

Submitted: 07/09/2011

Accepted: 19/10/2011

Published: 24/10/2011

Antibiogram and heavy metal tolerance of bullfrog bacteria in Malaysia

L.W. Tee* and M. Najiah

Faculty of Agrotechnology and Food Science, Universiti Malaysia Terengganu, 21030 Terengganu, Malaysia

Abstract

Bacterial isolates from 30 farmed bullfrogs (*Lithobates catesbeianus*) weighing 500-600 g at Johore, Malaysia with external clinical signs of ulcer, red leg and torticollis were tested for their antibiograms and heavy metal tolerance patterns. A total of 17 bacterial species with 77 strains were successfully isolated and assigned to 21 antibiotics and 4 types of heavy metal (Hg^{2+} , Cr^{6+} , Cd^{2+} , Cu^{2+}). Results revealed that bacteria were resistant against lincomycin (92%), oleandomycin (72.7%) and furazolidone (71.4%) while being susceptible to chloramphenicol and florfenicol at 97.4%. The multiple antibiotic resistance (MAR) index for *C. freundii*, *E. coli* and *M. morgani* was high with the value up to 0.71. Bacterial strains were found to exhibit 100 % resistance to chromium and mercury. High correlation of resistance against both antibiotics and heavy metals was found (71.4 to 100%) between bullfrog bacteria isolates, except bacteria that were resistant to kanamycin showed only 25% resistance against Cu^{2+} . Based on the results in this study, bacterial pathogens of bullfrog culture in Johore, Malaysia, were highly resistant to both antibiotics and heavy metals.

Keywords: Antibiotics, Heavy metal, Bullfrog, Bacteria.

Introduction

Overdosing of antibiotics in feed and excessive use of chemicals in prophylaxis has caused bacteria to become antibiotic-heavy metal resistant. Their residue may stay in the environment and could transfer to other bacteria via antibiotic resistance genes, which are often located in plasmids and transposons (Gillings *et al.*, 2008). Horizontal gene transfer among microorganisms is an important pathway for acquisition of antibiotic and heavy metal resistance in bacterial pathogens. Interaction by the co-resistance of the specific genes can confer resistance to both antibiotics and heavy metals (Baker-Austin *et al.*, 2006; Stepanauskas *et al.*, 2006). Resistance to antibiotics and heavy metals in frogs (*Rana ridibunda*) and rice frogs (*Fejervarya limnocharis*) have been reported (Vogiatzis and Loumbourdis, 1998; Othman *et al.*, 2009).

The hazards present in frog farm, mainly due to use of chemicals for treatment of diseases, stay inherently in the farmed products, and remain a health risk in public concerns (Boyd and Massaut, 1999).

According to the statistical data by FAO (2010), the global production of bullfrogs in 2009 was 1439 tons with the estimated value of 6,007,000 USD, with a 15.4% increase compared to the previous year. Besides, The United Nations' Commodity Trade Statistic Database (United Nations Statistics Division, 2008) reported that major exporting frog legs countries of bullfrog were Indonesia, China, Belgium and Luxembourg. The present study investigated the

antibiogram and heavy metal tolerances of bullfrog bacteria in Malaysia.

Materials and Methods

Samples

Thirty bullfrogs (*Lithobates catesbeianus*) weighing 500-600 g with external clinical signs such as ulcer, red leg and torticollis were brought from a bullfrog farm located at Johore (02°15.549' N, 102°39.261' E). Bullfrogs were euthanized by transdermal exposure to 1.0% solution of buffered MS-222 (McDaniel *et al.*, 2008). Internal organs (liver, kidney, spleen, heart, intestine, lung, ovary and gall bladder) were aseptically – excised and homogenized for 15 min in distilled physiological saline. Two-fold serial dilutions were plated in triplicates on Glutamate Starch Phenol Red Agar (GSP agar), MacConkey agar, Xylose Lysine Deoxycholate Agar (XLD agar), Thiosulfate-Citrate-Bile Salts-Sucrose Agar (TCBS agar), Baird-parker agar and Trypticase Soy Agar (TSA) (Oxoid, England). Plates were incubated at 28°C for 24 to 48 h and counted for colony forming units (cfu) per gram. A total of 17 bacterial species with 77 strains were identified by indole, oxidase, hemolysis tests on horse blood agar and commercial biochemical test, BBL Crystal TM Enteric/ Nonfermenter Identification System (Becton Dickinson, USA).

