


 Cite this: *RSC Adv.*, 2021, **11**, 14746

## Insights into antimicrobial agent sulfacetamide transformation during chlorination disinfection process in aquaculture water†

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Antibiotic addition and chlorination are two common processes in fishery culture. Antibiotic residues not only pollute aquaculture water, but are also one of the potential precursors of disinfection by-products (DBPs) during chlorination. The degradation kinetics, products identification and reaction mechanism of sulfacetamide (SFA), a new sulfonamides antibiotics, and potential formation of haloacetic acids (HAAs) in chlorination were explored. The results showed that the degradation of SFA followed pseudo first-order kinetic model, and chlorinating agent dose, pH of water, water temperature,  $\text{NH}_4^+$ ,  $\text{HCO}_3^-$  and humic acid (HA) had various effects on the degradation of SFA and the yields of HAAs. The presence of  $\text{Br}^-$  accelerated both the degradation rate of SFA and more formation of Br-DBPs. Through the identification of intermediate products, we proposed the transformation pathway of SFA during the chlorination disinfection process. Namely, in this NaClO disinfection system, the C-S bond between the sulfonyl group and benzene ring, and S-N bond between sulfonyl and acylamino of SFA were broken, and then the primary formed groups were further oxidized to produce intermediates, such as chloroanilines and chlorophenols. And then chlorophenols were subsequently chlorinated to form toxic HAAs. The present study might be of significance for the evaluation of effective degradation of SFA and potential production of halogenate-DBPs (H-DBPs) during the chlorination disinfection process in aquaculture water.

Received 1st March 2021

Accepted 13th April 2021

DOI: 10.1039/d1ra01605a

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### 1. Introduction

Aquaculture is a very large and mature industry in the world, and China is the largest producer and exporter of aquatic products as well.<sup>1</sup> Due to the occurrence of various aquatic diseases in the development of aquaculture, antibiotics have become one of the indispensable drugs in aquaculture.<sup>2</sup> China is the largest antibiotic user in the world in which animal consumption accounts for half of the total.<sup>3</sup> Nowadays, the most widely used antibiotics in aquaculture are quinolones, sulfonamides, macrolides,  $\beta$ -lactams, tetracyclines, furans, etc.<sup>4</sup> Antibiotics are added to the feed of aquatic animals in order to assure the safety in aquaculture, but they cannot be completely digested, therefore, the residual antibiotics would enter the aquaculture water body. After human consumption of aquatic

products contaminated by antibiotics, allergic symptoms and chronic intoxication might be caused.<sup>5</sup> For instance, sulfonamides and aureomycin may lead to fulminant liver necrosis, agranulocytosis, aplastic anemia and other diseases.<sup>6,7</sup> Ciprofloxacin may result in interstitial nephritis, hepatitis, liver necrosis, and furazolidone may cause hemolytic anemia and multiple neuritis.<sup>8</sup> Hence, more and more attention has been paid to the residue, transportation and transformation of antibiotics in aquaculture water, especially in the disinfection process.

Chlorination is the most common process in aquaculture water, with the disinfectants of chlorine compounds.<sup>9</sup> Residual antibiotics are degraded during the chlorination process; however, studies have found that antibiotics undergo side reactions with chlorine disinfectants to generate halogenated disinfection by-products (H-DBPs) such as trihalomethanes (THMs) and haloacetic acids (HAAs).<sup>10,11</sup> These H-DBPs have attracted extensive attention in the field of scientific research owing to the latent health risks.<sup>12</sup> In seawater, hypobromous acid (HOBr) can be generated by the chlorine oxidation to be dominant oxidant species, leading to preferential formation of various brominated by-products.<sup>13,14</sup> Rong *et al.*<sup>15</sup> reported that norfloxacin (NOR) in seawater was substituted with HOBr to form two brominated DBPs (Br-DBPs) during chlorination, and

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/d1ra01605a



the toxicity of Br-DBPs was far higher than that of chlorinated DBPs (Cl-DBPs). Marsà *et al.*<sup>16</sup> reported that bromoacetic acid was 89.8 times higher than that of chloroacetic acid in human urothelial cells. THMs exposure resulted in adverse effect for upgrowth and significant tail shortening in zebrafish, and tribromoacetic acid (TBAA) and dichloroacetic acid (DCAA) can significantly increase the deformity rate of zebrafish embryos.<sup>17,18</sup> In brief, possible derivative H-DBPs from antibiotics during chlorination disinfection process in aquaculture water might threaten health to aquatic animals, and subsequently intimidate the fishery economy. Therefore, the transformation fate of H-DBPs in aquaculture water during disinfection is significant for effective assessment of fishery health risk.

Sulfonamides (SAs) are broad-spectrum antibiotics used in aquaculture waters, and the most related researches focus on the degradation kinetics during chlorination disinfection process.<sup>19</sup> In contrast, the transformation mechanism and the potential generation of H-DBPs in the chlorination process still need to be further studied to explore the threat to fisheries aquaculture. Consequently, in the present study, we selected a new SA antibiotic (sulfacetamide, SFA) as the model substrate (the detailed information was shown in Table S1 in the ESI†). We studied the relationship between reaction kinetics of SFA in chlorination and the parameters, such as available chlorine concentration, pH, temperature, and other common components (*i.e.*  $\text{NH}_4^+$ ,  $\text{HCO}_3^-$  and HA) in aquaculture water. Based on the above analysis, the potential formation of H-DBPs represented by HAAs and transformation mechanism of SFA in the chlorination process were explored. The present research might provide technical support for the effective removal of SFA and the control of the production of HAAs during the chlorination process in aquaculture water.

## 2. Materials and methods

### 2.1 Chemicals and reagents

All chemicals were analytical grade except as noted. Mixed standard of 9 kinds of halogenated acetic acids (*i.e.* monochloroacetic acid (MCAA), dichloroacetic acid (DCAA), trichloroacetic acid (TCAA), monobromoacetic acid (MBAA), dibromoacetic acid (DBAA), tribromoacetic acid (TBAA), bromochloroacetic acid (BCAA), bromodichloroacetic acid (BDCAA), and chlorodibromoacetic acid (CDBAA)) were obtained from Accustandard (America). Humic acid (HA, BR) were purchased from Yuanye Biological Technology (Shanghai, China). Sodium hypochlorite solution (available chlorine,  $\text{HOCl}/\text{ClO}^-/\text{Cl}_2\text{O}$ , ~10% w/w) and ascorbic acid were purchased from Aladdin Co., Ltd. (Shanghai, China). Need of special note is the pH was maintained at 5 with 10 mM phosphate buffer except for the effect of pH in the examined experiments. Therefore,  $[\text{Cl}_2]$  was used to represent available chlorine species in the full text. Methyl *tert*-butyl ether (MTBE) and acetonitrile (HPLC grade) were obtained from Shanghai Anpu Co., Ltd., China. Sulfacetamide (SFA, 99.5%), concentrated sulfuric acid ( $\text{H}_2\text{SO}_4$ , 98%), sodium bicarbonate ( $\text{NaHCO}_3$ , AR), formic acid ( $\text{HCOOH}$ , AR), potassium dihydrogen phosphate ( $\text{KH}_2\text{PO}_4$ , AR),

dipotassium hydrogen phosphate ( $\text{K}_2\text{HPO}_4$ , AR), sodium hydroxide ( $\text{NaOH}$ , AR), sodium chloride (NaCl, AR), and sodium bromide (NaBr, AR) were all from Sinopharm Chemical Reagent Co., Ltd., China. All solutions were prepared using ultra-pure water (18.2 MΩ cm) produced by Hetai Co., Ltd. of Shanghai, China.

