost-effectiveness. Recently, ZIF-8 has been widely employed in gas storage, drug delivery, sensing, catalysis, and mixed matrix membranes.<sup>19–23</sup> Furthermore, ZIF composites have been prepared by modification with different chemical reagents, such as chitosan, thiourea, polydopamine, and enzymes, to enhance their adsorption abilities and confer other novel properties.<sup>24–27</sup>

Ionic liquids (ILs), produced by associating large organic cations and organic anions, are a class of liquid molten salts with melting points at room temperature. ILs are regarded as an alternative to organic solvents because of their unique chemical and thermal stabilities, extremely low vapour pressures, non-flammability, and high selectivity.<sup>28</sup> Accordingly, ILs have attracted widespread attention and have been widely used in catalysis, adsorption and separation, and modification.<sup>29–32</sup> According to the literature, ILs can be used in an immobilized or non-immobilized form for separation. Modification of a support material with ILs can increase the extraction efficiency by introducing different interactions between the sorbent and target analyte, including ion-exchange, hydrophobic, electrostatic, hydrogen bonding, and  $\pi$ – $\pi$  interactions.<sup>33,34</sup> Some IL-modified ZIFs have been reported in recent years, and these ZIFs are attractive in many research fields, especially for sample pretreatment.<sup>35</sup> To the best of our knowledge, no ZIF-8/IL-based composite adsorbent has been reported for the analysis of pyrethroids in a tea infusion.

A combination of magnetic ZIF-8 and hydrophobic IL could either promote the adsorption capacity of ZIF-8 for pyrethroids, or facilitate the applicability of IL in MSPE. Inspired by the above aspects, we sought to develop a simple, sensitive and reliable analytical technique using an IL-modified magnetic ZIF as an adsorbent for MSPE of pyrethroid insecticides from tea infusions. In this work, we prepared an IL-modified magnetic ZIF-8 nanocomposite *via* a simple synthesis by coating the IL on the surface of magnetic ZIF-8. Successful preparation of the magnetic nanocomposite was confirmed by material characterization. Furthermore, we optimized several experimental factors for the MSPE method. Finally, we applied the proposed method to the determination of four pyrethroids in tea infusions.

## 2. Materials and methods

### 2.1. Materials

The liquid standards (bifenthrin, permethrin, cypermethrin, and flucythrinate) were at a concentration of 1000 mg L<sup>-1</sup> and

were obtained from the Agro-Environmental Protection Institute, Ministry of Agriculture (Tianjin, China). A mixture of the four pyrethroids was prepared in methanol at a concentration of 100 mg L<sup>-1</sup> and stored at 4 °C in darkness. Acetonitrile, ethyl acetate, acetone, methanol, and *n*-hexane were purchased from Sigma-Aldrich (St. Louis, MO, USA). All of the organic solvents were of HPLC grade. Analytical grade 2-methylimidazole, zinc sulfate heptahydrate, ferric chloride hexahydrate (FeCl<sub>3</sub>·6H<sub>2</sub>O), ferrous chloride tetrahydrate (FeCl<sub>2</sub>·4H<sub>2</sub>O), ammonium hydroxide (mass fraction 28%), and thioglycolic acid were acquired from Aladdin Industrial Co. (Shanghai, China). Analytical grade ILs, including 1-hexyl-3-methylimidazolium hexafluorophosphate, 1-hexyl-3-methylimidazolium bis(trifluoromethanesulfonyl)imide, 1-octyl-3-methylimidazolium hexafluorophosphate, and 1-octyl-3-methylimidazolium bis(trifluoromethanesulfonyl)imide ([OMIM]NTf<sub>2</sub>), were purchased from Shanghai Yuanye Bio-Technology Co. (Shanghai, China). Ethanol, sodium chloride (NaCl), hydrochloric acid (HCl) and sodium hydroxide (NaOH) were of analytical grade and were obtained from Beijing Chemical Reagents Co. (Beijing, China).

### 2.2. Preparation of magnetic ZIF-8/IL nanocomposites

Fe<sub>3</sub>O<sub>4</sub> nanoparticles were synthesized by a co-precipitation method as reported previously.<sup>36</sup> In a typical procedure, 1.8 g of FeCl<sub>3</sub>·6H<sub>2</sub>O and 0.8 g of FeCl<sub>2</sub>·4H<sub>2</sub>O were dissolved in 240 mL of ultrapure water and stirred under a N<sub>2</sub> atmosphere at 80 °C for 30 min. Then, ammonium hydroxide (25%, 10 mL) was added dropwise, and the mixture was stirred for another 30 min. After that, precipitates were collected using an external magnetic field and washed three times each with ultrapure water and ethanol. Subsequently, the obtained precipitates were dried at 60 °C in a vacuum oven for 24 h.

Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 was prepared using a modification of a previously reported method.<sup>37</sup> Briefly, 0.1 g of Fe<sub>3</sub>O<sub>4</sub> nanoparticles were added to 20 mL of an ethanol solution of thioglycolic acid (20 mM) with shaking for 1 h at room temperature. The product was recovered by an external magnetic field and washed three times each with ultrapure water and ethanol. Afterwards, the obtained thioglycolic acid-functionalized Fe<sub>3</sub>O<sub>4</sub> nanoparticles were re-dispersed in 240 mL of ultrapure water/ethanol (1 : 1, v/v) and shaken for another 1.5 h at room temperature. Then, an ethanol solution (20 mL) of 0.84 g of 2-methylimidazole was added to the mixture, followed by stirring for 8 h at room temperature. Finally, the Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 product was retrieved by an external magnetic field and washed three times each with ultrapure water and ethanol. The final product was dried at 60 °C in a vacuum oven for 24 h.

The Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL nanocomposite was prepared by immobilizing an IL on the surface of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 *via* a simple direct immersion method based on the “physisorption” mechanism, and the entire process followed a modification of a previously reported method.<sup>38</sup> Briefly, an IL (1 g) was added to a centrifuge tube containing 6 mL of ethanol, followed by vortexing for 30 s. After dispersion, different quantities of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 (0.5, 0.6, 0.8, 1, 1.2, and 1.5 g) were transferred into the solution, which was kept under mechanical stirring for 30 min at room temperature. Then, the products were collected with an external



magnet and washed three times with ultrapure water and ethanol. Finally, the prepared products were dried at 60 °C in a vacuum oven for 24 h.

### 2.3. Characterisation of magnetic ZIF-8/IL nanocomposites

Transmission electron microscopy (TEM) and scanning electron microscopy (SEM) were conducted using a JEM-200CX microscope (JEOL, Tokyo, Japan) and a JSM-6300 microscope (JEOL, Tokyo, Japan), respectively. X-ray powder diffraction (XRD) measurements were performed on a D8 Advance diffractometer (Bruker, Karlsruhe, Germany). Fourier transform infrared (FT-IR) spectra were recorded on an FT-IR-8400 spectrometer (Shimadzu, Kyoto, Japan). Vibrating sample magnetometer measurements were conducted on a Versalab-7410 (Lake Shore, Columbus, USA). Nitrogen (N<sub>2</sub>) adsorption and desorption isotherm measurements were performed on an ASAP2460 surface area and porosity analyser (Micromeritics, Norcross, USA).

A Shimadzu GC-2010 plus equipped with a Shimadzu TQ8040 triple-quadrupole MS (Shimadzu, Kyoto, Japan) was applied for the analyses. An Rtx-5Ms capillary column from RESTEK (Bellefonte, PA, USA, 30 m × 0.25 mm i.d., 0.25 μm film thickness) was used for separation. The injector temperature was 250 °C. The carrier gas was helium at a constant flow rate of 1 mL min<sup>-1</sup>. The column temperature was initially maintained at 40 °C for 4 min, then increased to 125 °C at 25 °C min<sup>-1</sup>, increased to 300 °C at 10 °C min<sup>-1</sup>, and held at 300 °C for 6 min. The injection volume was 1.0 μL in splitless mode. The specific multiple reaction monitoring (MRM) transitions for all four pyrethroids and other parameters are given in Table 1.

### 2.4. Preparation of tea infusions

Various types of tea, such as Pu-erh, Tieguanyin, Da Hong Pao, and Sichuan Maofeng, were purchased from a local market. Tea infusions were prepared according to an established method.<sup>39</sup> Briefly, a tea sample (2 g) was infused in boiling water (100 mL) for 5 min. After that, the infusion was transferred through a funnel to a glass beaker. Finally, the tea infusion was filtered through a 0.45 μm polytetrafluoroethylene membrane filter and transferred to a dark-amber glass bottle. Method validation was performed by analysing ultrapure water and a blank tea infusion spiked with the standard mixture. All samples were stored at 4 °C before analysis.

### 2.5. MSPE procedure

The extraction of the four pyrethroids was studied using the obtained Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL nanocomposite as a sorbent. First, 10 mg of sorbent was dispersed into 10 mL of the sample solution (NaCl, 5% mass fraction) in a 15 mL centrifuge tube and vortex-mixed for 1.5 min to perform the extraction. After that, the black sorbents were gathered by a magnet, and the clear supernatant was discarded. Subsequently, 0.8 mL of acetonitrile was added to elute the analytes from the magnetic sorbent under vortex mixing for 1 min. After the magnetic sorbent was retrieved, the eluent was transferred to another 5 mL centrifuge tube. The same desorption experiment was conducted again. Finally, the desorption solvents were combined and evaporated to dryness under a gentle stream of N<sub>2</sub> at 40 °C. After that, the residue was redissolved in 0.5 mL of acetone, and a 1.0 μL aliquot was injected into the GC-MS/MS system for analysis.

## 3. Results and discussion

### 3.1. Characterization of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL

The morphology of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL was confirmed by SEM and TEM. Fig. 1A shows that the Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL nanoparticles were in nanoscale, with slight aggregation. Fig. 1B shows a highly porous block-shaped structure for the Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL nanocomposites. This porous structure could be good for the promotion of the ability of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL nanocomposites to adsorb pyrethroids.

FT-IR spectroscopy was used to study the surface properties of the prepared magnetic materials, including Fe<sub>3</sub>O<sub>4</sub>, Fe<sub>3</sub>O<sub>4</sub>/ZIF-8, and Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL.<sup>40</sup> For all three synthetic magnetic materials (Fig. 1C), the characteristic Fe–O vibration at 558 cm<sup>-1</sup> indicates the successful incorporation of Fe<sub>3</sub>O<sub>4</sub> into the ZIF-8 nanocomposites.<sup>41</sup> The peaks between 993 and 1421 cm<sup>-1</sup> were assigned to the Zn–N stretching vibration of ZIF-8. The vibration of the C–H bond of the alkyl groups at the nitrogen atoms of the imidazole ring can be observed from the bands at 878, 2825 and 2960 cm<sup>-1</sup>. The peaks at 1059 and 1138 cm<sup>-1</sup> can be attributed to the asymmetric and symmetric stretching of O=S–O<sup>-</sup>, whereas those at 1197 and 1350 cm<sup>-1</sup> were assigned to the asymmetric and symmetric stretching of O=S=O.<sup>39</sup> These results may indicate the feature structure of IL.