They were *Acinetobacter lwoffii*, *Aeromonas hydrophila*, *Aeromonas caviae*, *Chryseobacterium indologenes*, *Citrobacter freundii*, *Citrobacter amalonaticus*, *Edwardsiella tarda*, *Elizabethkingia meningoseptica*, *Escherichia coli*, *Escherichia*

*Corresponding Author: Faculty of Agrotechnology and Food Science, Universiti Malaysia Terengganu (UMT), 21030 Terengganu, Malaysia. Tel.: +609 6683508. Fax: +609 6683434. Email: leawee_t@yahoo.com

hermannii, *Morganella morganii*, *Pantoea agglomerans*, *Plesiomonas shigelloides*, *Pseudomonas aeruginosa*, *Serratia liquefaciens*, *Shewanella putrefaciens* and *Stenotrophomonas maltophilia*, as described by Cappuccino and Sherman (2002).

Antibiogram

A total of 21 commercial antibiotic discs were used as follows: ampicillin (AMP 25 µg), amoxicillin (AML 10 µg), chloramphenicol (C 30 µg), colistin sulphate (CT 25 µg), doxycycline (DO 30 µg), erythromycin (E 15 µg), florfenicol (FFC 30 µg), flumequine (UB 30 µg), fosfomycin (FOS 50 µg), furazolidone (FR 15 µg), kanamycin (K 30 µg), lincomycin (MY 15 µg), nalidixic acid (NA 30 µg), nitrofurantoin (F 50 µg), novobiocin (NV 30 µg), oleandomycin (OL 15 µg), oxolinic acid (OA 2 µg), oxytetracycline (OT 30 µg), spiramycin (SP 100 µg), tetracycline (TE 30 µg), and sulphamethoxazole (RL 25 µg) (Oxoid, England). Bacterial suspensions were adjusted to 0.5 McFarland. The antibiotic discs were placed on the surface of the medium by using sterile forceps and incubated at 28°C for 24 h. Diameter of inhibition zones around the discs were measured in millimeter (mm) and characterized as Sensitive (S), Intermediate (I) and Resistance (R) according to Clinical and Laboratory Standard Institute (CLSI, 2006).

Multiple Antibiotic Resistance (MAR) Test

The multiple antibiotic resistance (MAR) index of bacterial strains against antibiotics was calculated based on method used by Krumperman (1983) as follow: $MAR\ index = X / (Y \times Z)$. Where, X: Total bacteria resistant to antibiotics; Y: Total antibiotic used and Z: Total isolates. MAR index value less than 0.20 indicated that the antibiotics are seldom and never used, whereas a value greater than 0.20 suggests that the antibiotics are exposed to the bacteria.

Heavy Metal Tolerance Test

In heavy metal studies, bacterial cultures were grown for 24 h at 37°C on plates containing Trypticase Soy Agar (Oxoid, England) supplemented with Mercuric Chloride (HgCl₂) (Amresco, Ohio) at 2.5 µg/ml, 5.0 µg/ml, 10.0 µg/ml and 20.0 µg/ml; Potassium Dichromate (K₂Cr₂O₇) (Hamburg, Germany) and Cadmium Chloride Anhydrous (CdCl) (Fluka, USA) at 25 µg/ml, 50 µg/ml, 100 µg/ml, 200 µg/ml and 400 µg/ml; Cooper II Sulphate (CuSO₄) (Nacalai Tesque, Japan) at 150 µg/ml, 300 µg/ml, 600 µg/ml, 1200 µg/ml and 2400 µg/ml.

Heavy metal resistant indicative values were: 10 µg/ml for mercury; 100 µg/ml for cadmium and chromium; and 600 µg/ml for copper, respectively (Miranda and Castillo, 1998).