### 2.2 Experimental procedure

For SFA oxidation by  $[\text{Cl}_2]$ , the experiments were conducted using a 150 mL conical flask with 100 mL SFA solution of 1 mg L<sup>-1</sup>. The experiments were started by adding certain volumes of the  $[\text{Cl}_2]$  stock solution. At fixed points in time, 1 mL of samples was rapidly quenched with excessive ascorbic acid to stop the reaction, and then transferred into a high performance liquid chromatography (HPLC) instrument to perform analyses of the data. The pH of all experiments was maintained with 10 mM phosphate buffer. The temperature of all experiments was maintained by DF-101S constant temperature magnetic stirrer. The effects of  $[\text{Cl}_2]$  (50–125 μM), reaction pH (5–9), reaction temperature (10–40 °C),  $\text{NH}_4^+$  (0–5 mg L<sup>-1</sup>),  $\text{HCO}_3^-$  (0–100 mg L<sup>-1</sup>), HA (0–15 mg L<sup>-1</sup>) and  $\text{Br}^-$  (0–65 mg L<sup>-1</sup>) on SFA degradation were examined. In order to investigate the oxidation of SFA and the formation of HAAs in different types of aquaculture water during chlorination disinfection process, we also prepared water samples with specific  $\text{Cl}^-$  and  $\text{Br}^-$  concentrations at pH = 5 as models of marine culture water (containing 6.6 g L<sup>-1</sup> of  $\text{Cl}^-$  and 22 mg L<sup>-1</sup> of  $\text{Br}^-$ ) and seawater (containing 19 g L<sup>-1</sup> of  $\text{Cl}^-$  and 65 mg L<sup>-1</sup> of  $\text{Br}^-$ ). All the experiments were conducted at least duplicate, and the error bars were provided in the figures.

To examine the potential formation of HAAs, 40 mL of chlorinated water sample for 24 h was transferred to a 100 mL centrifuge tube, and then 2 mL of concentrated sulfuric acid was added to make the pH of the solution lower than 0.5, and subsequently 9 g of baked sodium chloride was quickly added to dissolve with shaking. The water samples were then extracted twice with 4 mL MTBE, and each extraction was shaken for 5 min and left to stand for 5 min, ultimately the extracts were combined. Afterwards, 3 mL of the extract was transferred into a 50 mL centrifuge tube, and after adding 3 mL of a newly prepared 10% of sulfuric acid methanol solution, mixed well and placed it into 50 °C water bath for 2 h for derivatization. After the derivative water sample was cooled, 7 mL of 250 g L<sup>-1</sup> sodium chloride solution was added, quickly followed shaken up. After leaving the water sample quietly for 5 min, a pipette was used to remove the water phase, and 1 mL of saturated sodium bicarbonate solution was slowly added to the organic layer to exhaust the gas. Finally, 1 mL of the upper organic phase was placed in a 1.5 mL brown sample bottle for GC-ECD analysis.

### 2.3 Analytical methods

A Shimadzu 20A HPLC equipped with a C18 column (4.6 mm × 150 mm, 5 μm) and a UV detector was used to determine the concentration changes of SFA in the experiments. The mobile phase used was 0.1% formic acid and acetonitrile ( $V_a/V_b = 70/$



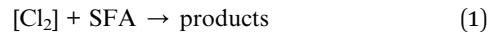
30) at a flow rate of  $1.0 \text{ mL min}^{-1}$ . The detection wavelength was set at  $275 \text{ nm}$ , the column temperature was  $30 \text{ }^\circ\text{C}$ , and the injection volume was  $20 \mu\text{L}$  (see HPLC chromatogram of SFA in Fig. S1†). The concentration change of HA was monitored by UV-2600 spectrophotometer of Shimadzu company (Japan), and the analytical wavelength was set at  $254 \text{ nm}$ .

The analytical method to determine HAAs refers to 'water quality-determination of haloacetic acids-gas chromatography' (HJ 758-2015), a national environmental protection standard of the People's Republic of China. An Agilent 7890A gas chromatography spectrometer (GC) equipped with a DB-5 column ( $30 \text{ m} \times 0.32 \text{ mm} \times 0.25 \mu\text{m}$ ) and an electron capture detector (ECD) was used to determine the concentrations of HAAs in NaClO disinfection process. The injections were made in the splitless mode using an injection temperature of  $210 \text{ }^\circ\text{C}$  and the injection volume was  $1 \mu\text{L}$ . Nitrogen was used as carrier gas and make-up gas, the flow rate of carrier gas was  $2.5 \text{ mL min}^{-1}$ . The temperature of ECD detector was  $260 \text{ }^\circ\text{C}$ . The GC column was operated in a temperature programmed mode with an initial temperature of  $40 \text{ }^\circ\text{C}$  held for 5 min, ramp first to  $65 \text{ }^\circ\text{C}$  with a  $5 \text{ }^\circ\text{C min}^{-1}$  rate, then to  $75 \text{ }^\circ\text{C}$  with  $1 \text{ }^\circ\text{C min}^{-1}$  rate, then to  $135 \text{ }^\circ\text{C}$  with  $5 \text{ }^\circ\text{C min}^{-1}$  rate, and then to  $280 \text{ }^\circ\text{C}$  with  $20 \text{ }^\circ\text{C min}^{-1}$  rate and held at that temperature for 5 min. The external standard method was used for quantitative analyses of halogenated methyl acetate derivatives of the nine HAAs mixed standards after methyl esterification pretreatment. The organic intermediates produced during the SFA degradation were identified by gas chromatography-mass spectrometry (GC-MS, Shimadzu GCMS-QP2010 plus) and the details are shown in Text S1.†

### 3. Results and discussion

#### 3.1 Chlorination kinetics of SFA

The chlorination of SFA is much complex in the actual disinfection of aquaculture water, but the main reaction process can be roughly described as eqn (1):



The rate equation of reaction (1) should be:

$$-\frac{d[\text{SFA}]}{dt} = k \cdot [\text{SFA}]^m \cdot [\text{Cl}_2]^n \quad (2)$$

where  $k$  is the reaction rate constant and  $m, n$  represent the reaction order with respect to the corresponding substrate.<sup>20</sup>

The degradation of SFA at a concentration of  $5 \mu\text{M}$  was examined at initial  $[\text{Cl}_2]$  concentrations of  $50, 75, 100$ , and  $125 \mu\text{M}$ . Fig. S2† and 1 show that the degradation effect of SFA increased with the increase of the concentration of  $[\text{Cl}_2]$ , and the degradation of SFA satisfied the pseudo first-order kinetics with  $R^2 > 0.998$  (Fig. 1a). In the actual disinfection of aquaculture water, the concentration of disinfectant is usually much more than that of antibiotic.<sup>21</sup> In the present study, the initial mole ratio of SFA to  $[\text{Cl}_2]$  was set at lower than  $1 : 10$ . Therefore, the concentration of  $[\text{Cl}_2]$  was far excessive and can be approximated as constant compared to SFA in the whole chlorination process. Therefore, the reaction kinetics between disinfectants  $[\text{Cl}_2]$  and antibiotics SFA are recognized as pseudo first-order model. As such, eqn (2) can be simplified to eqn (3).<sup>22</sup>

$$-\frac{d[\text{SFA}]}{dt} = k_{\text{obs}} \cdot [\text{SFA}] \quad (3)$$

where  $k_{\text{obs}}$  is the pseudo first-order reaction rate constant,  $k_{\text{obs}} = k \cdot [\text{Cl}_2]$ .