Table 1 Acquisition and chromatographic parameters for four pyrethroids

Triazoles	<i>t</i> <sub>R</sub> (min)	MRM1 ( <i>m/z</i> )	CE1 <sup>a</sup> (eV)	MRM2 ( <i>m/z</i> )	CE2 (eV)
Bifenthrin	18.001	181.10 > 166.10	12	181.10 > 153.10	8
Permethrin-1	19.777	183.10 > 168.10	14	183.10 > 165.10	14
Permethrin-2	19.902	183.10 > 168.10	14	183.10 > 165.10	14
Cypermethrin-1	20.630	163.10 > 127.10	6	163.10 > 91.00	14
Cypermethrin-2	20.733	163.10 > 127.10	6	163.10 > 91.00	14
Cypermethrin-3	20.793	163.10 > 127.10	6	163.10 > 91.00	14
Cypermethrin-4	20.831	163.10 > 127.10	6	163.10 > 91.00	14
Flucythrinate-1	20.794	199.10 > 157.10	10	199.10 > 107.10	22
Flucythrinate-2	20.985	199.10 > 157.10	10	199.10 > 107.10	22

<sup>a</sup> CE means collision energy.



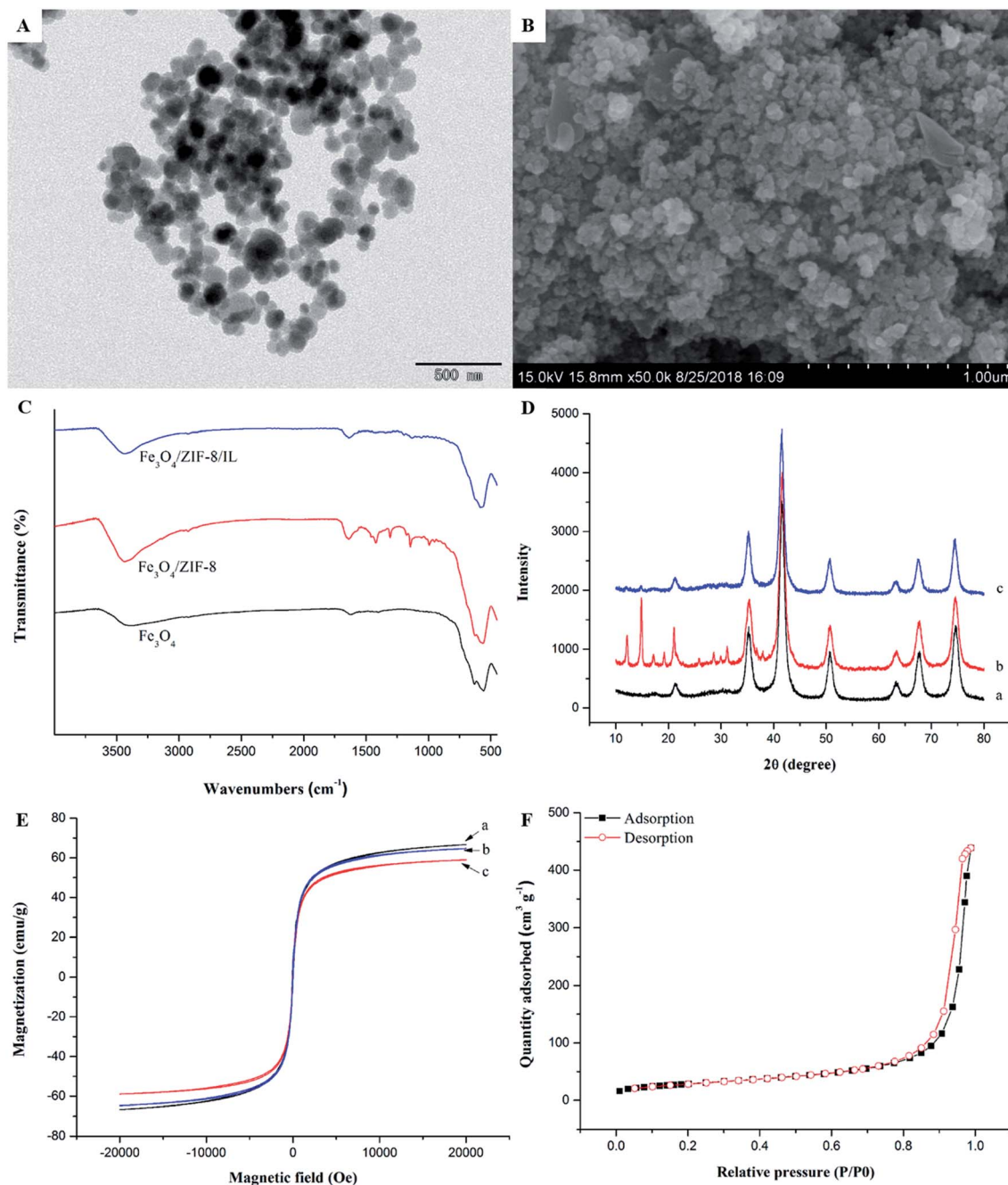


Fig. 1 Characterization of magnetic materials: (A) TEM image and (B) SEM image of  $\text{Fe}_3\text{O}_4/\text{ZIF-8}/\text{IL}$ ; (C) FT-IR spectra of magnetic materials; (D) XRD patterns of (a)  $\text{Fe}_3\text{O}_4$ , (b)  $\text{Fe}_3\text{O}_4/\text{ZIF-8}$ , and (c)  $\text{Fe}_3\text{O}_4/\text{ZIF-8}/\text{IL}$ ; (E) magnetic hysteresis loops of (a)  $\text{Fe}_3\text{O}_4$ , (b)  $\text{Fe}_3\text{O}_4/\text{ZIF-8}$ , and (c)  $\text{Fe}_3\text{O}_4/\text{ZIF-8}/\text{IL}$ ; and (F)  $\text{N}_2$  adsorption–desorption isotherms of  $\text{Fe}_3\text{O}_4/\text{ZIF-8}/\text{IL}$ .

