Brief Report

A Universal LC-MS/MS Method for Simultaneous Detection of Antibiotic Residues in Animal and Environmental Samples

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Abstract: Detecting and monitoring the usage of antibiotics is a critical aspect of efforts to combat antimicrobial resistance. Antibiotic residue testing with existing LC-MS/MS methods is limited in detection range. Current methods also lack the capacity to detect multiple antibiotic residues in different samples simultaneously. In this study, we demonstrate a methodology that permits simultaneous extraction and detection of antibiotic residues in animal and environmental samples. A total of 30 different antibiotics from 13 classes could be qualitatively detected with our methodology. Further study to reduce analytes' matrix effect would allow for quantification of antibiotic residues.

Keywords: antibiotic residue; solid-phase extraction; LC-MS/MS

1. Introduction

Antibiotics have been in use for nearly a century and have been an important means to treat and prevent bacterial infection in both humans and animals [1]. However, the misuse and overuse of antibiotics have driven the rapid development of drug-resistant bacteria [2]. This has contributed to the bigger problem of antimicrobial resistance (AMR), which is currently a severe global public health issue. In order to tackle the issue of AMR, a One Health holistic approach, which covers human, animal, and environmental sectors, is necessary, due to inter-sectoral transmission [3].

Under the One Health framework, governments and organizations have taken a primary approach to mitigating AMR by reducing antimicrobial use (AMU) in human and animal sectors [4,5]. However, AMU and antimicrobial consumption have been difficult to measure. AMU survey or antimicrobial procurement data serve as a proxy to measure AMU but are limited by data availability and reliability. Thus, testing of antibiotic residues would be a complementary or alternative option.

In antibiotic residue testing, most of the fundamental laboratory detection methods involve an initial extraction followed by liquid chromatography tandem mass spectrometry (LC-MS/MS) (Figure 1). Although the fine details of the methodology differ between different studies, this fundamental approach has been used to detect the chemical composition of and antibiotic residues in different samples, including porcine muscle [6], duck meat [7], aquaculture products [8], bovine milk [9], milk [10], honey [11], natural water [12,13], swine manure [14], and distiller grains [15].
However, the common limitations of antibiotic residue testing in these studies are the lack of proven applicability of the testing protocol to detect different sample types and the wide number of antibiotic classes used in human and animals. For example, effectively extracting target chemicals from solid samples is different from that of water samples, and the classes of antibiotic that can be detected vary. Moreover, antibiotics and their residues can be unstable. For example, these substances may be unstable in water [16] and could be unstable based on their storage conditions [17,18]. To address these shortcomings, we demonstrate how a single extraction and detection method can be used in AMU surveillance to simultaneously detect multiple antibiotics in both animal and environmental samples.

2. Materials and Methods

2.1. Chemicals and Reagents

Acetonitrile, ammonia solution, citric acid monohydrate, dibasic sodium hydrogen phosphate, and LC-MS grade methanol were purchased from VWR (Radnor, PA, USA) for chemical extraction. The methanol that was used as the solvent in LC-MS was procured from Wako Chemicals (Osaka, Japan). Ethylenediaminetetraacetic acid disodium salt (Na₂EDTA) was purchased from BDH (Radnor, PA, USA), while formic acid was purchased from Fisher Scientific (Waltham, MA, USA). A total of 43 antibiotics belonging to 16 different antibiotic classes were used for this study (Table 1). Amoxicillin, ceftazidime, cefuroxime sodium salt, chloramphenicol, ciprofloxacin hydrochloride, colistin sulfate, doxycline monohydrate, levofloxacin, florfencicol, metronidazole, oxytetracycline, spectinomycin hydrochloride pentahydrate, and sulfadiazine were purchased from Abcam (Cambridge, UK). Mequindox was ordered from Huawen Chemical (Henan, China), while tylvalosin was purchased from Santa Cruz Biotechnology (Dallas, TX, USA). Erythromycin, gentamicin sulfate, tylosin tartrate salt, and trimethoprim were purchased from MP Biomedicals (Irvine, CA, USA). Ampicillin, caffeine-(trimethyl-¹³C₃) solution (used as internal standard), cefalexin, cefquinome sulfate, cefotiofur sodium, chlorotetracycline hydrochloride, clindamycin phosphate, enrofloxacin, kanamycin sulfate, lincomycin hydrochloride, meropenem, neomycin trisulfate salt hydrate, norfloxacin, ofloxacin, penicillin G sodium salt, streptomycin sulfate salt, sulfachloropyridazine, sulfadimidine, sulfamethoxazole, sulframomethoxine, tetracycline, tiamulin, tilimicosin, and vancomycin were purchased from Sigma-Aldrich (Darmstadt, Germany).
Table 1. List of antibiotics tested.