We can also observe that at pH 5 and temperature of  $20 \text{ }^\circ\text{C}$ , as the concentration of chlorine disinfectant increased, the reaction rate constant  $k_{\text{obs}}$  were  $0.01044, 0.02298, 0.03783$ , and  $0.05596 \text{ s}^{-1}$ , respectively, and there was a linear relationship between  $k_{\text{obs}}$  and  $[\text{Cl}_2]$  dose ( $R^2 = 0.993$ ) (Fig. 1b). Hence, it can be noted that the concentration of  $[\text{Cl}_2]$  was an important factor for accelerating reaction kinetics.

The degradation of SFA in NaClO disinfection system at pH 5–9 was presented in Fig. S3† and 2, and the results of the experiment showed that the reaction rate of SFA was the fastest under neutral condition, and slowed down under weak acid or

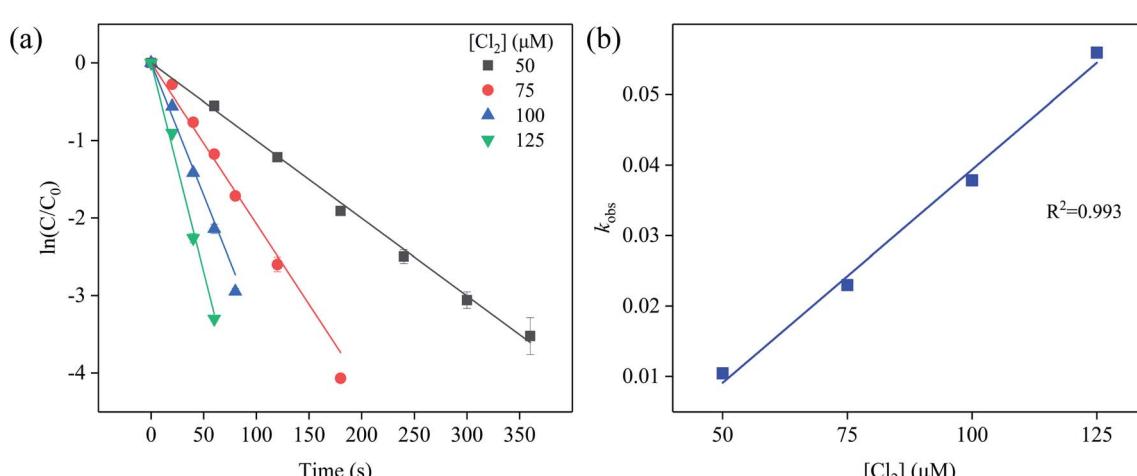


Fig. 1 (a) Effect of  $[\text{Cl}_2]$  dose on the degradation of SFA, and (b) the linear relationship between  $k_{\text{obs}}$  and  $[\text{Cl}_2]$  dose. Experimental conditions:  $[\text{SFA}]_0 = 5 \mu\text{M}$ ,  $\text{pH}_{\text{ini}} = 5$ ,  $T = 20 \text{ }^\circ\text{C}$ .



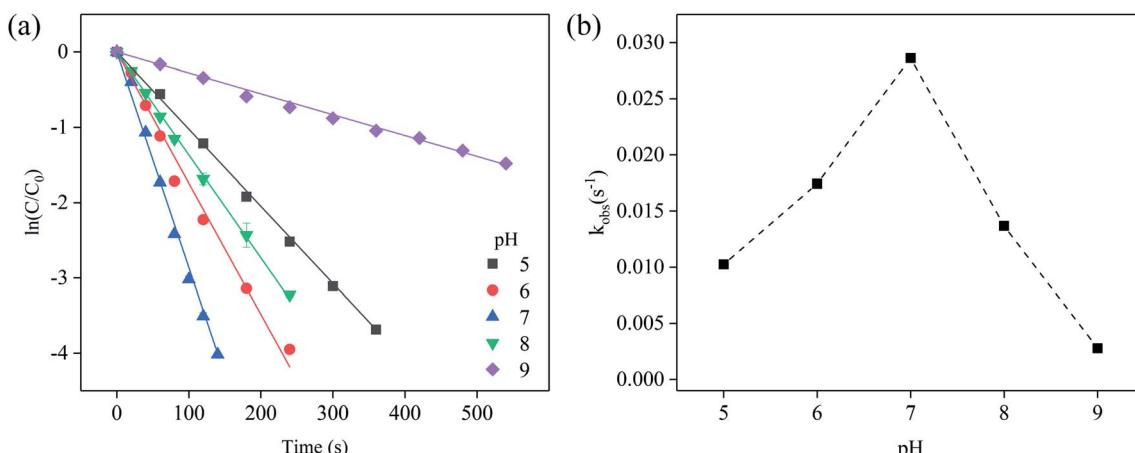


Fig. 2 (a) Oxidation of SFA by  $[Cl_2]$ , and (b) the variation of  $k_{obs}$  at different pH values. Experimental conditions:  $[SFA]_0 = 5 \mu M$ ,  $[Cl_2]_0 = 50 \mu M$ ,  $T = 20^\circ C$ .

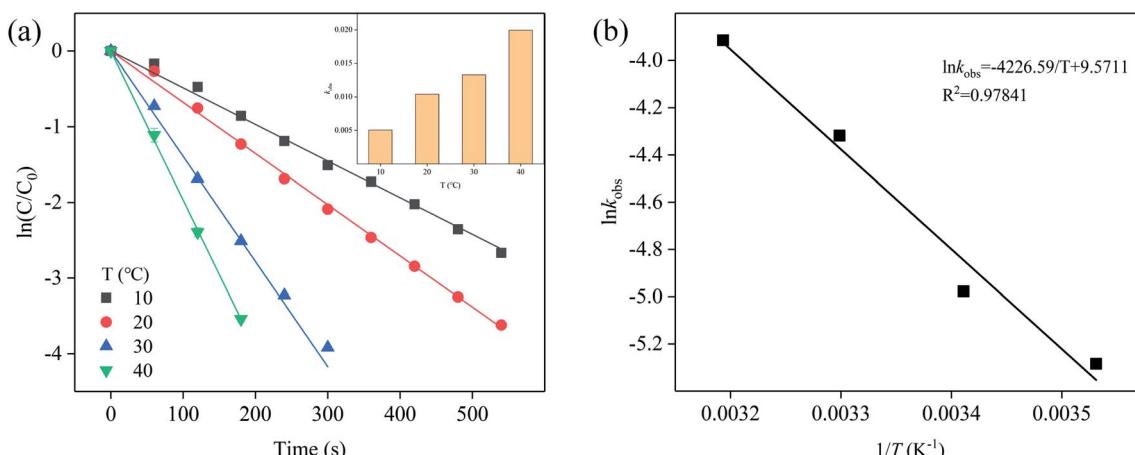


Fig. 3 (a) Effect of temperature, and (b) Arrhenius plot. Experimental conditions:  $[SFA]_0 = 5 \mu M$ ,  $[Cl_2]_0 = 50 \mu M$ ,  $pH_{ini} = 5$ .