The synthetic magnetic nanocomposite was also characterized by XRD. As shown in Fig. 1D, the crystal structure of ZIF-8 can be confirmed by the characteristic peaks at 10.0, 12.2, 14.9, 17.1, 19.2, 25.8 and 28.5°, and the  $\text{Fe}_3\text{O}_4$  crystal structure contains the typical peaks at 35.3, 41.6, 50.6, 67.7 and 74.6°. Comparison of the XRD patterns of  $\text{Fe}_3\text{O}_4/\text{ZIF-8}$  and  $\text{Fe}_3\text{O}_4/\text{ZIF-8}/\text{IL}$  revealed that the diffraction peaks of the latter were slightly weaker than those of the former. This observation indicates that  $\text{Fe}_3\text{O}_4/\text{ZIF-8}/\text{IL}$  exists as an aggregate and is coated with the hydrophobic IL.

The magnetic properties of the synthetic materials were investigated using a vibrating sample magnetometer (Fig. 1E). The saturation magnetization value of  $\text{Fe}_3\text{O}_4$  was 66.7  $\text{emu g}^{-1}$ . After the  $\text{Fe}_3\text{O}_4$  nanoparticles were coated with ZIF-8, the saturation magnetization value of  $\text{Fe}_3\text{O}_4/\text{ZIF-8}$  decreased slightly to 64.7  $\text{emu g}^{-1}$ . After modification with the IL, the saturation magnetization value of  $\text{Fe}_3\text{O}_4/\text{ZIF-8}/\text{IL}$  decreased further to 59.0  $\text{emu g}^{-1}$ . The magnetic hysteresis loops of the three prepared magnetic materials showed that both their remanence and coercivity values were zero. All of these results suggest that the



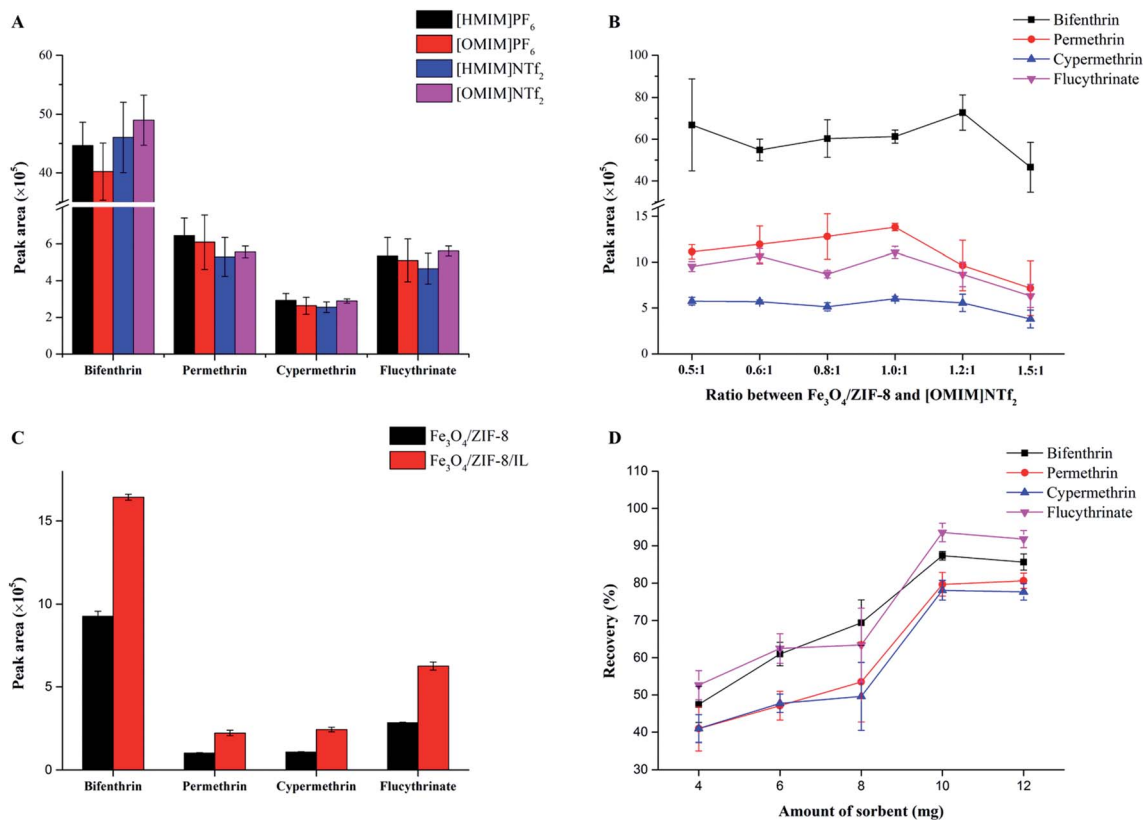


Fig. 2 Effect of (A) different types and (B) different mass ratios of ILs on the Fe<sub>3</sub>O<sub>4</sub>/ZIF-8; [HMIM]PF<sub>6</sub>, 1-hexyl-3-methylimidazolium hexafluorophosphate; [OMIM]PF<sub>6</sub>, 1-octyl-3-methylimidazolium hexafluorophosphate; [HMIM]NTf<sub>2</sub>, 1-hexyl-3-methylimidazolium bis(trifluoromethanesulfonyl)imide; (C) comparison of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 and Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL in the extraction of pyrethroids from ultrapure water; (D) effect of the sorbent mass on the extraction efficiencies of pyrethroids from ultrapure water.

prepared Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL is superparamagnetic and can be rapidly separated by an external magnetic field.

The pore properties of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL were studied by N<sub>2</sub> adsorption measurements at 77 K. As shown by the N<sub>2</sub> adsorption-desorption isotherm of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL in Fig. 1F, the Brunauer-Emmett-Teller (BET) surface area was 104 m<sup>2</sup> g<sup>-1</sup>, and the total pore volume was 0.68 cm<sup>3</sup> g<sup>-1</sup>. The pore size was also calculated by the Barrett-Joyner-Halenda method, and the distribution range was 1.7 to 300 nm, indicating the coexistence of mesopores and macropores.<sup>43</sup> These results suggest that Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL has a large surface area and pore volume, which are beneficial for facilitating the adsorption of pyrethroids.

