Target Analysis of Antibiotic Drugs in Poultry Feedstuff by Solid Phase Extraction and Ultra-High Performance Liquid Chromatography-Tandem Mass Spectrometry

Abdulraaq O. Oyedeji, Nsikak U. Benson*, Akan B. Williams, Titus A. M. Msagati

1Department of Science Laboratory Technology, The Federal Polytechnic, Ilaro, Nigeria.
2Department of Chemistry, Covenant University, Km 10 Idiroko Road, Ota, Ogun State, Nigeria.
3College of Science Engineering and Technology, University of South Africa, Roodepoort, South Africa.
*Corresponding author’s email: nsikak.benson@covenantuniversity.edu.ng

Abstract. Seventy-five poultry feed samples of two feed types were analysed for antibiotic drugs using a simple generic solid phase extraction procedure with dichloromethane-acetonitrile after delipidation with n-hexane. Analytical separation was performed on a Waters Acquity C18 column with gradient elution consisting of water and acetonitrile. Liquid chromatography–tandem mass spectrometry (LC-MS/MS), with positive and negative electrospray ionization methods in the multiple reaction monitoring modes (MRM), was used for the quantification of 21 compounds from six classes including fluoroquinolones, sulfonamides, lincosamides, anthelmintics, macrolides and the β-lactams in a single chromatographic run of 14 minutes. All the six classes of the drugs were found in the two feed types at concentration ranging between 0.22 – 1505 ng/g. Sulfadimethoxine, sulfaguanidine, sulfamerazine, and sulfamethoxazole were the major sulfonamides in the two feed types with concentration at the part per million levels. Albendazole, penicillin-G, sulfadiazine, sulfamethoxazine and sulfisoxazole were not detected in the layers mash exclusively fed to laying birds; also, sulfamethazine and sulfamoxazole were the only two drugs not detected in the growers mash meant for birds raised for meat. Ciprofloxacin, sulfadimethoxine, sulfamethoxazine, sulfamerazine, and sulfaguanidine were the most prominent antibiotic drugs in the two feed types. Results from the present study suggest that feed millers surreptitiously fortify their feeds with antibiotics without declaring same, thus exposing poultry chickens to sub-therapeutic dosages of the drugs. It is evident that self-regulation for safety in the poultry industry should be discouraged thus relevant authorities must take steps to reduce and control the use of antibiotics to protect public health.

Keywords: Poultry feedstuff, Nigeria, Antibiotic drugs, LC-MS/MS
1. Introduction
The use of antibiotics in poultry feeds results in increased growth rate, improvement in egg production and hatchability [1]. Poultry birds, however, retain bacteria strains resistant to the antibiotics they are fed thus flourishing in the intestinal flora as well as the muscle and final transfer to human along the food chain [2]. Several countries have put a ban on the use of antibiotics in feeds because it portends a long-term health hazard to humans by promoting antibiotic-resistant pathogens, an important threat to modern medicine [1,3]. As part of efforts to raise awareness on the incidence of antimicrobial resistance and deliver solutions to fight antibiotic-resistant pathogens, this study examined antibiotic drugs in poultry feeds since the food chain is a significant source of antimicrobial resistance and drug allergic reactions [4].

2. Materials and method
2.1 Standards
Stock solutions of targeted analytes were prepared from commercially available standard antibiotics. 10 mg of each pure standard was dissolved in 10 mL of solvent to prepare 1 mg/mL solution. Working solutions of antibiotic mix were prepared from the stock stored at 4°C.

2.2 Sample Preparation
The samples were prepared in a three stage previously validated method as described by Mu et al. [5]. 0.50 g of feedstuff sample was placed in a mortar separately with 0.05 g Na₂-EDTA and 0.05 g oxalic acid was added to the mortar and gently ground with the sample to obtain a uniform blend.

2.3 SPE Clean-up
Samples were cleaned-up with an ENVI-8® SPE cartridge that had been previously conditioned with 5 mL analytical grade n-hexane (EMSURE®). The sample blend was afterwards introduced onto the cartridge and drained with 6 mL of the n-hexane, and the analytes eluted with 8 mL acetonitrile (LiChrosolv®) – dichloromethane (1:1, v/v). The eluate was evaporated to near dryness under a gentle stream of nitrogen, and the residue re-dissolved in 1 mL 10% methanol (LiChrosolv®) and vortexed. The final solution was filtered with a 0.45 µm disposable syringe (Acrodisc®) and 10 µL of the filtrate injected into the Nexera UHPLC system [5].