weak base condition. The observed pseudo first-order kinetic constants of SFA degradation first increased from  $0.01044$  to  $0.03003 \text{ s}^{-1}$  with the pH extended from 5 to 7, and then decreased to  $0.00273 \text{ s}^{-1}$  when pH further reached to 9. The macroscopic  $K_{a1}$  and  $K_{a2}$  constants of SFA (Fig. S4†) are related to amino and amide groups, respectively. The  $pK_{a1}$  and  $pK_{a2}$  of SFA were 1.76 and 5.22 (Table S1†),<sup>23</sup> and the  $pK_a$  of HOCl was 7.54,<sup>24</sup> indicating both SFA and some available chlorine species existed in molecular state (*i.e.* SFA and HOCl) at pH 5, respectively. Although HOCl had strong oxidizability under acidic conditions, the molecular state of SFA had relatively weaker ability to provide electrons for molecular HOCl. When the pH value was in the range from 5.22 to 7.49, SFA was in the form of deprotonated state, and HOCl existed in the molecular state, thus, the deprotonated amide group of SFA can provide electrons more easily to HOCl with higher oxidizability. Furthermore, trace  $Cl_2O$ , potent chlorinating agent at  $pH \lesssim 8$  dependent on the existence of HOCl (eqn (4)), is more potent electrophiles than HOCl/ $ClO^-$ .<sup>25</sup> As such, the electrophilic action to SFA by slight  $Cl_2O$  can explain the effective removal of SFA and the increased reaction rate under neutral conditions.

In alkaline conditions ( $pH > 7.49$ ), SFA existed in deprotonic state and the available chlorine species mainly existed in the form of  $ClO^-$ , however, the oxidative capability and electrophilicity of  $ClO^-$  was much weaker than that of HOCl and  $Cl_2O$ ,<sup>24,25</sup> which led to the slower degradation of SFA.



The temperature had a great influence on the degradation rate of SFA, and the degradation of SFA at high temperature was faster than that at low temperature (Fig. S5†). Fig. 3a shows the pseudo first order kinetic constants of SFA increased from  $0.00507$  to  $0.01994 \text{ s}^{-1}$  with the temperature elevated from 10 to 40 °C. The effect of temperature on chemical reaction rate can also be expressed by Arrhenius equation (eqn (5)).<sup>26</sup>

$$\ln k_{obs} = \ln A - \frac{E_a}{RT} \quad (5)$$

where  $A$  is the Arrhenius constant,  $\text{s}^{-1}$ ;  $R$  is the molar gas constant,  $8.314 \times 10^{-3} \text{ kJ (mol K)}^{-1}$ ;  $T$  is the absolute temperature, K;  $E_a$  is the reactive activation energy,  $\text{kJ mol}^{-1}$ .

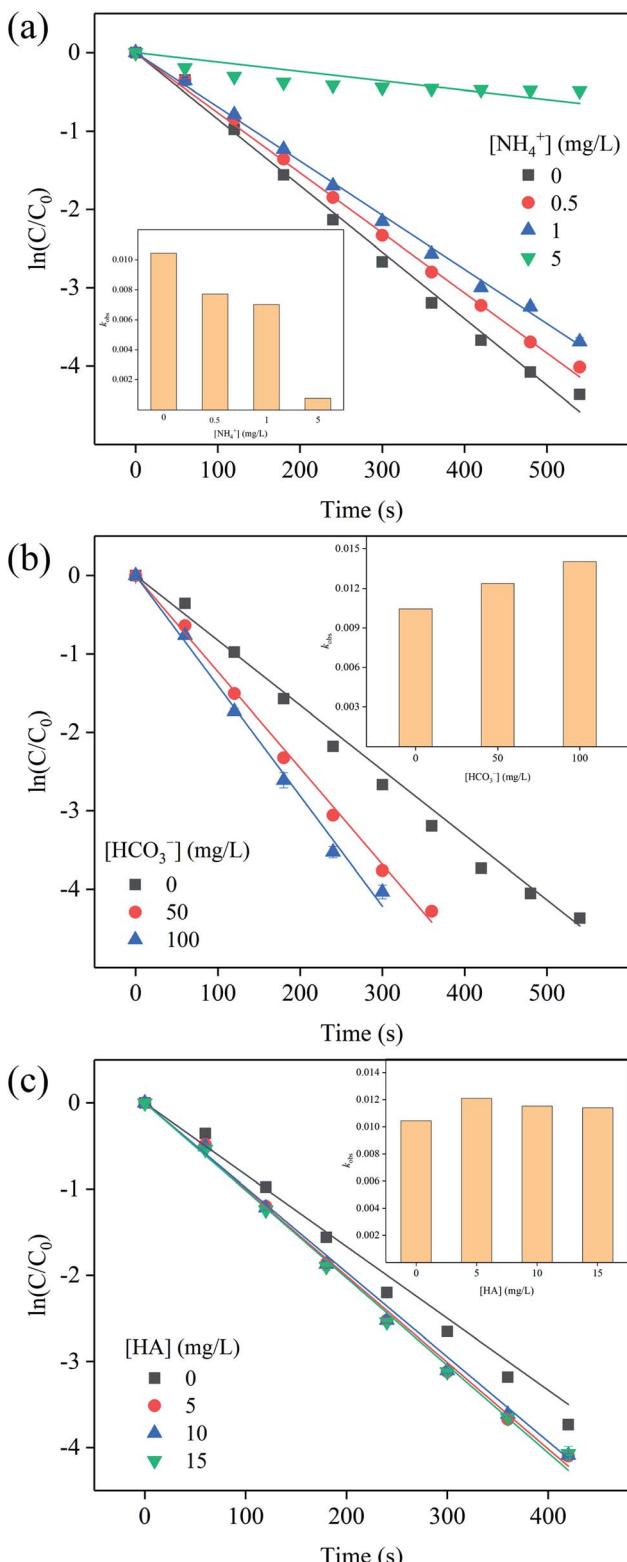


Fig. 4 Effects of concentrations of (a)  $NH_4^+$ , (b)  $HCO_3^-$ , and (c) HA on the oxidation of SFA by  $[Cl_2]$ . Experimental conditions:  $[SFA]_0 = 5 \mu M$ ,  $[Cl_2]_0 = 50 \mu M$ ,  $pH_{ini} = 5$ ,  $T = 20^\circ C$ .