### 3.2. Optimization of the extraction conditions

Next, we used single-factor experiments to investigate several experimental conditions affecting the extraction performance of MSPE of pyrethroids. These conditions were the type and content of the IL, type and mass of sorbent, adsorption time, ionic strength, and pH of the sample solution.

As previously reported, the extraction efficiency of an IL for a pesticide residue depends on the physicochemical characteristics of the IL.<sup>39</sup> Therefore, selection of an appropriate IL and optimization of its content are essential for optimization of the extraction conditions. In the present work, we studied four ILs, 1-hexyl-3-methylimidazolium hexafluorophosphate, 1-hexyl-3-

methylimidazolium bis(trifluoromethanesulfonyl)imide, 1-octyl-3-methylimidazolium hexafluorophosphate, and [OMIM]NTf<sub>2</sub>, as potential reagents for modifying Fe<sub>3</sub>O<sub>4</sub>/ZIF-8. The extraction efficiency was selected to evaluate the effect of modification of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 with the four ILs. The extraction efficiencies of the four IL-modified Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 sorbents for the four pyrethroids were at the same level (Fig. 2A). This result indicates that the length of the alkyl chain of the cation and the type of anion do not greatly affect the extraction efficiencies for the target analytes. However, the results for the [OMIM]NTf<sub>2</sub> treatment group had the best standard deviations (SDs); hence, [OMIM]NTf<sub>2</sub> was adopted as the best IL for Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 modification. Next, we investigated mass ratios of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 to [OMIM]NTf<sub>2</sub> of 0.5 : 1, 0.6 : 1, 0.8 : 1, 1 : 1, 1.2 : 1, and 1.5 : 1 to optimize the IL content in Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL. A mass ratio of 1 : 1 gave the best extraction efficiencies and SDs (Fig. 2B). Therefore, the mass ratio of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 to [OMIM]NTf<sub>2</sub> was set at 1 : 1.

It is well known that adsorbents play a key role in determining extraction efficiency. Consequently, we compared the extraction efficiencies of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 with those of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL. Compared to Fe<sub>3</sub>O<sub>4</sub>/ZIF-8, Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL had better extraction efficiencies for the target analytes (Fig. 2C). This result suggests that modification with [OMIM]NTf<sub>2</sub> can improve the extraction efficiencies of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 for pyrethroids. To achieve satisfactory extraction performance with Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL for