| Classes          | Antibiotics                                |
|------------------|--------------------------------------------|
| Aminoglycosides  | Gentamicin                                 |
|                  | Kanamycin sulfate                           |
|                  | Neomycin trisulfate salt hydrate            |
|                  | Spectinomycin hydrochloride pentahydrate    |
|                  | Streptomycin sulfate salt                  |
| Amphenicols      | Chloramphenicol                             |
|                  | Florfenicol                                |
| Antifolate       | Trimethoprim                               |
| Carabapenems     | Meropenem                                  |
| Cephalosporins   | Cefalexin                                  |
|                  | Cefquinome sulfate                         |
|                  | Ceftazidime                                |
|                  | Ceftiofur sodium                           |
|                  | Cefuroxime                                 |
| Fluoroquinolones | Ciprofloxacin                              |
|                  | Enrofloxacin                               |
|                  | Levofloxacin                               |
|                  | Norfloxacin                                |
|                  | Ofloxacin                                  |
| Glycopeptides    | Vancomycin                                 |
| Lincosamides     | Clindamycin phosphate                      |
|                  | Lincomycin hydrochloride                   |
| Macrolides       | Erythromycin                               |
|                  | Tilmicosin                                 |
|                  | Tylosin tartrate salt                      |
|                  | Tylosaloin                                 |
| Nitroimidazole   | Metronidazole                              |
| Penicillins      | Amoxicillin                                |
|                  | Ampicillin                                 |
|                  | Penicillin G sodium salt                   |
| Pleuromutilins   | Tiamulin                                   |
| Polymyxins       | Colistin A                                 |
|                  | Colistin B                                 |
| Quinoxaline 1,4-di-N-oxides (QdNOs) | Mequindox                                  |
| Sulfonamides     | Sulfoxaminypyradizine                      |
|                  | Sulfadiazine                               |
|                  | Sulfadimidine                              |
|                  | Sulfamethoxazole                           |
|                  | Sulfamonomethoxine                         |
| Tetracyclines    | Chlortetracycline hydrochloride             |
|                  | Doxycycline                                |
|                  | Oxytetracycline                            |
|                  | Tetracycline                               |
2.2. Sample Preparation

Swine feces were collected from local farms and freeze-dried before processing. Pork was purchased from local markets. River water, animal drinking water, and Milli-Q water were used as water samples. An antibiotic mixture (~0.01 mg/mL concentration) containing 43 different antibiotics was used for the spike-in. For solid samples, the antibiotic mixture was directly spiked into 1 g of solid sample (i.e., fecal sample and meat sample) by adding the aqueous antibiotic mixture into a Falcon tube containing the solid sample and then mixing by vortex. The sample was allowed to stand for 1 h to allow for the antibiotic mixture to be absorbed into the solid sample. For liquid samples, the antibiotic mixture was spiked into 100 mL of liquid sample at three volumes: 1 mL of 0.01 mg/mL (i.e., 10 µg), 0.5 mL of 0.01 mg/mL (i.e., 5 µg), and 0.1 mL of 0.01 mg/mL (i.e., 1 µg). All samples were stored at −20 °C and extracted within 1 week.

2.3. Chemical Extraction

Chemical extraction was carried out according to the literature [14] with modifications. The chemical extraction buffer that was used was a mixture of Na$_2$EDTA-McIlvaine buffer solution (10 mL), which contains 10.93 mg/mL anhydrous dibasic sodium phosphate, 12.93 mg/mL citric acid monohydrate, and 37.22 mg/mL Na$_2$EDTA, and 100% methanol (10 mL). For extraction, 20 mL of this chemical extraction buffer was added into 1 g of sample (fecal sample or meat sample). Extraction was carried out in a shaking incubator at 300 rpm for 30 min at room temperature. Supernatants were collected after centrifugation at 3200 g for 5 min at 4 °C. Residues were further extracted two more times with the same protocol. Supernatants of the same sample were then combined and stored at −20 °C until solid-phase extraction (SPE) was performed. For liquid samples, chemical extraction was not performed. All samples were filtered through a 0.2 µm PES syringe filter, purchased from Sartorius (Göttingen, Germany), prior to solid-phase extraction.

2.4. Solid-Phase Extraction (SPE)

An Oasis MAX 6cc (150 mg) cartridge (cat. no.: 186000369), Oasis PRiME HLB 6cc (200 mg) cartridge (cat. no.: 186008057), and Oasis PRiME MCX 6cc (150 mg) cartridge (cat. no.: 186008919) were purchased from Waters (Milford, MA, USA) and used to form a MAX-HLB-MCX combined cartridge for solid-phase extraction. All cartridges were preconditioned with acetonitrile and Milli-Q water separately. To investigate the retention of antibiotics in different SPE cartridges, 10 mL of the antibiotic mixture containing 43 targeted antibiotics was loaded to the MAX-HLB-MCX combined cartridge. For solid samples, supernatants collected from chemical extractions were diluted with Milli-Q water to reduce the methanol content to less than 5% (v/v). All of the diluted supernatant flowed through the MAX-HLB-MCX combined cartridge at a flow rate of 3 mL/min. For liquid samples, all of the samples were passed through the MAX-HLB-MCX combined cartridge directly at a flow rate of 3 mL/min. Analytes were eluted from the MAX cartridge, HLB cartridge, and MCX cartridge separately. To elute the analytes, 4 mL of 2% formic acid in methanol, acetonitrile/methanol (60%/40%; v/v), and 5% ammonia solution in methanol were used, respectively. The elution process was repeated three times. The ratio in which elutes were mixed from the MAX, HLB, and MCX cartridges was 1:1:1. Elutes from the same sample were combined for LC-MS/MS analysis.