According to Fig. 3b, the calculated observed activation energy was  $35.14 \text{ kJ mol}^{-1}$ . Zhang *et al.*<sup>15</sup> reported that the observed activation energies of the reactions of NOR and

roxithromycin (ROX) with  $[Cl_2]$  were  $105.00$  and  $79.34 \text{ kJ mol}^{-1}$ , respectively. In contrast, the obtained observed activation energy of  $[Cl_2]$  to oxidize SFA is much lower, indicating the reaction rate is faster under the same experimental conditions, and the reaction between SFA and  $[Cl_2]$  is easier to occur from the perspective of activation energy as well.

$NH_4^+$  and  $HCO_3^-$  are common cation and anion in aquaculture water.<sup>27</sup> The presence of  $NH_4^+$  can inhibit the oxidation of SFA by  $[Cl_2]$  (Fig. S6a† and 4a). With the concentration of  $NH_4^+$  increased from  $0$  to  $5 \text{ mg L}^{-1}$ , the  $k_{obs}$  decreased from  $0.01044$  to  $0.00076 \text{ s}^{-1}$ . When the concentration of  $NH_4^+$  reached  $5 \text{ mg L}^{-1}$ , only less than  $40\%$  of SFA was degraded within  $10 \text{ min}$  (Fig. S6a†). These phenomena might be attributed that HOCl could react with  $NH_4^+$  to form  $NH_2Cl$ ,  $NHCl_2$  and  $NCl_3$  (eqn (6)–(9)),<sup>28,29</sup> whose oxidation abilities were weaker than that of HOCl,<sup>30</sup> leading to the decreased oxidation of SFA (Fig. S6a†).

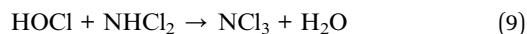
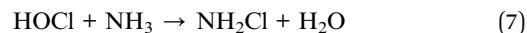
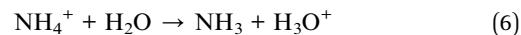


Fig. S6b† and 4b showed the  $HCO_3^-$  had little effect on SFA degradation. With the concentration of  $HCO_3^-$  increased from  $0$  to  $100 \text{ mg L}^{-1}$ ,  $k_{obs}$  varied much more slightly and negligibly, *i.e.* only from  $0.01044$  to  $0.01403 \text{ s}^{-1}$ . This phenomenon indicated that  $HCO_3^-$  had neglected influence on the transformation of SFA.

Humic acid (HA) is the main component of natural organic matter, which accounts for  $50\text{--}90\%$ .<sup>31,32</sup> HA is commonly found in natural water bodies, and the concentration in water bodies ranges from  $0$  to  $30 \text{ mg L}^{-1}$ .<sup>33</sup> Therefore, the effects of HA on the degradation of SFA by  $[Cl_2]$  were also evaluated. Fig. S6c† and 4c showed the negligible effect of altered concentrations of HA on the oxidation of SFA by  $[Cl_2]$ . When the concentration of HA increased from  $0$  to  $15 \text{ mg L}^{-1}$ , the changes of  $k_{obs}$  were ignorable (Fig. 4c). At the same time, we measured the concentration changes of HA during the reaction. The results showed that HA hardly degraded in  $420 \text{ s}$ .  $[Cl_2]$  with relatively low redox potential (*e.g.*  $E_{HOCl} = 740 \text{ mV}$ ), has limited oxidation capacity to degrade HA in the examined  $420 \text{ s}$ .<sup>34</sup>

### 3.2 Influence of different water bodies

According to the characteristics of water quality, there are not only freshwater aquaculture models, but also mariculture models. The intricacy of chemical characteristics and features of chlorination in seawater greatly differ from those of freshwater, as a result of the oxidation of bromide ( $\sim 65 \text{ mg L}^{-1}$ ) present in seawater, thereby resulting in HOBr as the dominant oxidant species, unlike HOCl in freshwater.<sup>35</sup> Fig. S7a† and 5a showed the effects of different water bodies on the degradation of SFA by  $[Cl_2]$ . Most of SFA was degraded in  $100 \text{ s}$  in marine



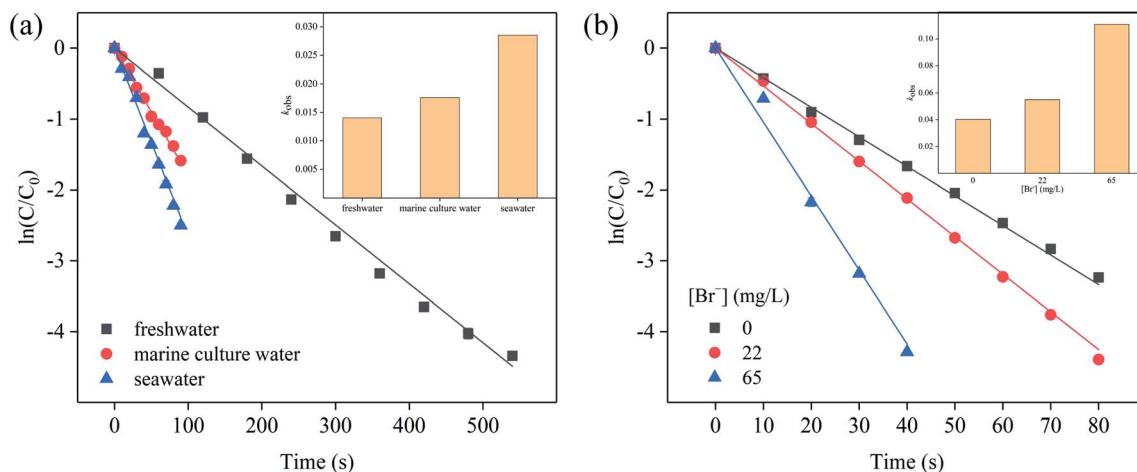


Fig. 5 Effects of (a) different water bodies and (b)  $\text{Br}^-$  concentration on SFA degradation. Experimental conditions:  $[\text{SFA}]_0 = 5 \mu\text{M}$ ,  $[\text{Cl}_2]_0 = 50 \mu\text{M}$ ,  $\text{pH}_{\text{ini}} = 5$ ,  $T = 20^\circ\text{C}$ .

culture water and seawater. However, in freshwater, it took about 550 s to be degraded completely. The degradation rate of SFA in various media was decreased in the order of seawater >

marine culture water > freshwater, *i.e.* the  $k_{\text{obs}}$  of them were in the sequence of  $0.02854 \text{ s}^{-1} > 0.01762 \text{ s}^{-1} > 0.01044 \text{ s}^{-1}$  through the fitting of the first-order kinetic model (eqn (3)). Considering