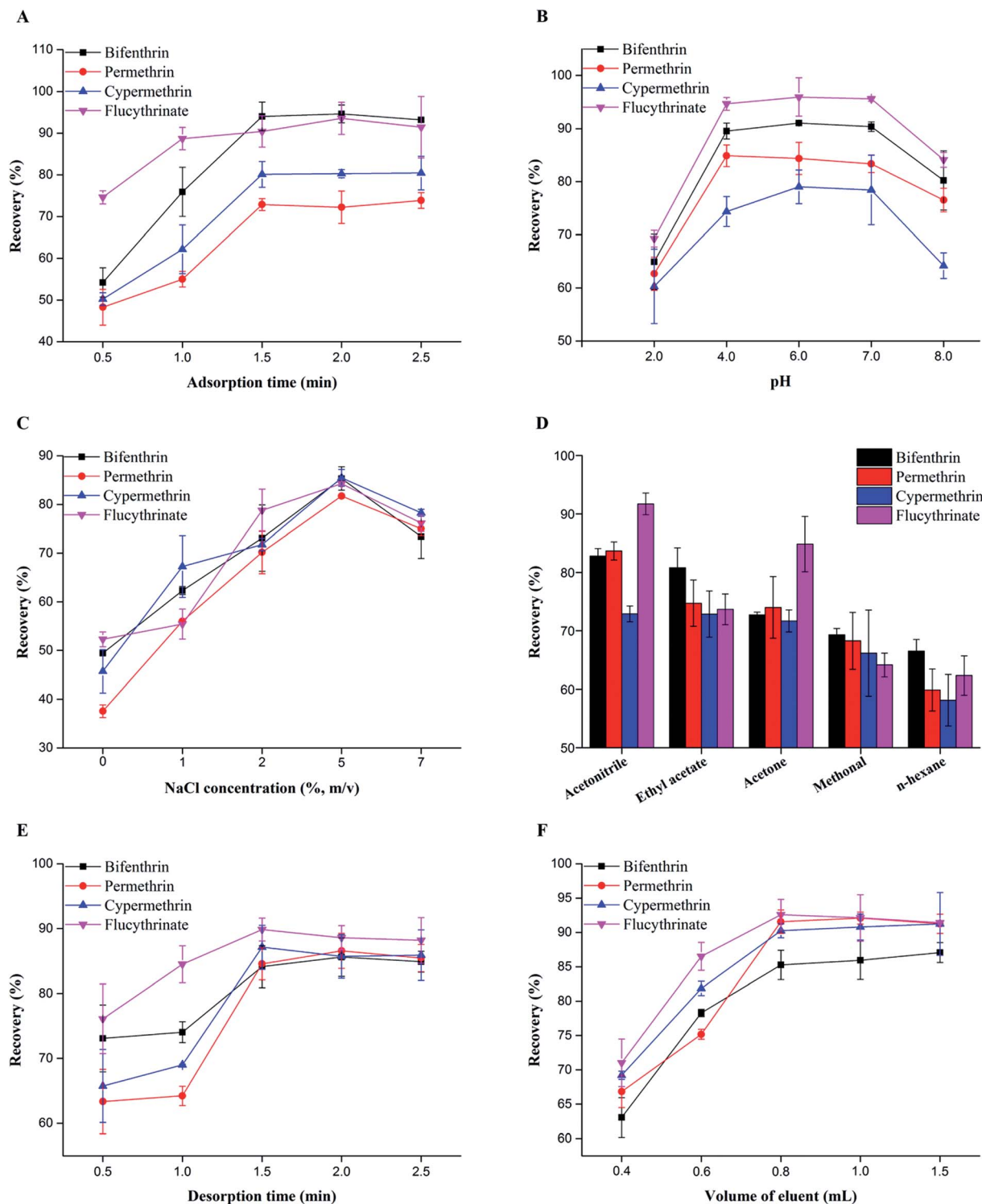


Fig. 3 Effects of different parameters on the MSPE of pyrethroids from ultrapure water: (A) adsorption time, (B) pH of the sample solution, and (C) ionic strength of the sample solution. Desorption conditions: 0.8 mL of acetonitrile; vortex mixing time, 1.5 min; and the same desorption process was repeated one more time. (D) Desorption solvent, (E) desorption time, and (F) eluent volume. Extraction conditions: amount of adsorbent, 10 mg; extraction time, 1.5 min; ionic strength of the sample solution, 5% NaCl.

pyrethroids, we tested a series of  $\text{Fe}_3\text{O}_4/\text{ZIF-8}/\text{IL}$  masses ranging from 4 to 12 mg in 10 mL of sample solution. The extraction efficiency increased as the dose of sorbent was increased from 4

to 10 mg and then remained nearly constant with further increases in the sorbent mass (Fig. 2D). These results show that 10 mg of sorbent is adequate for MSPE of pyrethroids.



Table 2 Analytical parameters of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL as an adsorbent for MSPE of the four pyrethroids from ultrapure water samples

Pyrethroids	Regression equation	Linear range ( $\mu\text{g L}^{-1}$ )	$R^2$	LOD ( $\mu\text{g L}^{-1}$ )	RSD <sup>a</sup> (%) ( $n = 6$ )	
					Intraday	Interday
Bifenthrin	$Y = 1.5 \times 10^8 X + 2414$	0.5–50	0.9991	0.0040	0.69	4.26
Permethrin	$Y = 1.7 \times 10^7 X + 2202$	0.5–100	0.9991	0.0374	2.13	7.67
Cypermethrin	$Y = 1.4 \times 10^7 X - 9587$	0.5–100	0.9998	0.0409	0.88	5.52
Flucythrinate	$Y = 4.0 \times 10^7 X - 20766$	0.5–100	0.9998	0.0367	1.24	5.53

<sup>a</sup> The RSDs were determined by performing six replicate analyses of samples spiked with each pyrethroid at  $10 \mu\text{g L}^{-1}$ .

The extraction time is a nonnegligible factor that affects the adsorption equilibrium between the adsorbent and sample solution. Thus, 0.5–2.5 min of vortexing time was set to study the effect of the adsorption time on the MSPE of the analytes. As illustrated in Fig. 3A, the extraction efficiencies for the four pyrethroids increased as the adsorption time was increased from 0.5 to 1.5 min and then kept relatively constant with further increases in the adsorption time. Based on the above results, 1.5 min was chosen as the adsorption time for MSPE.

The pH of the sample solution can affect MSPE performance by altering the surface charge of the adsorbent and/or chemical form of the analyte. In this study, we adjusted the sample solution pH from 2.0 to 8.0 with HCl or NaOH to investigate the effect of pH on the extraction performance. As shown in Fig. 3B, the extraction efficiencies for analytes remained relatively constant as the sample solution pH from 4.0 to 7.0, but decreased with the sample solution pH adjusted to 2.0 or 8.0. This behaviour was possibly caused by instability of the pyrethroids under highly acidic and alkaline conditions. Because the majority of tea samples are slightly acidic, we did not adjust the pH values of the tea infusions before extraction.<sup>44</sup>

Salt is another crucial parameter that can affect extraction performance. Therefore, we added NaCl (0 to 7%, w/v) to the sample solution to investigate the effect of salt on the extraction efficiencies. As the NaCl content was increased from 0 to 5%, the extraction efficiencies of the sorbent for the analytes also increased and then decreased with further increases in the NaCl content (Fig. 3C). Considering the extraction efficiencies and SDs, we selected 5% (w/v) NaCl as the optimum content for MSPE.

### 3.3. Optimization of the desorption conditions

The desorption procedure is essential for MSPE of pyrethroids. To obtain acceptable desorption performance, we studied the

effects of changing the desorption solvent, desorption time, and eluent volume.

It is well known that the desorption solvent can control the extraction efficiency of MSPE. Accordingly, acetonitrile, ethyl acetate, acetone, methanol, and *n*-hexane were chosen as potential desorption solvents. Among these solvents, acetonitrile gave the best extraction efficiencies for the four pyrethroids (Fig. 3D). Moreover, we investigated changes in the desorption time by increasing the vortex mixing time from 0.5 to 2.5 min. A desorption time of 1.5 min was satisfactory for desorption of pyrethroids from the sorbent (Fig. 3E). Furthermore, the effect of desorption volumes between 0.4 and 1.5 mL was studied, and we found that 0.8 mL provided the optimum extraction efficiencies (Fig. 3F). In summary, the optimum process for eluting the pyrethroids used 0.8 mL of acetonitrile with vortex mixing for 1.5 min and was repeated one more time.

### 3.4. Method validation

The proposed method was characterized under the optimized conditions for linearity, limit of determination (LOD), and precision in both ultrapure water and a tea infusion, and the results are shown in Tables 2 and 3.