2.5. LC-MS/MS Analysis

The Acquity I-Class ultra-high-performance liquid chromatographic system by Waters (Milford, MA, USA), coupled with the QTRAP® 6500+ MS system from AB Sciex (Framingham, MA, USA), was used for LC-MS/MS analysis. A Phenomenex Synergi 4 µm Fusion-RP 80 Å (2 mm × 50 mm) column was used for separation, and the column oven temperature was set at 40 °C. The elution gradient (solvent A: 0.1% aqueous formic acid, solvent B: acetonitrile) was set up as follows: 0 min, 0% B; 0.1 min, 0% B; 1 min, 10% B; 6.5 min, 50% B; 7 min, 100% B; 8 min, 100% B. This gradient was re-equilibrated to 0% B for
2 min after each run. The flow rate was 0.5 mL/min, and the injection volume was 5 µL. In terms of electrospray ionization, the parameters were the following: (CUR), nitrogen, 12; collision gas (CAD), high; electrospray voltage, +5500 V; ion source temperature, 550 °C; curtain gas of 25, CAD gas medium, and gas 1 and 2 of 45 and 50 psi, respectively. Retention time and transitions are shown in Table 2.

Table 2. Liquid chromatography and mass spectrometry of 43 antibiotics.

| Antibiotics              | Retention Time (min) | Transition 1 (m/z)          | Transition 2 (m/z)          | Limit of Detection (ppb) |
|-------------------------|----------------------|-----------------------------|-----------------------------|--------------------------|
| Amoxicillin             | 0.83                 | 366.1 > 348.9               | 366.1 > 208                 | 8.51                     |
| Ampicillin              | 1.78                 | 350 > 191.9                 | 350 > 160                   | 0.49                     |
| Cefalexin               | 1.79                 | 348 > 158                   | 348 > 174                   | 0.9                      |
| Cefquinome sulfate      | 2.33                 | 529 > 396                   | 529 > 134                   | 1.11                     |
| Cefazidime              | 2.14                 | 547.1 > 467.8               | 547.1 > 396                 | 3.44                     |
| Ceftriaxone sodium      | 3.53                 | 524 > 241                   | 524 > 285                   | 0.31                     |
| Cefuroxime              | 2.57                 | 447 > 385.7                 | 447 > 342                   | 4.1                      |
| Chloramphenicol         | 2.56                 | 323.1 > 274.9               | 323.1 > 304.8               | 5.41                     |
| Chlortetracycline hydrochloride | 2.66 | 479 > 444                   | 479 > 462                   | 4.06                     |
| Ciprofloxacin           | 2.35                 | 332.1 > 313.9               | 332.1 > 231.1               | 0.4                      |
| Clindamycin phosphate   | 3.16                 | 505.1 > 457                 | 505.1 > 487.1               | 0.55                     |
| Colistin A              | 1.95                 | 585.6 > 535.5               | 585.6 > 576.4               | 862.53                   |
| Colistin B              | 1.77                 | 578.5 > 528.4               | 578.5 > 569.5               | 793.3                    |
| Doxycycline             | 2.94                 | 445.1 > 428                 | 445.1 > 267                 | 0.49                     |
| Enrofloxacin            | 2.58                 | 360 > 316                   | 360 > 245                   | 0.4                      |
| Erythromycin            | 3.83                 | 734.3 > 576.3               | 734.3 > 157.9               | 14.52                    |
| Florfenicol             | 2.26                 | 358 > 340                   | 358 > 241                   | 13.21                    |
| Gentamicin              | 2.79                 | 500.1 > 456                 | 500.1 > 227.1               | 320                      |
| Kanamycin sulfate       | 0.27                 | 485 > 324                   | 485 > 163                   | 6.54                     |
| Levofloxacin            | 2.36                 | 362.1 > 318.2               | 362.1 > 261.1               | 0.47                     |
| Lincomycin hydrochloride| 1.43                 | 407 > 126                   | 407 > 359                   | 0.45                     |
| Mequitidox              | 2.18                 | 219 > 143                   | 219 > 185                   | 1.7                      |
| Meropenem               | 1.79                 | 384.1 > 340.1               | 384.1 > 297.7               | 1.76                     |
| Metronidazole           | 1.06                 | 172 > 128.2                 | 172 > 82.1                  | 0.46                     |
| Neomycin trisulfate salt hydrate | 0.26 | 615 > 293                   | 615 > 161                   | 345.18                   |
| Norfloxacin             | 2.3                  | 320 > 302                   | 320 > 231.2                 | 0.54                     |
| Olofoxacin              | 2.35                 | 362 > 318                   | 362 > 261                   | 0.43                     |
| Oxytetracycline         | 2.02                 | 461 > 426                   | 461 > 444                   | 1.98                     |
| Penicillin G sodium salt | 2.36 | 335 > 160                   | 335 > 176                   | 10.79                    |
| Spectinomycin hydrochloride pentahydrate | 0.3 | 333 > 189                   | 333 > 140                   | 2.52                     |
| Streptomycin sulfate salt | 4.13 | 582 > 174                   | 582 > 156                   | 425.98                   |
| Sulfachloropyridazine   | 2.3                  | 285 > 156                   | 285 > 108                   | 0.6                      |
| Sulfadiazine            | 1.45                 | 251 > 156                   | 251 > 92                    | 0.45                     |
| Sulfadimidine           | 2.08                 | 279 > 186                   | 279 > 156                   | 0.36                     |
| Sulfamethoxazole        | 2.4                  | 254.1 > 155.8               | 254.1 > 108.2               | 0.43                     |
| Sulfamonomethoxine      | 2.38                 | 281 > 156                   | 281 > 126                   | 0.62                     |
| Tetracycline            | 2.14                 | 445 > 410                   | 445 > 269                   | 0.49                     |
Table 2. Cont.