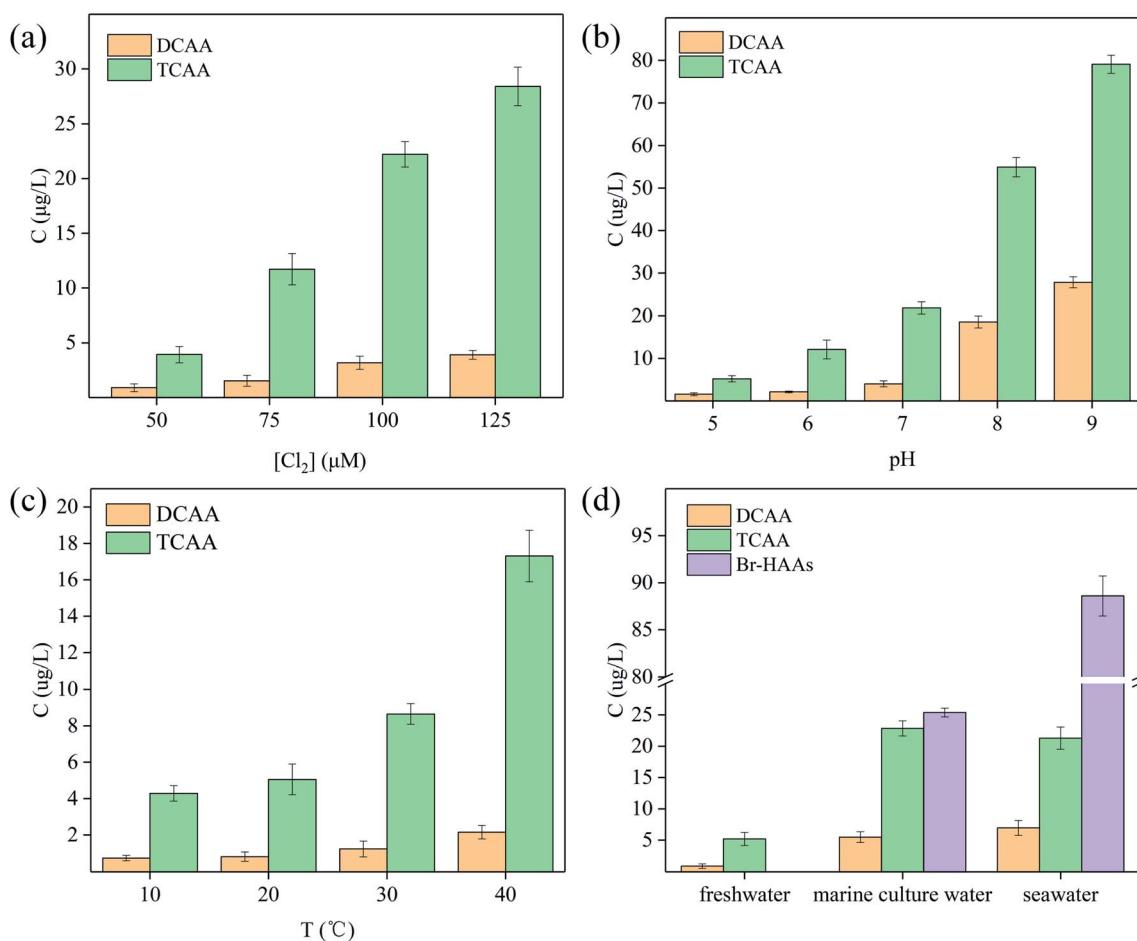
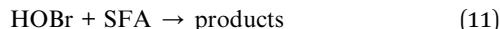
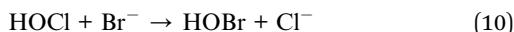


Fig. 6 Effects of (a) the concentration of  $[\text{Cl}_2]$ , (b) pH, (c) temperature and (d) various water bodies on the yields of HAAs. Experimental conditions:  $[\text{SFA}]_0 = 5 \mu\text{M}$ ,  $[\text{Cl}_2]_0 = 50 \mu\text{M}$  (except for a),  $\text{pH}_{\text{ini}} = 5$  (except for b),  $T = 20^\circ\text{C}$  (except for c). Note: marine culture water contained  $6.6 \text{ g L}^{-1}$  of  $\text{Cl}^-$  and  $22 \text{ mg L}^{-1}$  of  $\text{Br}^-$  and seawater contained  $19 \text{ g L}^{-1}$  of  $\text{Cl}^-$  and  $65 \text{ mg L}^{-1}$  of  $\text{Br}^-$ .

Table 1 Degradation products information of SFA

Number	Characteristic ions <i>m/z</i> (auxiliary ions <i>m/z</i> )	Structure	Number	Characteristic ions <i>m/z</i> (auxiliary ions <i>m/z</i> )	Structure
p1	49		p6	161(163)	
p2	109(80)		p7	143(80)	
p3	128(65)		p8	162(64)	
p4	109(80)		p9	97(132)	
p5	65(92)		p10	232(131)	

that the reaction rate constant is likely to depend on the change of the concentration of  $\text{Br}^-$ , therefore, the concentration of  $\text{Cl}^-$  was fixed at  $6.6 \text{ g L}^{-1}$ , and the  $\text{Br}^-$  concentration effect was examined. The results in Fig. S7b† and 5b illustrated that the degradation of SFA increased with the  $\text{Br}^-$  concentration amplified from 0 to  $65 \text{ mg L}^{-1}$ , and the reaction rate constant increased from  $0.04027$  to  $0.11078 \text{ s}^{-1}$ . This phenomenon indicated that the presence of  $\text{Br}^-$  accelerated the reaction rate. The oxidation rate of  $\text{Br}^-$  by  $\text{HOCl}$  to form  $\text{HOBr}$  was much higher than that of organic compounds with  $\text{HOCl}$ , thus generating a large amount of  $\text{HOBr}$  with the characteristic of higher electrophilicity, and subsequently, the amino and other electron donating groups in SFA react more easily and rapidly with  $\text{HOBr}$ , leading to promoted degradation of SFA (eqn (10) and (11)).<sup>36-38</sup>



### 3.3 Potential formation of HAAs

The formation potential of HAAs during the chlorination of SFA in the simulated aquaculture water was examined. It was found that two main kinds of HAAs, *i.e.* DCAA and TCAA, were produced in the chlorination of SFA in the fresh aquaculture water (Fig. 6), which bore a resemblance to the chlorination in drinking water.<sup>39</sup> With the increase of chlorine disinfectant, the generation of DCAA and TCAA gradually increased, and the total concentration of HAAs increased from  $5.11$  to  $31.33 \mu\text{g L}^{-1}$  (Fig. 6a), indicating that the dosage of chlorine disinfectants

was a key parameter affecting Cl-DBPs production. When the dosage of  $[\text{Cl}_2]$  is insufficient, the organic pollutants and some bacteria cannot be completely eliminated, further influencing the ideal disinfection effect. When the chlorine disinfectants dosage is immoderate, the yields of Cl-DBPs will increase, which in turn poses threatening risk for fishery. Therefore, in the actual disinfection process, the chlorine ratio should be strictly controlled to ensure the disinfection effect and the safety of aquaculture water.<sup>21</sup>

As can be seen from Fig. 6b, the formation of DCAA and TCAA increased with the pH rise, the reason for which might be alkaline pH promoted the reaction equilibrium movement towards the direction for formation of chloroacetic salts, further improving the generation of HAAs. In addition, the temperature also has a positive influence on the yield of HAAs (Fig. 6c). During the chlorination process, the concentrations of DCAA and TCAA gradually increased, and the total concentration of HAAs enlarged from  $4.63$  to  $18.92 \mu\text{g L}^{-1}$ , as the temperature increased from  $10$  to  $40^\circ\text{C}$ . The degradation kinetics of SFA indicated the obtained lower observed activation energy encouraged the easily occurred reaction between SFA and  $[\text{Cl}_2]$ , possibly supporting the susceptive formation of HAAs.