The linearity was determined by analysing ultrapure water and a blank tea infusion sample that were spiked with standard solutions of the pyrethroids at  $0.5\text{--}500 \mu\text{g L}^{-1}$  and plotting the peak area *versus* the concentration. For water sample, good linearities for permethrin, cypermethrin, and flucythrinate over the studied concentration range ( $0.5\text{--}100 \mu\text{g L}^{-1}$ ) were observed, with determination coefficients ( $R^2$ ) in the range of 0.9991–0.9998. For the blank tea sample, good linearities with  $R^2$  values ranging from 0.9990 to 0.9999 were obtained over the studied concentration range ( $0.5\text{--}500 \mu\text{g L}^{-1}$ ) for permethrin, cypermethrin, and

Table 3 Analytical parameters of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL as an adsorbent for MSPE of the four pyrethroids from tea infusions

Pyrethroids	Regression equation	Linear range ( $\mu\text{g L}^{-1}$ )	$R^2$	LOD ( $\mu\text{g L}^{-1}$ )	RSD <sup>a</sup> (%) ( $n = 6$ )	
					Intraday	Interday
Bifenthrin	$Y = 9.3 \times 10^7 X + 23133$	0.5–50	0.9997	0.0065	5.97	8.89
Permethrin	$Y = 6.0 \times 10^6 X + 11032$	0.5–500	0.9999	0.0369	5.00	8.96
Cypermethrin	$Y = 4.8 \times 10^6 X + 903$	0.5–500	0.9995	0.0935	9.70	11.07
Flucythrinate	$Y = 1.4 \times 10^7 X + 420$	0.5–500	0.9990	0.1017	8.90	11.95

<sup>a</sup> The RSDs were determined by performing six replicate analyses of samples spiked with each pyrethroid at  $10 \mu\text{g L}^{-1}$ .



Table 4 Comparison of different methods for the analysis of pyrethroids in tea samples

Method	Sorbent	Sample amount (mL)	Sorbent amount (mg)	Extraction time (min)	Volume of eluent (mL)	Linear range ( $\mu\text{g L}^{-1}$ )	LOD ( $\mu\text{g L}^{-1}$ )	Ref.
dSPE-HPLC-DAD	C <sub>12</sub> MIM-ATP <sup>a</sup>	8	40	0.5	Acetonitrile, 0.4	2–500	0.6	44
MSPE-DLLME-SFO <sup>b</sup> -GC-ECD	Fe <sub>3</sub> O <sub>4</sub> /MIL-101(Cr)	50	10	10	Methanol, 0.4	0.05–10	0.008–0.015	45
dSPE-UFLC-UV	Fe <sub>3</sub> O <sub>4</sub> /C/PANI microbowls	150	8	12	Methanol, 3	0.1–20	0.025–0.032	46
MSPE-HPLC-UV	Fe <sub>3</sub> O <sub>4</sub> -MCNTs	10	40	15	5% acetic acid acetonitrile, 3	0.05–25 $\mu\text{g g}^{-1}$	0.010–0.018 $\mu\text{g g}^{-1}$	47
MSPE-GC-MS/MS	Fe <sub>3</sub> O <sub>4</sub> /ZIF-8/IL	10	10	2	Acetonitrile, 1.6	0.5–500	0.0065–0.1017	This work

<sup>a</sup> 1-Dodecyl-3-methylimidazolium bromide-attapulgit. <sup>b</sup> Solidification of a floating organic drop.

flucythrinate. However, the response to bifenthrin were far higher than those to the other three pyrethroids and were saturated for the ultrapure water sample spiked at 100  $\mu\text{g L}^{-1}$  and for the tea infusion sample spiked at 200  $\mu\text{g L}^{-1}$ . Consequently, the linear ranges for bifenthrin in ultrapure water sample (0.5–50  $\mu\text{g L}^{-1}$ ) and tea infusion sample (0.5–50  $\mu\text{g L}^{-1}$ ) were smaller than that for the other three pyrethroids and had  $R^2$  values of 0.9991 and 0.9997, respectively. The LODs, calculated using S/N ratios of three, were 0.0040–0.0409  $\mu\text{g L}^{-1}$  in water and 0.0065–0.1017  $\mu\text{g L}^{-1}$  in the tea infusion. The method precision was evaluated by measuring the RSDs for samples spiked with the standard solution at 10  $\mu\text{g L}^{-1}$ . Six replicates were analysed for each experiment. For the water sample, the intra- and inter-day RSDs for the precision were in the range of 0.69–2.13% and 4.26–7.27%, respectively. For the tea infusion, the intra- and inter-day RSDs for the precision were in the range of 5.00–9.70% and 8.89–11.95%, respectively. These results indicate that the suggested method possesses high sensitivity, a wide linear range, and good repeatability.

### 3.5. Comparison of the Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL-based method with other reported methods

We compared our Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL-based method with several other methods for the determination of pyrethroids in tea samples (Table 4). Compared with dispersive SPE, the proposed Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL-based method does not require a time-consuming centrifugation step, which makes it more convenient. Moreover, the extraction time of the developed method is shorter than that of any other MSPE-based method. Regarding analytical performance, our method exhibits the widest linear range and lowest LOD among the available methods. Based on our comparison, we can conclude that our proposed Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL-MSPE method is a sensitive and simple technique for the analysis of four pyrethroid insecticides in tea infusions.

### 3.6. Real sample analysis

We applied the proposed Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL-based MSPE method to the analysis of four pyrethroid insecticides in infusions of Pu-

Table 5 Analytical results for the determination of pyrethroids in infusions of real tea samples

Matrix	Analyte	Spiked concentration ( $\mu\text{g L}^{-1}$ , $n = 3$ )				
		0	10	50	Recovery (%)	RSD (%)
Da Hong Pao	Bifenthrin	<LOD	87.1	6.5	88.8	4.0
	Permethrin	<LOD	88.3	6.4	95.3	3.0
	Cypermethrin	<LOD	78.4	2.5	88.2	2.2
	Flucythrinate	<LOD	80.8	6.5	96.8	2.2
Sichuan Maofeng	Bifenthrin	<LOD	73.5	9.3	81.5	4.2
	Permethrin	<LOD	78.6	5.2	83.7	5.5
	Cypermethrin	<LOD	72.9	4.0	89.4	1.3
	Flucythrinate	<LOD	75.7	1.4	98.1	3.1
Tieguanyin	Bifenthrin	<LOD	76.3	8.7	85.0	7.8
	Permethrin	<LOD	83.6	8.7	94.1	7.2
	Cypermethrin	<LOD	72.1	7.3	87.3	2.5
	Flucythrinate	<LOD	73.1	5.9	85.7	2.3
Pu-erh	Bifenthrin	<LOD	76.2	5.3	84.1	6.3
	Permethrin	<LOD	72.9	10.3	94.0	2.3
	Cypermethrin	<LOD	72.3	10.2	82.4	2.6
	Flucythrinate	<LOD	74.3	5.5	90.9	3.8