| Antibiotics              | Retention Time (min) | Transition 1 (m/z) | Transition 2 (m/z) | Limit of Detection (ppb) |
|--------------------------|----------------------|--------------------|--------------------|--------------------------|
| Tiamulin                 | 4.12                 | 494 > 192          | 494 > 119          | 0.59                     |
| Tilmicosin               | 3.38                 | 869.4 > 696        | 869.4 > 174        | 5.73                     |
| Trimethoprim             | 1.92                 | 291.1 > 230        | 291.1 > 260.9      | 0.41                     |
| Tylosin tartrate salt    | 4.13                 | 916.3 > 772        | 916.3 > 174        | 11.63                    |
| Tylvalosin               | 5.2                  | 1042.3 > 814       | 1042.3 > 174       | 64.27                    |
| Vancomycin               | 1.95                 | 726 > 144          | 725 > 144          | 26.63                    |

For transition ranges, each pure antibiotic compound purchased commercially was first injected into the LC-MS/MS instrument for preliminary testing. From this preliminary test, information about transition ranges was obtained. Two transitions of each antibiotic with sharp peaks shown were chosen as references to identify the antibiotic.

2.6. Data Analysis

Data analysis for LC-MS/MS was performed using the SciEX OS-Q Analysis Software (Framingham, MA, USA). Analytes were confirmed by comparing the retention time and the ratio of characteristic transitions between the sample and the standard.

3. Results

3.1. Limit of Detection

This study began with the determination of the detection limit (i.e., limit of detection, LOD) of the MS system. The LOD was determined by injecting a low concentration of antibiotic standard into the mass spectrometer directly and then reviewing the peak generated. If the signal was three times higher than the background base noise level, then we accepted the peak as an actual peak. Infusion was only used for optimizing the MRM parameter before LC-MS/MS. The results of LOD are shown in Table 2. Generally, the detection limit varied. Most of the 43 antibiotics were detectable at levels lower than 70 ppb, with only six antibiotics having a level of detection higher than 70 ppb. These six antibiotics were colistin A, colistin B, neomycin trisulfate, gentamicin, and streptomycin sulfate. Nevertheless, all 43 targeted antibiotics were detectable.

3.2. Solid-Phase Extraction

The percentage of recovery was calculated by comparing the concentration of antibiotic recovered after passing through SPE and without passing through SPE. In other words, the percentage of recovery = total concentration of antibiotic in elutes from SPE/concentration of antibiotic in antibiotic mixture before passing through SPE. The recoveries of amoxicillin, ampicillin, cefquinome sulfate, meropenem, and tiamulin from SPE were poor, with loss being >75% (Table 3). Apart from those which had a poor recovery, 11 antibiotics were retained in the MAX cartridge; five antibiotics were retained in the HLB cartridge; and 17 antibiotics were retained in the MCX cartridge. For chlortetracycline hydrochloride, doxycycline, and mequindox, the MAX cartridge could not completely retain all of the residues, and a significant portion flowed through the MAX cartridge and were retained in the HLB cartridge. For sulfadiazine and sulfadimidine, they could be detected in the elutes of all three cartridges in the MAX-HLB-MCX tandem. Overall, 38 out of the 43 antibiotics had a recovery that was satisfactory or good after SPE.
Table 3. Solid-phase extraction of 43 antibiotics mixture.

| Antibiotics                        | SPE Recovery | Mainly Retained |
|-----------------------------------|--------------|-----------------|
| Amoxicillin                       | Poor         | MAX             |
| Ampicillin                        | Poor         | MAX             |
| Cefalexin                         | Good         | MCX             |
| Cefquinome sulfate                | Poor         | MAX/HLB         |
| Ceftazidime                       | Good         | MAX             |
| Cefitofur sodium                  | Good         | MAX             |
| Cefuroxime                        | Good         | MAX             |
| Chloramphenicol                   | Good         | MAX             |
| Chlortetracycline hydrochloride   | Satisfactory | MAX/HLB         |
| Ciprofloxacin                     | Good         | MCX             |
| Clindamycin phosphate             | Good         | MAX             |
| Colistin A                        | Satisfactory | MCX             |
| Colistin B                        | Satisfactory | MCX             |
| Doxycycline                       | Satisfactory | MAX/HLB         |
| Enrofloxacin                      | Satisfactory | MCX             |
| Erythromycin                      | Good         | HLB             |
| Florfenicol                       | Good         | MAX             |
| Gentamicin                        | Satisfactory | MAX             |
| Kanamycin sulfate                 | Good         | MCX             |
| Levofloxacin                      | Good         | MCX             |
| Lincomycin hydrochloride          | Good         | MCX             |
| Mequindox                         | Satisfactory | MAX/HLB         |
| Meropenem                         | Poor         | MAX/HLB         |
| Metronidazole                     | Good         | MCX             |
| Neomycin trisulfate salt hydrate  | Good         | MCX             |
| Norfloxacin                       | Good         | MCX             |
| Ofloxacin                         | Good         | MCX             |
| Oxytetracycline                   | Satisfactory | MCX             |
| Penicillin G sodium salt          | Good         | MAX             |
| Spectinomycin hydrochloride       | Satisfactory | MCX             |
| Streptomycin sulfate salt hydrate | Good         | HLB             |
| Sulfachloropyridazine             | Good         | MAX             |
| Sulfadiazine                      | Good         | MAX/HLB/MCX     |
| Sulfadimidine                     | Good         | MAX/HLB/MCX     |
| Sulfamethoxazole                  | Good         | MAX             |
| Sulfamonomethoxine                | Good         | MAX             |
| Tetracycline                      | Good         | MCX             |
| Tiamulin                          | Poor         | MCX             |
| Tilmicosin                        | Satisfactory | HLB             |
| Trimethoprim                      | Good         | MCX             |
| Tylosin tartrate salt             | Good         | HLB             |
| Tylosinol                         | Good         | HLB             |
| Vancomycin                        | Good         | MCX             |