2.4 Chromatographic Separation
The chromatographic separation was performed on a Nexera UHPLC system with a Waters Acquity UPLC® BEH C18 column (2.1 mm X 100 mm, 1.7 µm, particle size) with the column compartment maintained at 40°C. Water and acetonitrile with 0.1% formic acid were the mobile phases in gradient elution with a flow rate of 0.2 mL/min as listed in Table 1.

| Table 1: Liquid Chromatography gradient |
|-----------------------------------------|
| Time (min) | A% | B% |
| 0          | 98 | 2  |
| 10         | 0  | 100|
| 12         | 0  | 100|
| 14         | 88 | 12 |

2.5 MS/MS Detection
Sample analysis was performed on Shimadzu Triple Quadrupole Mass Spectrometer (LC-MS 8040®) system using electrospray ionization (ESI) + and – polarity switch. Analytes were monitored in the MRM mode in a dwell time of 100 ms for each channel, and an event time of 0.309 sec to achieve optimal peak shapes and sensitivity.
IOP Conf. Series: Journal of Physics: Conf. Series 1299 (2019) 012102 IOP Publishing

3. Results and discussion

The presence of 21 antibiotic drugs from different classes were established in two poultry feed types collected between May and September, 2017 in Ogun State, Nigeria. Albendazole, penicillin-G, sulfadiazine, sulfaguanidine and sulfixozazole were in the growers mash at mean concentrations, ranging between 1.33±1.50 and 56.18±1.99 ng/g with albendazole and penicillin-G at both extremes. Sulfamethazine and sulfadoxine were present in the layers feed at mean concentrations of 105.68±63.99 and 120.05±206.32 ng/g, respectively. All other 17 antibiotics were in both the growers and layer mash at varying concentrations (Table 3).

Ciprofloxacin, sulfadimethoxine, sulfaguanidine and sulfamerazine were the most prevalent drugs in the two feed types and they occurred at concentrations above 200 ng/g except sulfaguanidine that was 104.96±19.72 and 140.74±10.28 ng/g in the growers and layers mash, respectively. Ciprofloxacin and sulfadimethoxine had higher mean values in layers mash compared with the growers mash. Only sulfamerazine occurred more in the growers mash among the major drugs that were determined. Sulfamethoxazole with a mean concentration of 241.51±206.32 ng/g was higher in the layers mash compared with the 31.53±41.04 ng/g in the growers mash. Sulfamerazin was 77.47±60.45 and 61.19±22.26 ng/g, in the growers and layers mash, respectively.