Results

Antimicrobial resistance patterns

Out of the 77 bacterial isolates tested, 71 isolates were resistant to lincomycin (92%), 56 isolates and 55

isolates were resistant to oleandomycin (72.7%) and furazolidone (71.4%), respectively. On the other hand, there were only two isolates (2.6%) resistant to chloramphenicol and florfenicol (Figure 1).

The majority of the bacteria such as *E. meningoseptica*, *C. freundii*, *E. coli*, *E. hermannii*, *E. tarda*, *M. morganii*, *P. agglomerans*, *P. shigelloides*, *P. aeruginosa* and *S. maltophilia* were found to be resistant to lincomycin, followed by oleandomycin (Table 1).

Among the effective antibiotics against bacteria tested were chloramphenicol, florfenicol, kanamycin, doxycycline, nalidixic acid, colistin sulphate and oxolinic acid.

Antibiogram showed that *P. agglomerans* and *S. liquefaciens* were susceptible to up to 16 and 15 antibiotics, respectively. While *P. aeruginosa* was 100% resistant against 5 antibiotics namely oleandomycin, novobiocin, furazolidone, amoxicillin and lincomycin. Besides, it was susceptible to nalidixic acid, colistin sulphate, doxycycline, kanamycin, tetracycline, oxytetracycline and flumequine. *S. putrefaciens* was susceptible to all antibiotics except fosfomycin (Table 2).

Multiple Antibiotic Resistant (MAR) Index

The lowest MAR index value was seen with *S. putrefaciens*, and the highest was found in *C. freundii*, *E. coli* and *M. morganii* as high as 0.71 (Table 3).

The MAR value for *E. meningoseptica*, *E. coli*, *E. hermannii*, *M. morganii* and *P. aeruginosa* isolates was higher than 0.20.

Heavy Metal Tolerance of Bacteria

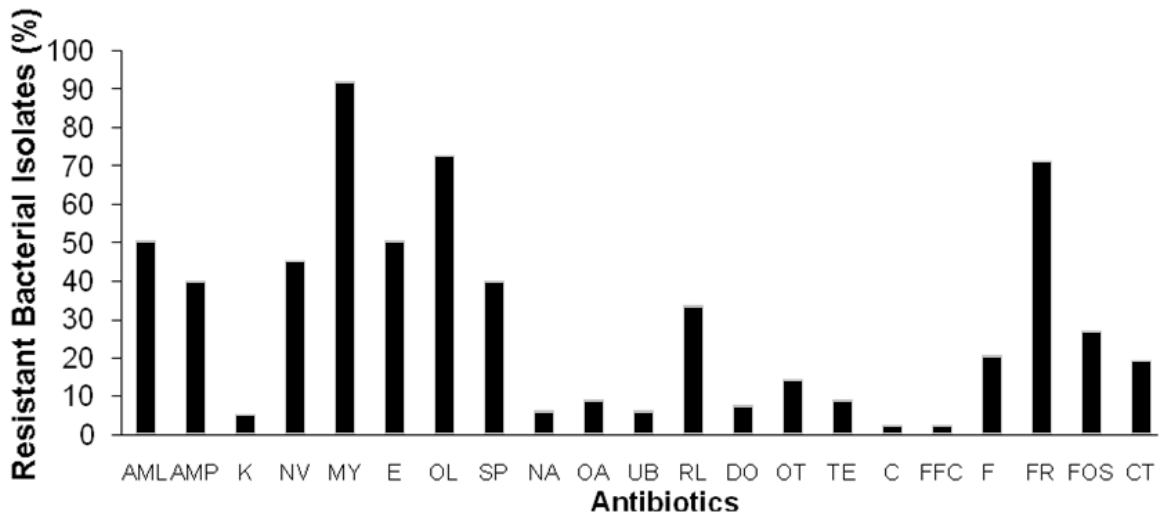
In bullfrog farm, antibiotic multiple-resistance in isolates was distinctly associated with tolerance among heavy metals (Hg²⁺, Cr⁶⁺, Cd²⁺, Cu²⁺).