Fig. 6d showed the effect of different types of aquaculture waters on the generation potential of HAAs. The production of DCAA and TCAA in marine culture water and seawater were much more than that in freshwater. Furthermore, the Br-HAAs, such as BCAA, BDCAA, DBAA, CDBAA and TBAA, can be detected in both marine culture water and seawater (Fig. 6d and S8†), which might attribute to high concentration  $\text{Br}^-$  participating in the generation of higher oxidative  $\text{HOBr}$  (eqn (9)), further accelerating the formation of Br-HAAs (Fig. S8†).<sup>40</sup>



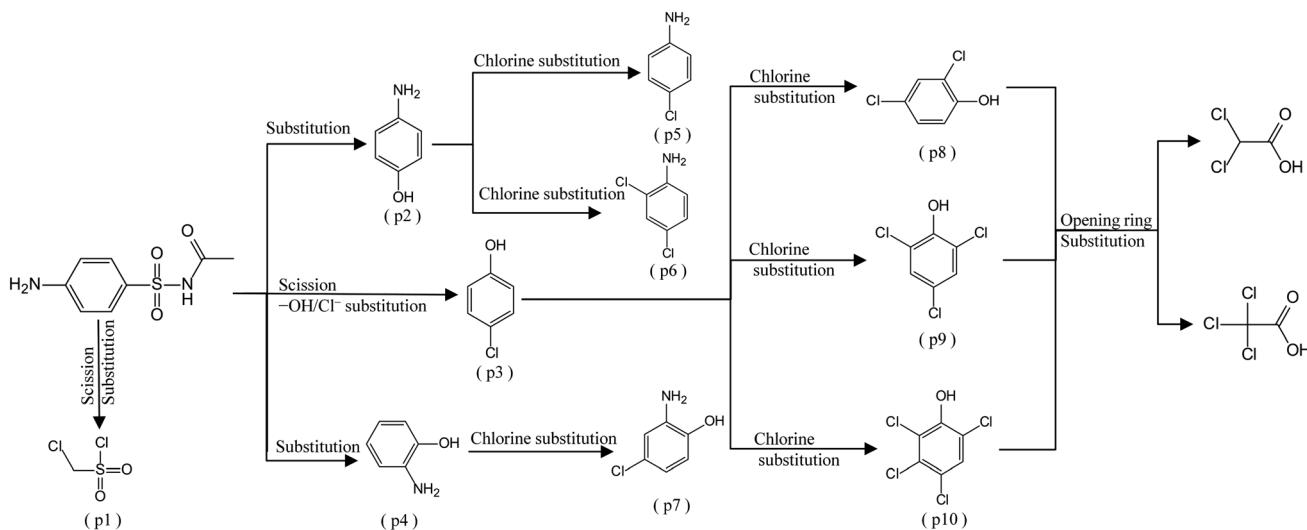


Fig. 7 Possible reaction pathway of SFA during chlorination process in fresh aquaculture water.

### 3.4 Proposed reaction pathway of SFA during the chlorination process

In addition to the maternal SFA and the final two DBPs (DCAA, TCAA), we also detected the intermediate products of SFA in NaClO disinfection system by GC-MS. Based on the mass spectrum data, several main transformation products were detected, and listed in Table 1, Fig. S9–S20.† Based on the reasonable analysis of the experimental results,<sup>20,41–44</sup> the degradation route of SFA and potential formation pathway of HAAs were proposed in Fig. 7.

The transformation process of SFA in NaClO disinfection system can be divided into three steps: (1) with the oxidation of  $[Cl_2]$ , the C-S bond between sulfonyl group and benzene ring and S-N bond between sulfonyl and acylamino groups of SFA were scissored, and then the intermediate sulfonyl group formed p1(chloromethanesulfonyl chloride), and the rest intermediate aminobenzene were oxidized to produce p2 (4-hydroxyaniline), p3 (4-chlorophenol) and p4 (2-hydroxyaniline).<sup>20,43</sup> (2) Due to the chlorination mechanism in NaClO system, the intermediate products p2 and p4 were substituted by chlorine to produce p5 (4-chloroaniline), p6 (2,4-dichloroaniline) and p7 (2-amino-4-chlorophenol). The intermediate product p3 was substituted by chlorine to form poly-chlorophenols (p8–p10).<sup>44</sup> (3) The chlorophenols could be cleaved under the action of  $[Cl_2]$ , and then fully combined with Cl in the subsequent chlorination process to produce HAAs and other products.<sup>12,41,42</sup>

## 4. Conclusions

This study demonstrated that the reaction of SFA with  $[Cl_2]$  followed pseudo first-order kinetic model. The results showed that chlorine disinfectant dosage and temperature had positive effects on the degradation of SFA and Cl-HAAs formation; and neutral pH favored the degradation of SFA, but alkaline pH was more beneficial to generate Cl-HAAs. The common  $NH_4^+$  cation

addition inhibited removal of SFA; in comparison,  $HCO_3^-$  and HA had a negligible effect on the degradation of SFA. The presence of  $Br^-$  not only accelerated the degradation rate of SFA, but also led to the production of Br-HAAs. In the case of excess  $[Cl_2]$  ( $[Cl_2]/[SFA] > 10$ ), C-S and S-N bonds around sulfonyl group of SFA were scissored, and then the primary formed groups were further oxidized to produce intermediate products, such as chloroanilines and chlorophenols. Then the chlorophenols were further chlorinated to form HAAs with toxicity.

## Author contributions

Yaoguang Guo: investigation, visualization, writing original draft, and funding acquisition. Zhiyuan Liu: methodology, investigation, and writing original draft. Xiaoyi Lou: funding acquisition, writing – review & editing, supervision. Changling Fang: software, and formal analysis. Pu Wang: methodology, and investigation. Genying Wu: methodology. Jie Guan: resources, funding acquisition, and supervision.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The present work received financial support from Shanghai Sailing Program (18YF1429900 and 19YF1459900), Shanghai Natural Science Foundation (20ZR1421100), Natural Science Foundation of China (52070127), the Central-Public interest Scientific Institution Basal Research Fund (2019T13, 2019T14), and postgraduate Program Fund of Shanghai Polytechnic University (EGD19YJ0079). Dr Y. Guo also appreciates the Cultivation Discipline Fund of Shanghai Polytechnic University (XXKPY1601), the Project of Key Undergraduate Courses



(Instrumental Analysis) from Shanghai Municipal Education Commission.