### Table 2: Analytes, retention time (RT) and MRM transition with collision energies (CE)

| Antibiotics        | RT (Min) | Transition 1 (amu) Ch 1 | Transition 2 (amu) Ch 2 | Transition 3 (amu) Ch 3 |
|---------------------|----------|-------------------------|-------------------------|-------------------------|
| Albendazole*        | 7.243    | 264.5/232.15(8.0)       | 264.5/264.50(26.0)      | 264.5/264.50(35.0)      |
| Ampicillin          | 7.447    | 351.10/160.05(-14.0)    | 351.10/160.15(30.0)     | 351.10/160.70(-25.0)    |
| Azithromycin        | 6.076    | 749.50/591.30(-32.0)    | 749.50/116.10(-48.0)    | 749.50/158.10(-42.0)    |
| Ciprofloxacin       | 3.194    | 333.10/315.15(-21.0)    | 333.10/232.10(38.10)    | 333.10/289.20(-17.0)    |
| Erythromycin        | 6.969    | 734.60/158.15(-33.0)    | 734.60/576.30(-21.0)    | 734.60/83.00(-53.0)     |
| Levofloxacin        | 5.594    | 362.50/261.15(-29.0)    | 362.50/318.15(-19.0)    | 362.50/219.05(-43.0)    |
| Lincomycin          | 5.606    | 407.50/126.15(-30.0)    | 407.50/359.30(-18.0)    | -                       |
| Mebendazole         | 7.419    | 297.10/265.05(-35.0)    | 297.10/256.20(-6.0)     | 297.10/105.05(21.05)    |
| Mebendazole*        | 7.382    | 294.50/262.20(9.0)      | 294.50/294.50(40.0)     | 294.5/294.5(52.0)       |
| Penicillin G        | 4.856    | 367.50/160.0(16.0)      | 367.50/91.10(-48.0)     | 367.50/114.10(-36.0)    |
| Phenoxydazole       | 8.294    | 306/268.00(-21.0)       | 300/159.0(-37.0)        | 300/131.05(-50.0)       |
| Phenoxazol          | 8.294    | 298.2/266.10(11.0)      | 298.2/189.10(31.0)      | 298.2/160.50(50.0)      |
| Sulfadiazine        | 5.875    | 252/156.0(-15.0)        | 252/157.0(-16.0)        | 252/153.10(-29.0)       |
| Sulfadimethoxine    | 7.148    | 311.90/156.05(-23.0)    | 311.90/157.05(-22.0)    | 311.90/105.05(-30.0)    |
| Sulfadimethoxine*   | 7.372    | 309.5/66.15(38.0)       | 309.5/154.0(27.0)       | 309.5/122.0(45.0)       |
| Sulfaguanidine      | 5.162    | 216.0/93.10(-26.0)      | 216.0/157.0(-13.0)      | 216.0/65.50(-17.0)      |
| Sulfamerazine       | 6.104    | 266.0/156.95(-16.0)     | 266.0/93.10(-34.0)      | 266.0/64.5(49.0)        |
| Sulfamerzin         | 6.398    | 282.0/92.15(-30.0)      | 282.0/93.20(-30.0)      | 282.0/157.0(-18.0)      |
| Sulfamethazine      | 6.578    | 280/187.0(-17.0)        | 280/186.0(-17.0)        | 280.0/125.05(-23.0)     |
| Sulfamethoxazone    | 6.578    | 255/93.05(-30.0)        | 255/157.0(-16.0)        | 255/192.15(-30.0)       |
| Sulfamethoxazone*   | 6.496    | 269/156.90(-16.0)       | 269/155.95(-15.0)       | 269/92.05(-30.0)        |
| Sulfamethoxazol     | 6.024    | 301.90/156.05(-18.0)    | 301.90/108.25(-26.0)    | 301.90/92.10(-30.0)     |
| Sulfamethoxazol*    | 5.534    | 299.5/144.15(35.0)      | 299.5/142.20(34.0)      | 299.5/208.30(25.0)      |
| Sulfasalazine       | 7.006    | 400.0/382.10(-20.0)     | 400.0/224.05(-30.0)     | 400.0/119.05(-45.0)     |
| Sulfasalazine*      | 7.006    | 398.1/198.15(24.0)      | 398.1/197.20(35.0)      | 398.1/192.00(50.0)      |
| Sulfadoxine         | 2.639    | 269.00/156.95(-13)      | 269.00/93.00(-27.0)     | 269.00/155.90(-14.0)    |
| Sulfadoxine*        | 2.254    | 267.2/172.1(21.0)       | 267.2/240.1(17.0)       | 267.2/171.05(19.0)      |
| Tylosin             | 6.952    | 916.50/174.05(-41.0)    | 916.50/101.05(-51.0)    | 916.50/145.0(39.0)      |