Isolates were found to be tolerant to different concentrations of heavy metals, ranging from 2.5 to 2400 µg/ml. In our study, heavy metal resistance varies as in the pattern of Hg-Cr>Cd>Cu (Table 4). All the isolates showed 100% resistant to mercury and chromium. There were 89.6% and 76.6% isolates resistance to cadmium and copper, respectively. The maximum heavy metal tolerance of bacteria was found at > 400 µg/ml for copper, and minimum for mercury (20 µg/ml).

Mercury was found to be the most toxic heavy metal with the inhibition concentration of 40 µg/ml for 12 bacterial isolates.

High percentages of resistant patterns among heavy metals and antibiotics were observed. Isolates resistant to heavy metals were also resistant to nalidixic acid, flumequine, doxycycline, chloramphenicol and florfenicol.

The 100% of double-resistant strains were: mercury and chromium to all antibiotics; cadmium to NV, SP, NA, OA, UB, DO, OT, C, FFC, F and CT; and lastly copper to NA, UB, DO, C and FFC (Table 5).



AML:Amoxicillin; AMP:Ampicillin; K:Kanamycin;NV:Novobiocin; MY:Lincomycin; E:Erythromycin; OL: Oleandomycin; SP: Spiramycin; NA: Nalidixic Acid; OA:Oxolinic Acid; UB:Flumequine; RL: Sulphamethoxazole; DO:Doxycycline;OT:Oxytetracycline; TE:Tetracycline; C:Chloramphenicol; FFC:Florfenicol; F:Nitrofurantoin; FR:Flumequine; FOS:Fosfomycin; CT:Colistin Sulphate.

Fig. 1. Percentage of bacterial strains resistance to antibiotics.

Table 1. Percentage of bacterial resistance to various antibiotics.

Organism	Antibiotic Resistance profile (%)																				
	NA	CT	AMP	DO	OL	SP	K	OA	NV	FFC	TE	FR	FOS	OT	AML	F	UB	MY	C	E	RL
<i>A. Freundii</i> (n=1)	0	0	0	0	100	0	0	0	0	0	0	0	0	0	0	0	0	0	0	100	100
<i>A. hydrophila</i> (n=9)	0	0	67	11	89	22	11	11	67	0	0	78	0	11	67	11	11	89	0	30	11
<i>A. caviae</i> (n=1)	0	0	0	0	0	0	0	0	0	100	100	0	0	0	0	0	100	0	0	0	
<i>C. indologenes</i> (n=2)	0	0	50	0	0	0	0	0	0	0	0	0	0	50	0	0	0	0	0	100	
<i>E. meningoseptica</i> (n=4)	0	50	75	0	50	0	50	50	25	0	50	100	50	50	100	50	0	100	0	0	25
<i>C. amalonaticus</i> (n=3)	0	33	0	0	33	33	0	0	0	0	0	67	33	0	0	0	0	67	0	67	0
<i>C. freundii</i> (n=17)	12	6	18	6	82	41	0	6	71	0	6	82	12	12	35	12	6	100	6	41	29
<i>E. coli</i> (n=4)	50	50	75	50	100	75	0	50	50	0	25	75	25	50	75	25	50	100	0	100	75
<i>E. hermannii</i> (n=1)	0	0	100	0	100	100	0	0	100	0	0	0	0	0	100	0	0	100	0	100	0
<i>E. tarda</i> (n=8)	13	25	13	13	100	75	0	0	25	0	13	88	0	13	25	0	13	100	0	88	0
<i>M. morgani</i> (n=10)	0	70	90	10	90	90	0	0	40	0	10	80	90	10	90	70	0	100	0	90	90
<i>P. agglomerans</i> (n=4)	0	0	25	0	0	0	0	0	0	0	0	25	50	0	25	0	0	100	0	0	0
<i>P. shigelloides</i> (n=3)	0	0	33	0	100	0	0	0	33	0	0	33	0	33	67	0	0	100	0	0	33
<i>P. aeruginosa</i> (n=3)	0	0	33	0	100	33	0	33	100	67	0	100	67	0	100	67	0	100	33	67	33
<i>S. liquefaciens</i> (n=2)	0	0	0	0	50	0	0	0	50	0	0	50	50	0	0	0	0	100	0	0	50
<i>S. putrefaciens</i> (n=1)	0	0	0	0	0	0	0	0	0	0	0	0	100	0	0	0	0	0	0	0	0
<i>S. maltophilia</i> (n=4)	0	0	25	0	25	25	25	0	25	0	0	50	25	0	25	25	0	100	0	25	25