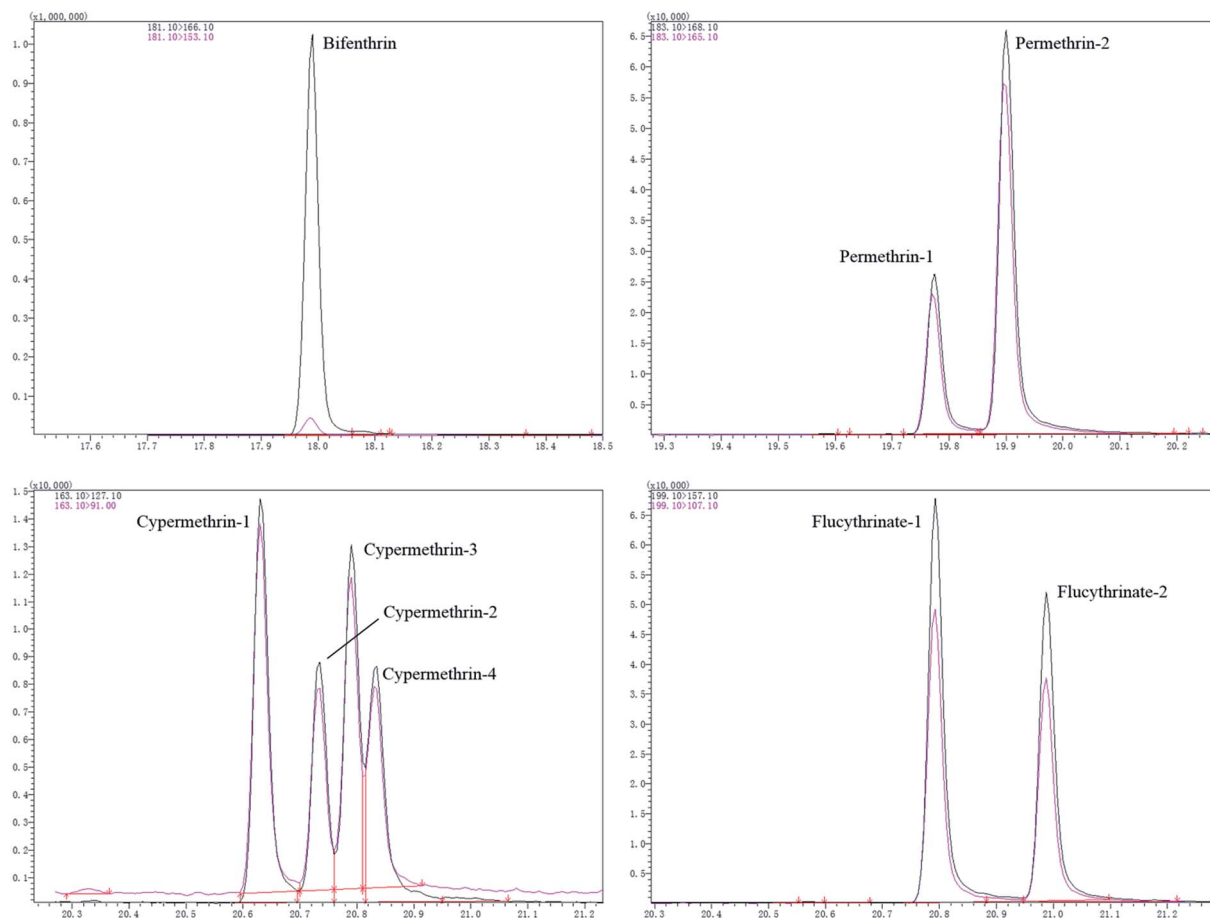


Fig. 4 Extracted total ion chromatograms obtained by the proposed method for pyrethroids from a real tea sample (Sichuan Maofeng) spiked at  $10 \mu\text{g L}^{-1}$ .

erh, Tieguanyin, Da Hong Pao, and Sichuan Maofeng teas. The method showed good accuracy (recovery), precision (RSD), and utility for the determination of the pyrethroid insecticides in the tea infusions (Table 5). No pyrethroids were found in the four tea infusions. The extracted total ion chromatograms of pyrethroids obtained from the Sichuan Maofeng tea after spiking at  $10 \mu\text{g L}^{-1}$  are shown in Fig. 4.

## 4. Conclusions

In this study, a novel magnetic adsorbent was prepared by immobilizing [OMIM]NTf<sub>2</sub> on the surface of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8 via a simple direct immersion method based on the “physisorption” mechanism. The obtained magnetic nanocomposite possessed high magnetism, a satisfactory BET surface area, and a large pore volume. Combined with GC-MS/MS, the adsorbent was first employed as an adsorbent for MSPE of pyrethroid insecticides from tea infusions. Under the optimized conditions, the Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL-based method offers a wide linear range, excellent sensitivity, good accuracy, and good precision. Modification promoted the adsorption of pyrethroids by Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL and gave satisfactory MSPE performance for pyrethroids in tea infusions. The development of Fe<sub>3</sub>O<sub>4</sub>/ZIF-8/IL-

MSPE could be extended to other IL-modified MOFs for the enrichment and determination of many other organic pollutants or pesticide residues in beverages.

## Conflicts of interest

There are no conflicts to declare.

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