Precursor (amu)/Product (amu) *Negative electrospray ionization (CE is positive in MeV)
The maximum residue limit for tylosin in feedstuff is 100 mg/kg and has been reported as the most abundant residue in chicken liver [6]; meanwhile, some EU members including Sweden have banned the use of antibiotics in feeds arising from concerns linked to multidrug resistance [7]. Some reports have linked the presence of antibiotic-resistant organisms in poultry products to the use of feeds supplemented with antibiotics [8], and tylosin, erythromycin, penicillin, sulfonamides, fluoroquinolones and lincomycin have been reported to be present at 1 – 200 g per ton of poultry feed [9]. Tylosin was reported to be available in poultry feedstuffs from Belgium at a concentration of 0.85 – 6.32 mg/kg by Huebra and Holst [10], while Poucke et al. [11] had 0.29 – 0.41 mg/kg tylosin in feed samples from the same area. Tylosin is usually used as a growth promoter. Even though, tylosin has been banned by the EU[12], it is allowed as feed additive in China and the United States [13]. Tylosin is, however, present in feedstuffs examined in the present study. According to Diarra et al. [14], supplementation of broiler feedstuff with penicillin resulted in increased body weight among broiler chickens. The drug reportedly improved feed efficiency with reduced feed intake and this improvement was significant, thus, the addition of different antibiotics into poultry rations as seen in the present study may lead to the same effects on the birds. Meanwhile, various types of bacteria have been reported to be resistant against penicillin in chicken and the drug had refused to kill E. coli. Due to increase in antibiotic resistance in foodborne pathogens, there is a campaign to reduce the use of antibiotics at sub-therapeutic levels as growth promoters [15], and approval for the use of antibiotics as growth promoters has been withdrawn in the European Union [16].

The use of antibiotics in feeds could result to selectivedevelopment of resistanceby disease-causing bacteria, and when used as growth-promoters it imposes a selection pressure for bacteria that are resistant to antibiotics that may be used in clinical or veterinary practice, thus compromising the continued use of antibiotics for therapeutic purposes [15,17,18]. Boix et al. [19] reported the presence of α – nandrolone, chlortetracycline, tetracycline, trimethoprim and salicylic acid in poultry feedstuff from Spain. Chlortetracycline, oxytetracycline, and doxycycline were found in poultry feedstuffs from Poland [3]. Poultry feedstuff analysed from the Valencia Region in Spain as part of a residual control plan also showed the presence of growth promoting agents [20]. Growth promoting agents were, however, not determined in the present study. Ampicillin residue was reported in tissues of poultry fed ampicillin medicated drug at 40 mg/kg after a 2-day withdrawal period suggesting that drug residues in animal tissues could arise from their presence in the feedstuff. However, the concentration of the drug was below the 30 ng/g acceptable maximum residue limit for the drug in Japan [21].
Table 3: Distribution of Antibiotic Residues (n=34) in Poultry Feeds by LC-MS/MS

| Antibiotics     | Growers mash | Layers mash | f   | p   |
|-----------------|--------------|-------------|-----|-----|
|                 | Mean±SD      | Min         | Max |     |     |
| Albendazole     | 1.33±1.50    | 0.22        | 2.52| ND  | -   |
| Ampicillin      | 41.49±25.65  | 4.31        | 58.54| 48.49±9.30 | 37.55 | 58.54 | 0.26 | 0.63 |
| Azithromycin    | 10.87±1.10   | 10.29       | 12.02| 23.79±28.59 | 1.8  | 56.11 | 0.61 | 0.48 |
| Ciprofloxacin   | 383.89±410.95| 35.51       | 836.67| 546.12±358.20 | 145.9 | 836.67 | 0.27 | 0.63 |
| Enrofloxacin    | 46.26±13.23  | 32.11       | 58.33| 63.33±7.64 | 56.67 | 71.67 | 3.75 | 0.13 |
| Lincomycin      | 7.73±3.45    | 4.3         | 11.2 | 6.43±1.45 | 4.77  | 7.44  | 0.3  | 0.58 |
| Mebendazole     | 22.67±7.22   | 12.5        | 28.39| 25.74±23.00 | 12.26 | 60.11 | 0.07 | 0.81 |
| Phebendazole    | 4.1±1.10     | 2.85        | 4.89 | 7.63±3.10 | 4.72  | 10.89 | 3.47 | 0.14 |
| Penicillin-G    | 56.18±1.99   | 53.88       | 57.33| ND  | -    | -    | -   | -   |
| Sulfadiazine    | 10.98±0.92   | 10.26       | 12.01| ND  | -    | -    | -   | -   |
| Sulfadimethoxine| 603.56±786.76| 61.311      | 1505.92| 628.31±458.37 | 126.97 | 1025.62 | 0.002 | 0.97 |
| Sulfaguanidine  | 104.96±19.72 | 90.18       | 127.35| 140.74±10.28 | 129.22 | 148.79 | 7.77 | 0.05 |
| Sulfamerazine   | 206.66±29.00 | 173.61      | 227.9 | 105.79±85.37 | 29.83 | 221.29 | 3.7  | 0.11 |
| Sulfamerazine   | 77.47±60.45  | 19.46       | 140.1 | 61.19±22.26 | 44.57 | 86.48 | 0.19 | 0.68 |
| Sulfamethazine  | ND           | -           | -    | 105.68±63.99 | 35.52 | 160.84 | NA  | NA  |
| Sulfamethoxazole| 31.53±14.02  | 6.6         | 78.9 | 241.51±206.32 | 4.89  | 385.67 | 3.00 | 0.16 |
| Sulfamozole     | ND           | -           | -    | 120.05±5.84 | 113.49 | 124.67 | -   | -   |
| Sulfaquinoxaline| 18.96±15.66  | 8.74        | 36.99| ND  | -    | -    | -   | -   |
| Sulfasalazine   | 15.54±8.07   | 6.22        | 27.14| 15.82±9.59 | 0.801 | 24.45 | 0.003 | 0.96 |
| Sulfixozazole   | 16.97±1.02   | 16.25       | 17.69| ND  | -    | -    | -   | -   |
| Tylosin         | 4.04±1.7     | 2.75        | 5.04 | 3.3±0.19 | 3.09  | 3.42  | 1.18 | 0.34 |