AMP: ampicillin (25 µg); AML: amoxicillin (10 µg), C: chloramphenicol (30 µg), CT: colistin sulphate (25 µg), DO: doxycycline (30 µg), E: erythromycin (15 µg), FFC: florfenicol (30 µg), UB: flumequine (30 µg), FOS: fosfomycin (50 µg), FR: furazolidone (15 µg), K: kanamycin (30 µg), MY: lincomycin (15 µg), NA: nalidixic acid (30 µg), F: nitrofurantoin (50 µg), NV: novobiocin (30 µg), OL: oleandomycin (15 µg), OA: oxolinic acid (2 µg), OT: oxytetracycline (30 µg), SP: spiramycin (100 µg), TE: tetracycline (30 µg), RL: sulphamethoxazole (25 µg) (Oxoid, England).

Table 2. Antibiogram patterns of bullfrog bacteria.

Bacteria Species	Antibiotic Resistance Profiles
<i>Acinetobacter lwoffii</i>	OL, E, RL
<i>Aeromonas hydrophila</i> 1	OL, NV, MY, E
<i>Aeromonas hydrophila</i> 2	AMP, OL, SP, NV, FR, AML, MY
<i>Aeromonas hydrophila</i> 3	AMP, DO, OL, SP, NV, FR, OT, AML, MY
<i>Aeromonas hydrophila</i> 4	AMP, OL, FR, AML, MY
<i>Aeromonas hydrophila</i> 5	NV, FR, MY, E, RL
<i>Aeromonas hydrophila</i> 6	AMP, OL, NV, FR, AML, MY
<i>Aeromonas hydrophila</i> 7	AMP, OL, NV, FR, AML, MY
<i>Aeromonas hydrophila</i> 8	AMP, OL, K, AML, MY
<i>Aeromonas hydrophila</i> 9	OL, OA, FR, F, UB, E
<i>Aeromonas caviae</i>	TE, FR, MY
<i>Chryseobacterium indologenes</i> 1	RL
<i>Chryseobacterium indologenes</i> 2	AMP, AML, RL
<i>Elizabethkingia meningoseptica</i> 1	OL, NV, FR, AML, MY, E
<i>Elizabethkingia meningoseptica</i> 2	CT, AMP, K, OA, TE, FOS, OT, AML, F, MY
<i>Elizabethkingia meningoseptica</i> 3	CT, AMP, OL, K, OA, TE, FR, FOS, OT, AML, F, MY
<i>Elizabethkingia meningoseptica</i> 4	AMP, FR, AML, MY, RL
<i>Citrobacter amalonaticus</i> 1	CT, OL, SP, FR, MY, E
<i>Citrobacter amalonaticus</i> 2	FOS, MY
<i>Citrobacter amalonaticus</i> 3	FR, E
<i>Citrobacter freundii</i> 1	OL, SP, MY
<i>Citrobacter freundii</i> 2	OL, SP, NV, FR, AML, MY, E
<i>Citrobacter freundii</i> 3	AMP, OL, SP, NV, FR, AML, MY, E
<i>Citrobacter freundii</i> 4	NA, AMP, DO, OL, SP, OA, NV, TE, FR, OT, AML, UB, MY, E, RL
<i>Citrobacter freundii</i> 5	CT, AMP, OL, SP, FR, FOS, AML, F, MY, E, RL
<i>Citrobacter freundii</i> 6	OL, SP, NV, FR, OT, MY, E, RL
<i>Citrobacter freundii</i> 7	FR, F, MY
<i>Citrobacter freundii</i> 8	MY
<i>Citrobacter freundii</i> 9	OL, SP, NV, FR, AML, MY
<i>Citrobacter freundii</i> 10	NA, OL, NV, FR, MY
<i>Citrobacter freundii</i> 11	NV, FR, MY
<i>Citrobacter freundii</i> 12	OL, NV, FR, MY