Concentration of antibiotics mixture: 10 mL, 0.01 mg/mL. Recovery < 25% is considered to be “Poor”; recovery ≥25% and ≤60% is considered to be “Satisfactory”; and recovery >60% is considered to be “Good”.
3.3. Chemical Extraction

The integrated recovery of antibiotic residues (i.e., including the limitation of chemical extraction, SPE, and LOD) is shown in Table 4. Using our methodology, 30 out of 43 targeted antibiotics could be detected (Table 4). The 13 antibiotics that could not be detected were: amoxicillin, cefquinome sulfate, ceftazidime, cefuroxime, ciprofloxacin, colistin (A and B), gentamicin, kanamycin sulfate, meropenem, neomycin trisulfate, norfloxacin, and vancomycin. Among these 13 undetectable antibiotics, amoxicillin, cefquinome sulfate, and norfloxacin were found to have a poor recovery from SPE.

Table 4. Detection of antibiotic mixture for spike-in of water, fecal, and meat sample.

| Antibiotics | Water Sample | Fecal Sample | Meat Sample |
|-------------|--------------|--------------|-------------|
| Amoxicillin | N.D.         | N.D.         | N.D.        |
| Ampicillin  | Detected     | Detected     | Detected    |
| Cefalexin   | Detected     | Detected     | Detected    |
| Cefquinome sulfate | N.D. | N.D. | N.D. |
| Ceftazidime | N.D.         | N.D.         | N.D.        |
| Ceftriaxone sodium | Detected | Detected | Detected |
| Cefuroxime  | N.D.         | N.D.         | N.D.        |
| Chloramphenicol | Detected | Detected | Detected |
| Chloramphenicol hydrochloride | Detected | Detected | Detected |
| Ciprofloxacin | N.D. | N.D. | N.D. |
| Clindamycin phosphate | Detected | Detected | Detected |
| Colistin A  | N.D.         | N.D.         | N.D.        |
| Colistin B  | N.D.         | N.D.         | N.D.        |
| Doxycycline | Detected     | Detected     | Detected    |
| Enrofloxacin| Detected     | Detected     | Detected    |
| Erythromycin| Detected     | Detected     | Detected    |
| Florfenicol | Detected     | Detected     | Detected    |
| Gentamicin  | N.D.         | N.D.         | N.D.        |
| Kanamycin sulfate mixture of kanamycin A (main component) and kanamycin B and C | N.D. | N.D. | N.D. |
| Levofloxacin| Detected     | Detected     | Detected    |
| Lincomycin hydrochloride | Detected | Detected | Detected |
| Mequitoxin | Detected     | Detected     | Detected    |
| Meropenem   | N.D.         | N.D.         | N.D.        |
| Metronidazole | Detected | Detected | Detected |
| Neomycin trisulfate salt hydrate | N.D. | N.D. | N.D. |
| Norfloxacin | N.D.         | N.D.         | N.D.        |
| Oflofoxacin | Detected     | Detected     | Detected    |
Table 4. Cont.

| Antibiotics                        | Water Sample | Fecal Sample | Meat Sample |
|------------------------------------|--------------|--------------|-------------|
| Oxytetracycline                    | Detected     | Detected     | Detected    |
| Penicillin G sodium salt           | Detected     | Detected     | Detected    |
| Spectinomycin hydrochloride pentahydrate | Detected   | Detected     | Detected    |
| Streptomycin sulfate salt         | Detected     | Detected     | Detected    |
| Sulfachloropyridazine              | Detected     | Detected     | Detected    |
| Sulfadiazine                       | Detected     | Detected     | Detected    |
| Sulfadimidine                      | Detected     | Detected     | Detected    |
| Sulfamethoxazole                   | Detected     | Detected     | Detected    |
| Sulfamonomethoxime                 | Detected     | Detected     | Detected    |
| Tetracycline                       | Detected     | Detected     | Detected    |
| Tiamulin                           | Detected     | Detected     | Detected    |
| Tilmicosin                         | Detected     | Detected     | Detected    |
| Trimethoprim                       | Detected     | Detected     | Detected    |
| Tylosin tartrate salt             | Detected     | Detected     | Detected    |
| Tylvalosin                         | Detected     | Detected     | Detected    |
| Vancomycin                         | N.D.         | N.D.         | N.D.        |

1 mL of antibiotic mixture (0.01 mg/mL) was spiked into 1 g of solid sample or 100 mL of liquid sample. N.D. represents “not detected”.

3.4. Sensitivity

The sensitivities of antibiotic classes were calculated by taking the raw data and then summing the data for all of the tests performed for all antibiotics in a class. Overall, the sensitivity of the protocol in this study was high (i.e., >60%) for 30 out of the 43 antibiotic residues (Table 5). When the spike-in concentration was high, i.e., 10 µg, our approach showed a sensitivity of 100% in most of the antibiotics in all three types of sample. For ampicillin, ceftiofur, chloramphenicol, chlortetracycline, doxycycline, erythromycin, metronidazole, penicillin, spectinomycin, streptomycin, tetracycline, tilmicosin, tylosin tartrate, and tylvalosin, the sensitivities of the protocol were only reduced when the spiked-in antibiotic content was reduced to 1 µg (Table S1).