4. Conclusions

Results from the present study showed the presence of antibiotic drugs in poultry feeds at sub-therapeutic levels. To address the exposure to these drugs, feed millers must reduce and refine the use of antibiotics, replace antibiotics with alternatives to reduce the threat of antibiotic resistance. Control measures including improvements in food hygiene must be taken by poultry farmers to reduce the spread of zoonotic bacteria to humans via the food chain.

Acknowledgment

The financial assistance provided by the Tertiary Education Trust Fund (Tetfund) of Nigeria to the first author for his doctoral study at Covenant University, Ota is acknowledged. This publication is part of the thesis submitted by the first author for his Ph.D. The publication support by Covenant University is acknowledged.

Conflicts of Interest: The authors declare no conflict of interest. The founding sponsors had no role in the design of the study; in the collection, analyses, or interpretation of data; in the writing of the manuscript, and in the decision to publish the results.
References

[1] Spellberg, B., Bartlett, J.G. and Gilbert, D.N., 2013. The future of antibiotics and resistance. New England Journal Medicine, 368(4), 299-302.

[2] Dye, J., 2012. FDA must act to remove antibiotics from animal feed: Judge. https://www.manitobacooperator.ca/2012/03/29/judge-orders-fda-to-%E2%80%98remove-antibiotics-from-feed%E2%80%99/. Retrieved 21/11/2018.

[3] Patyra, E. and Kwiatek, K., 2017. Development and validation of multi-residue analysis for tetracycline antibiotics in feed by high performance liquid chromatography coupled to mass spectrometry. Food Additives and Contaminants Part A 34(9), 1553-1561.

[4] Chowdhury, R., Haque, M.N., Islam, K.M.S. and A. B. M. Khaleduzzaman, A.B.M., 2009. A review on antibiotics in an animal feed. Bangladesh Journal of Animal Science 38(1&2), 22 – 32.

[5] Mu, H., Yu, H. and Hu, Y-M., 2012. Determination of fluoroquinolones, sulfonamides, and tetracyclinesmultiresidues simultaneously in porcine tissue by MSPD and HPLC-DAD. Journal of Pharmaceutical Analysis 2(1), 76 – 81.

[6] WHO, 2011. Tackling antibiotic resistance from a food safety perspective in Europe, http://www.euro.who.int/__data/assets/pdf_file/0005/136454/e94889.pdf .

[7] Cheng, G., Hao, H., Xie, S., Wang, X., Dai, M., Huang, L. and Yuan, Z., 2014. Antibiotic alternatives: the substitution of antibiotics in animal husbandry? Frontiers in Microbiology 5(217), 1–15.