## References

- X. B. Bai, Z. T. Fu, N. Li, S. Stankovski, X. S. Zhang and X. X. Li, *J. Clean. Prod.*, 2021, **288**, 125633.
- M. Ramesh, T. Thilagavathi, R. Rathika and R. K. Poopal, Antioxidant status, *Aquaculture*, 2018, **491**, 10–19.
- X. Liu, J. C. Steele and X. Z. Meng, *Environ. Pollut.*, 2017, **223**, 161–169.
- L. Santos and F. Ramos, *Trends Food Sci. Technol.*, 2016, **52**, 16–30.
- R. C. Okocha, I. O. Olatoye and O. B. Adedeji, *Publ. Health Rev.*, 2018, **39**, 1–22.
- N. Hamid, M. Junaid and D. S. Pei, *J. Hazard. Mater.*, 2020, **382**, 121106.
- M. Hajji, H. Jebali, A. Mrad, Y. Blel, N. Brahmi, R. Kheder, S. Beji, L. B. Fatma, W. Smaoui, M. Krid, F. B. Hmida, L. Rais and M. K. Zouaghi, *Drug Safety – Case Reports*, 2018, vol. 5.
- B. H. Ali, Pharmacological, *Vet. Res. Commun.*, 1999, **23**, 343–360.
- Z. H. Pan, Y. J. Zhu, M. Wei, Y. Y. Zhang and K. K. Yu, *J. Environ. Sci.*, 2021, **102**, 170–184.
- H. Sanawar, Y. H. Xiong, A. Alam, J. P. Croué and P. Y. Honga, *Aquaculture*, 2017, **480**, 94–102.
- H. C. Hong, F. Q. Huang, F. Y. Wang, L. X. Ding, H. J. Lin and Y. Liang, *J. Hydrol.*, 2013, **476**, 274–279.
- H. C. Hong, Y. J. Xiong, M. Y. Ruan, F. L. Liao, H. J. Lin and Y. Liang, *Sci. Total Environ.*, 2013, **444**, 196–204.
- R. K. Padhi, S. Subramanian, A. K. Mohanty, S. N. Bramha, M. V. R. Prasad and K. K. Satpathy, *Environ. Eng. Res.*, 2021, **17**, 57–62.
- R. K. Padhi, S. Subramanian, A. K. Mohanty and K. K. Satpathy, *Environ. Monit. Assess.*, 2019, **191**, 1–13.
- Y. Y. Zhang, C. Rong, Y. Q. Song, Y. H. Wang, J. Y. Pei, X. Y. Tang, R. J. Zhang and K. F. Yu, *Chemosphere*, 2017, **182**, 245–254.
- A. Marsà, C. Cortés, A. Hernández and R. Marcos, *Toxicol. Appl. Pharmacol.*, 2018, **347**, 70–78.
- T. Elisabet, P. Esther, G. L. Javier, J. M. Llobet and G. C. Jesús, *J. Water Health*, 2015, **13**, 54–66.
- Y. Huang, H. Li, Q. Zhou, A. Li, C. Shuang, Q. Xian, B. Xu and Y. Pan, *Water Res.*, 2018, **146**, 298–306.
- F. L. Dong, C. Li, J. Crittenden, T. Q. Zhang, Q. F. Lin, G. L. He, W. Q. Zhang and J. M. Luo, *J. Hazard. Mater.*, 2019, **366**, 88–97.
- V. d. J. Gaffney, V. V. Cardoso, M. J. Benoliel and C. M. M. Almeida, *J. Environ. Manage.*, 2016, **166**, 466–477.
- S. Q. Zhou, Y. S. Shao, N. Y. Gao, S. M. Zhu, Y. Ma and J. Deng, *Ecotoxicol. Environ. Saf.*, 2014, **107**, 30–35.
- P. Wang, Y. L. He and C. H. Huang, *Water Res.*, 2011, **45**, 1838–1846.
- M. J. B. Mengelers, P. E. Hougee, L. H. M. Janssen and A. S. J. P. A. M. V. Miert, *J. Vet. Pharmacol. Therapeut.*, 1997, **20**, 276–283.
- B. Li and T. Zhang, *Water Res.*, 2012, **46**, 3703–3713.
- M. R. Rose, S. S. Lau, C. Prasse and J. D. Sivey, *Environ. Sci. Technol. Lett.*, 2020, **7**, 360–370.
- S. Norzaee, M. Taghavi, B. Djahed and F. K. Mostafapour, *J. Environ. Manage.*, 2018, **21**, 316–323.
- L. J. Binkowski and B. Rzonca, *Water, Air, Soil Pollut.*, 2014, **225**, 2217.
- Y. J. Zhu, M. Wei, Z. H. Pan, L. Y. Li, J. Y. Liang, K. F. Yu and Y. Y. Zhang, *Sci. Total Environ.*, 2020, **705**, 135960.
- F. X. Tian, W. K. Ye, B. Xu, X. J. Hu, S. X. Ma, F. Lai, Y. Q. Gao, H. B. Xing, W. H. Xia and B. Wang, *Chem. Eng. J.*, 2020, **398**, 125570.
- L. Jing and B. Ernest, *Environ. Sci. Technol.*, 2009, **43**, 60–65.
- M. R. Yazdani, N. Duimovich, A. Tiraferri, P. Laurell, M. Borghei, J. B. Zimmerman and R. Vahalac, *J. Mol. Liq.*, 2019, **291**, 104353.
- X. X. Cheng, H. Liang, A. Ding, F. S. Qu, S. L. Shao, B. Liu, H. Wang, D. J. Wu and G. B. Li, *J. Membr. Sci.*, 2016, **505**, 15–25.
- X. Wang, B. Qiong, H. L. Zhai, M. Y. Xiong and Y. Liu, *Food Chem.*, 2016, **190**, 1033–1039.
- X. F. Wang, B. H. Zhou and X. Shao, *J. Environ. Sci. Health, Part C: Environ. Carcinog. Ecotoxicol. Rev.*, 2019, **54**, 528–535.
- R. K. Padhi, S. Subramanian and K. K. Satpathy, *Chemosphere*, 2019, **218**, 540–550.
- Y. Du, X. T. Lv, Q. Y. Wu, D. Y. Zhang, Y. T. Zhou, L. Peng and H. Y. Hu, *J. Environ. Sci.*, 2017, **58**, 51–63.
- J. Criquet, E. M. Rodriguez, S. Allard, S. Wellauer, E. Salhi, C. A. Joll and U. V. Gunten, *Water Res.*, 2015, **85**, 476–486.
- H. Y. Dong, Z. M. Qiang, X. J. Yuan and A. J. Luo, *Sep. Purif. Technol.*, 2018, **193**, 415–420.
- Z. L. Chen, A. Li, S. J. Wang, Y. Han, F. Liu and J. M. Shen, *Adv. Mater. Res.*, 2014, **250–253**, 3425–3428.
- J. Wang, Z. N. Hao, F. Q. Shi, Y. G. Yin, D. Cao, Z. W. Yao and J. F. Liu, *Environ. Sci. Technol.*, 2018, **5**, 5662–5670.
- T. Y. Zhang, B. Xu, S. J. Yao, Y. R. Hu, K. F. Lin, H. Ye and C. Z. Cui, *Water Res.*, 2019, **160**, 188–196.
- V. d. J. Gaffney, V. V. Cardoso, M. J. Benoliel and C. M. M. Almeida, *J. Environ. Manage.*, 2016, **166**, 466–477.
- R. Nassar, A. Rifai, A. Trivella, P. Mazellier, S. Mokh and M. A. Iskandarani, *J. Mass Spectrom.*, 2018, **53**, 614–623.
- W. H. Gan, V. K. Sharma, X. Zhang, L. Yang and X. Yang, *J. Hazard. Mater.*, 2015, **292**, 197–204.