Out of 16 classes of antibiotic tested, antibiotics from 13 classes could be detected (Table 6). The sensitivity of detecting antibiotics from the antifolate, lincosamide, pleuromutilin, quinoxaline 1,4-di-N-oxide (QdNO), and sulfonamide classes was relatively high in all three types of sample at three different concentrations. The sensitivity of detection of antibiotics from the amphenicol, macrolide, nitroimidazole, and tetracycline classes was high (around 100%) when the spiked-in content was 10 µg and 5 µg, but it was less sensitive when the spiked-in content was reduced to 1 µg. The sensitivity of our detection method was relatively lower for aminoglycosides, cephalosporins, fluoroquinolones, and penicillins, while carbapenems, glycopeptides, and polymyxins could not be detected.
Table 5. Sensitivity of overall protocol for 43 antibiotics at spike-in concentrations of 10 µg, 5 µg, and 1 µg.

| Antibiotics         | Sensitivity 10 µg Spiked-in | Sensitivity 5 µg Spiked-in | Sensitivity 1 µg Spiked-in |
|---------------------|-----------------------------|-----------------------------|-----------------------------|
|                     | Water Sample | Fecal Sample | Meat Sample | Water Sample | Fecal Sample | Meat Sample | Water Sample | Fecal Sample | Meat Sample |
| Amoxicillin         | 0            | 0            | 0           | 0            | 0            | 0           | 0            | 0            | 0           |
| Ampicillin          | 100          | 100          | 100         | 100          | 100          | 100         | 66.7         | 0            | 33.3        |
| Cefalexin           | 100          | 100          | 100         | 100          | 100          | 100         | 100          | 100          | 100         |
| Cefquinome sulfate | 0            | 0            | 0           | 0            | 0            | 0           | 0            | 0            | 0           |
| Ceftazidime         | 0            | 0            | 0           | 0            | 0            | 0           | 0            | 0            | 0           |
| Ceftiofur sodium    | 100          | 100          | 100         | 100          | 100          | 100         | 66.7         | 66.7         | 0           |
| Cefuroxime          | 0            | 0            | 0           | 0            | 0            | 0           | 0            | 0            | 0           |
| Chloramphenicol     | 100          | 100          | 100         | 100          | 100          | 100         | 100          | 100          | 66.7        |
| Chlorotetracycline hydrochloride | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 66.7 |
| Ciprofloxacin       | 0            | 0            | 0           | 0            | 0            | 0           | 0            | 0            | 0           |
| Clindamycin phosphate | 100         | 100          | 100         | 100          | 100          | 100         | 100          | 100          | 100         |
| Colistin A          | 0            | 0            | 0           | 0            | 0            | 0           | 0            | 0            | 0           |
| Colistin B          | 0            | 0            | 0           | 0            | 0            | 0           | 0            | 0            | 0           |
| Doxycycline         | 100          | 100          | 100         | 100          | 100          | 100         | 0            | 33.3         | 0           |
| Enrofloxacin        | 100          | 100          | 100         | 100          | 100          | 100         | 100          | 100          | 100         |
| Erythromycin        | 100          | 100          | 100         | 100          | 33.3         | 100         | 33.3         | 100          | 0           |
| Florfenicol         | 100          | 100          | 100         | 100          | 100          | 100         | 100          | 100          | 100         |
| Gentamicin          | 0            | 0            | 0           | 0            | 0            | 0           | 0            | 0            | 0           |
| Kanamycin sulfate   | 0            | 0            | 0           | 0            | 0            | 0           | 0            | 0            | 0           |
| Antibiotics                        | Sensitivity          | 10 µg Spiked-in | 5 µg Spiked-in | 1 µg Spiked-in |
|-----------------------------------|----------------------|-----------------|----------------|---------------|
|                                   | Water Sample | Fecal Sample | Meat Sample | Water Sample | Fecal Sample | Meat Sample | Water Sample | Fecal Sample | Meat Sample |
| Levofloxacin                      | 100          | 100           | 100         | 100           | 100           | 100         | 100           | 100           | 100          |
| Lincomycin hydrochloride          | 100          | 100           | 100         | 100           | 100           | 100         | 100           | 100           | 100          |
| Mequindox                         | 100          | 100           | 100         | 100           | 100           | 100         | 100           | 100           | 100          |
| Meropenem                         | 0            | 0             | 0           | 0             | 0             | 0           | 0             | 0             | 0            |
| Metronidazole                     | 100          | 100           | 100         | 100           | 100           | 100         | 100           | 33.3          | 66.7         |
| Neomycin trisulfate salt hydrate  | 0            | 0             | 0           | 0             | 0             | 0           | 0             | 0             | 0            |
| Norfloxacin                       | 0            | 0             | 0           | 0             | 0             | 0           | 0             | 0             | 0            |
| Ofloxacin                         | 100          | 100           | 100         | 100           | 100           | 100         | 100           | 100           | 100          |
| Oxytetracycline                   | 100          | 100           | 100         | 100           | 100           | 100         | 100           | 100           | 100          |
| Penicillin G sodium salt          | 100          | 100           | 100         | 100           | 33.3          | 100         | 0             | 0             | 0            |
| Spectinomycin hydrochloride penta hydrate | 100       | 66.7          | 100         | 66.7          | 66.7          | 100         | 0             | 0             | 0            |
| Streptomycin sulfate salt         | 66.7         | 100           | 100         | 33.3          | 0             | 66.7        | 0             | 0             | 0            |
| Sulfachloropyridazine             | 100          | 100           | 100         | 100           | 100           | 100         | 100           | 100           | 100          |
| Sulfadiazine                      | 100          | 100           | 100         | 100           | 100           | 100         | 100           | 100           | 100          |
| Sulfadimidine                     | 100          | 100           | 100         | 100           | 100           | 100         | 100           | 100           | 100          |
| Sulfamethoxazole                  | 100          | 100           | 100         | 100           | 100           | 100         | 100           | 100           | 100          |
| Sulfamonomethoxime                | 100          | 100           | 66.7        | 100           | 100           | 100         | 100           | 100           | 100          |
| Tetracycline                      | 100          | 100           | 100         | 100           | 100           | 100         | 100           | 100           | 66.7         |
Table 5. Cont.