[8] Diarra, M. S. and Malouin, F., 2014. Antibiotics in Canadian poultry productions and anticipated alternatives. Frontier in Microbiology 5(282), 1 – 15.

[9] Baldrias, L. R., 2015. Antibiotic residues in meat and meat products, implications on human health. Livestock Nutritional Biotechnology: Pre and Pro-biotics in Food Animals. http://www.nast.ph/index.php/downloads/category/62-livestock?download=211:antibiotic-residues-in-meat-and-meat-products-implications-on-human-health. Accessed 15.08.2018.

[10] Huebra, M. J. G and Holst, U. V. C., 2007. Sample preparation strategy for the simultaneous determination of macrolide antibiotics in animal feedstuffs by liquid chromatography with electrochemical detection (HPLC-ECD). Journal of Pharmaceutical and Biomedical Analysis 43, 1628 – 1637.

[11] Poucke, C. V, Dumoulin, F. D., De Keyser, K., Elliott, C., and Van Peteghem, C., 2004. Tylosin Detection in animal feed by Liquid Chromatography -Tandem Mass Spectrometry with enzymatic hydrolysis of the tylosin urea adducts. Journal of Agricultural and Food Chemistry 52, 2803 – 2806.

[12] Situ, C. and Elliott, C.T., 2005. Simultaneous and rapid detection of five banned antibiotic growth promoters by immunoassay. AnalyticaChimicaActa 529, 89 – 96.

[13] Wang, Z., Song, X., Zhou, T., Bian, K., Zhang, F., He, L. and Liu, Q., 2015. Simultaneous determination of ten macrolides drugs in feeds by high performance liquid chromatography with evaporation light scattering detection. RSC Advances, 5, 1491–1499.

[14] Diarra, M. S., Silversides, F. G., Diarrassouba, F., Pritchard, J., Masson, L., Brousseau, . . . and Topp, E., 2007. Impact of feed supplementation with antimicrobial agents on growth performance of broiler chickens, Clostridium perfringens and Enterococcus counts, and antibiotic resistance phenotypes and distribution of antimicrobial resistance determinants in Escherichia coli isolates. Applied and Environmental Microbiology 73(20), 6566–6576.
[15] Kumar, S., Chen, C., Indugu, N, Werlang, G. O., Singh, M., . . . and Kim, W. K., 2018. Effect of antibiotic withdrawal in feed on chicken gut microbial dynamics, immunity, growth performance and prevalence of foodborne pathogens. PLoS ONE 13(2), e0192450. https://doi.org/10.1371/journal.pone.0192450.

[16] Castanon, J. I. R., 2007. History of the use of antibiotic as growth promoters in European poultry feeds. Poultry Science 86, 2466–2471.

[17] Azuma, T.Email Author, Otomo, K., Kunitou, M., Shimizu, M., Hosomaru, K., Mikata, S., Ishida, M., Hisamatsu, K., Yunoki, A., Mino, Y., Hayashi, T., 2019. Environmental fate of pharmaceutical compounds and antimicrobial-resistant bacteria in hospital effluents, and contributions to pollutant loads in the surface waters in Japan, Science of the Total Environment, 657, 476-484.

[18] Wang, Q., Li, X., Yang, Q., Chen, Y., Du, B., 2019. Evolution of microbial community and drug resistance during enrichment of tetracycline-degrading bacteria, Ecotoxicology and Environmental Safety, 171, 746-752.

[19] Boix, C., Ibanez, M. V., Sancho, J. V., Leon, N., Yusa, V., and Hernandez, F., 2014. Qualitative screening of 116 veterinary drugs in feed by liquid chromatography-high resolution mass spectrometry: potential application to quantitative analysis. Food Chemistry 160, 313-20.

[20] Leon, N., Pastor, A. and Yusa, V., 2016. Target analysis and retrospective screening of veterinary drugs, ergot alkaloids, plant toxins and other undesirable substances in feed using liquid chromatography–high resolution mass spectrometry, Talanta 149, 43 – 52.

[21] Hamamoto, K. and Mizuno, Y., 2017. LC-MS/MS measurement of ampicillin residue in chicken tissues at 2 days after in feed administration. Journal of Veterinary Medical Science, 79(3), 474–478.