| Antibiotics         | Sensitivity (10 µg Spiked-in) | Sensitivity (5 µg Spiked-in) | Sensitivity (1 µg Spiked-in) |
|---------------------|-------------------------------|-----------------------------|-----------------------------|
|                     | Water Sample                  | Fecal Sample                | Meat Sample                 | Water Sample                  | Fecal Sample                | Meat Sample                 | Water Sample                  | Fecal Sample                | Meat Sample                 |
| Tiamulin            | 100                           | 100                         | 100                         | 100                           | 100                          | 100                         | 100                           | 100                          | 100                          |
| Tilmicosin          | 66.7                          | 100                         | 100                         | 66.7                          | 100                          | 100                         | 0                             | 33.3                         | 33.3                         |
| Trimethoprim        | 100                           | 100                         | 100                         | 100                           | 100                          | 100                         | 100                           | 100                          | 100                          |
| Tylosin tartrate salt | 100                           | 100                         | 100                         | 100                           | 100                          | 100                         | 66.7                          |                              |                              |
| Tylvalosin          | 100                           | 100                         | 100                         | 100                           | 100                          | 100                         | 33.3                          | 100                          | 100                          |
| Vancomycin          | 0                             | 0                           | 0                           | 0                             | 0                            | 0                           | 0                             | 0                            | 0                            |

Sensitivity = \[\frac{\text{number of true positives}}{\text{number of true positives} + \text{number of false negatives}}\] × 100%. Note: the number of tests performed to calculate detection sensitivity was 3.
Table 6. Sensitivity of overall protocol for 16 groups of antibiotics at spike-in concentrations of 10 µg, 5 µg, and 1 µg.

| Antibiotics Group | Sensitivity | Sensitivity | Sensitivity | Sensitivity |
|------------------|-------------|-------------|-------------|-------------|
|                  | 10 µg Spiked-in | 5 µg Spiked-in | 1 µg Spiked-in |
|                  | Water Sample | Fecal Sample | Meat Sample | Water Sample | Fecal Sample | Meat Sample | Water Sample | Fecal Sample | Meat Sample |
| Aminoglycosides  | 33.3 | 33.3 | 40 | 20 | 13.3 | 33.3 | 0 | 0 | 0 |
| Amphenicols      | 100 | 100 | 100 | 100 | 100 | 100 | 50 | 50 | 83.3 |
| Antifolate       | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 |
| Carbapenems      | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Cephalosporins   | 40 | 40 | 40 | 40 | 40 | 40 | 33.3 | 33.3 | 20 |
| Fluoroquinolones | 60 | 60 | 60 | 60 | 60 | 60 | 60 | 60 | 60 |
| Glycopeptides    | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Lincosamides     | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 |
| Macrolides       | 91.7 | 100 | 100 | 91.7 | 83.3 | 100 | 41.7 | 58.3 | 58.3 |
| Nitroimidazole   | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 33.3 | 66.7 |
| Penicillins      | 66.7 | 66.7 | 66.7 | 66.7 | 44.4 | 66.7 | 22.2 | 0 | 11.1 |
| Pleuromutilins   | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 |
| Polymyxins       | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Quinoxaline 1,4-di-N-oxides (QdNOs) | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 |
| Sulfonamides     | 100 | 100 | 93.3 | 100 | 100 | 100 | 100 | 100 | 100 |
| Tetracyclines    | 100 | 100 | 100 | 100 | 100 | 100 | 75 | 83.3 | 58.3 |

Sensitivity = [number of true positives/(number of true positives + number of false negatives)] × 100%.
4. Discussion

In this study, we have developed a LC-MS/MS-based working protocol that could detect residues of 30 antibiotics from 13 classes in animal meat and environmental samples. Although using LC-MS/MS to detect antibiotic residues in food and water samples is not a novel technique, our protocol has two significant improvements. The first improvement is that we have developed a methodology that can cover different sample types. Previous developed methods can detect 34 veterinary drugs from six distinct groups in porcine muscle [6]; 75 antibiotics from six groups in meat and aquaculture products [8]; 63 pharmaceuticals in natural water [12]; 58 antibiotics from eight groups in milk [10]; and 20 antibiotics from three different groups in honey [11]. Most of these methods targeted food samples and are not sufficient in monitoring antibiotic contaminations, particularly in various types of environmental sample.

The second improvement is that we have developed a methodology that can cover a wider range of different antibiotics. Different antibiotics have different chemical properties, and they may require different extraction and detection methods [19,20]. In our study, we used a single extraction and detection protocol to cover 30 antibiotics from 13 families, which makes it an improvement over existing methodologies in terms of efficiency and convenience [7,9,13–15]. To provide an insight into why we could make the aforementioned two improvements in detecting antibiotic residues, we provide an interpretation of the results after SPE and chemical extraction were conducted in our protocol.

In theory, SPE is a common procedure performed to clean up the samples before LC-MS/MS analysis [21,22]. However, the major purpose of using SPE in this study was to concentrate the analytes from large-volume but low-concentration samples, such as environmental water and wastewater with diluted concentrations of antibiotic residues. Thus, in order to increase the sensitivity of detection, SPE was an essential step of our developed method. The chemical nature of the antibiotics and the type of SPE cartridge could influence the final recovery of antibiotic residues for detection. Considering that 43 antibiotics with varying properties were targeted in this study, we used a MAX-HLB-MCX tandem consisting of the MAX cartridge, the MCX cartridge, and the HLB cartridge to retain as much of the antibiotic residues as possible. The rationale behind this tandem formation was that the different cartridges would target compounds with different pH properties: the MAX cartridge targets acidic compounds; the MCX cartridge targets basic compounds; and the HLB cartridge targets relatively neutral compounds (i.e., those that are neither basic nor acidic).

Our SPE method was able to recover a majority of the antibiotics (38 out of 43 antibiotics) with satisfactory or good performance, indicating its ability to successfully detect antibiotics of different pH properties. For those five antibiotics that had a poor recovery after SPE (i.e., >75% loss), we hypothesized that the chemical structure of these antibiotics may not be compatible with the tandem formation that we designed. Most of these belonged to the β-lactam class of antibiotics, of which amoxicillin, ampicillin, cefquinome sulfate, and meropenem possess a chemically unstable β-lactam ring that spontaneously undergoes hydrolysis [23]. Thus, this is a plausible reason explaining why it was very difficult to recover these antibiotics in water samples.

Generally, the extraction method of using Na₂EDTA-McIlvaine buffer solution/methanol (1:1; v/v) provided a detection of 30 different antibiotics. There were three antibiotics that were found to have a poor SPE recovery and could not be detected in spiked-in samples: amoxicillin, cefquinome sulfate, and meropenem. It would not be possible to determine the reason of zero sensitivity for these three antibiotics, since it could be related to the chemical extraction method, poor SPE recovery, or both. Thus, it is difficult to determine whether the three antibiotics could be extracted using our chemical extraction protocol. For the rest of the nine antibiotics that could not be detected, it seems that the extraction method was insufficient in recovering these antibiotics. We arrived at this conclusion because we observed a satisfactory recovery rate from SPE for these antibiotics, and the spike-in concentrations were higher than the LOD. Based on these results, the extraction method may not be suitable.
for extracting the residues of ceftazidime, cefuroxime, ciprofloxacin, colistin, gentamicin, kanamycin sulfate, neomycin trisulfate, norfloxacin, and vancomycin from samples. Further study for modifying the chemical extraction method to further increase the range of recovery may be necessary.

The proposed method has notable limitations, such as the inability to conduct precise quantitative analysis and a decline in sensitivity with lowering the spiked-in concentration. Although SPE is commonly performed to remove impurities from analytes in order to reduce a potential matrix effect, the use of the MAX-HLB-MCX tandem resulted in strong matrix effects. Moreover, it is difficult to obtain corresponding isotopes for all tested antibiotics to correct the matrix effect of each antibiotic during quantification. However, an estimation of the antibiotic concentrations may still be made by performing calculations. From LC-MS/MS results, one will have the information of the total amount of antibiotics in the elute of each sample. The concentration of an antibiotic can be estimated by the following equation: (concentration of antibiotic in sample = amount of antibiotic/sample weight or sample volume). Thus, the difference in volume between solid and liquid samples during the whole process does not affect the estimation of the antibiotic concentration in samples after calculation.

Only 15 out of the 30 detectable antibiotics could maintain a sensitivity of 100% when the spiked-in content was reduced from 10 µg to 5 µg and 1 µg for all three types of sample. It seems that, when the spiked-in concentrations were gradually lowered, the sensitivity of the protocol decreased. From the results in Tables 3 and 4, the loss of antibiotic residues during the extraction process and SPE was expected. The explanation is that the likelihood of a false negative result increased when the residue content decreased. Having stated these limitations, our method could be useful in AMU surveillance in livestock farms as a first-line qualitative assessment tool—especially for detecting residues in farm waste.

5. Conclusions

In conclusion, 30 different antibiotics from 13 classes could be detected with high sensitivity with our sample processing method when the residue content was 10 ppm or above. When the residue content was reduced to 1 ppm, 27 different antibiotics could still be detected, and 21 of them had a sensitivity higher than 50%. The developed chemical extraction method, together with SPE, allowed us to detect at least 30 antibiotic residues from 13 families qualitatively in foods and environmental samples at the same time. Nevertheless, further study to reduce the matrix effect of analytes is necessary so that quantification of antibiotic residues could be possible.

Supplementary Materials: The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/antibiotics11070845/s1, Table S1. Reduction in sensitivity of overall protocol for 43 antibiotics at spike-in concentrations of 10 µg, 5 µg, and 1 µg.

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