<i>Citrobacter freundii</i> 13	OL, FR, AML, MY, E
<i>Citrobacter freundii</i> 14	OL, NV, FR, FOS, MY, C
<i>Citrobacter freundii</i> 15	OL, NV, FR, MY, RL
<i>Citrobacter freundii</i> 16	OL, NV, FR, MY
<i>Citrobacter freundii</i> 17	OL, NV, MY, E, RL
<i>Escherichia coli</i> 1	NA, AMP, DO, OL, SP, OA, NV, TE, FR, OT, AML, UB, MY, E, RL
<i>Escherichia coli</i> 2	NA, AMP, DO, OL, SP, OA, NV, FR, OT, AML, F, UB, MY, E, RL
<i>Escherichia coli</i> 3	CT, OL, FR, MY, E
<i>Escherichia coli</i> 4	CT, AMP, OL, SP, FR, FOS, AML, MY, E, RL
<i>Escherichia hermannii</i>	AMP, OL, SP, NV, AML, MY, E
<i>Edwardsiella tarda</i> 1	OL, SP, NV, FR, AML, MY
<i>Edwardsiella tarda</i> 2	NA, AMP, DO, OL, SP, TE, FR, OT, AML, UB, MY, E
<i>Edwardsiella tarda</i> 3	OL, SP, FR, MY, E
<i>Edwardsiella tarda</i> 4	CT, OL, SP, FR, MY, E
<i>Edwardsiella tarda</i> 5	CT, OL, MY, E
<i>Edwardsiella tarda</i> 6	OL, SP, FR, MY, E
<i>Edwardsiella tarda</i> 7	OL, SP, FR, MY, E
<i>Edwardsiella tarda</i> 8	OL, NV, FR, MY, E
<i>Morganella morganii</i> 1	CT, AMP, OL, SP, NV, FR, FOS, AML, MY, E, RL
<i>Morganella morganii</i> 2	CT, AMP, OL, SP, FR, FOS, AML, F, MY, E, RL
<i>Morganella morganii</i> 3	CT, OL, SP, MY, E
<i>Morganella morganii</i> 4	CT, AMP, DO, OL, SP, NV, TE, FR, FOS, OT, AML, F, MY, E, RL
<i>Morganella morganii</i> 5	CT, AMP, OL, SP, FR, FOS, AML, F, MY, E, RL
<i>Morganella morganii</i> 6	CT, AMP, OL, SP, FR, FOS, AML, F, MY, E, RL
<i>Morganella morganii</i> 7	AMP, OL, SP, NV, FR, FOS, AML, F, MY, E, RL
<i>Morganella morganii</i> 8	AMP, OL, SP, FR, FOS, AML, F, MY, E, RL
<i>Morganella morganii</i> 9	CT, AMP, OL, SP, FR, FOS, AML, F, MY, E, RL
<i>Morganella morganii</i> 10	AMP, NV, FOS, AML, MY, RL
<i>Pantoea agglomerans</i> 1	FOS, MY
<i>Pantoea agglomerans</i> 1	MY
<i>Pantoea agglomerans</i> 1	FOS, MY
<i>Pantoea agglomerans</i> 1	AMP, FR, AML, MY
<i>Plesiomonas shigelloides</i> 1	OL, OT, MY, RL
<i>Plesiomonas shigelloides</i> 2	AMP, OL, NV, FR, AML, MY
<i>Plesiomonas shigelloides</i> 3	OL, AML, MY
<i>Pseudomonas aeruginosa</i> 1	AMP, OL, SP, OA, NV, FFC, FR, FOS, AML, F, MY, C, E
<i>Pseudomonas aeruginosa</i> 2	OL, NV, FR, AML, MY, RL
<i>Pseudomonas aeruginosa</i> 3	OL, NV, FFC, FR, FOS, AML, F, MY, E
<i>Serratia liquefaciens</i> 1	FOS, MY
<i>Serratia liquefaciens</i> 2	OL, NV, FR, MY, RL
<i>Shewanella putrefaciens</i>	OT
<i>Stenotrophomonas maltophilia</i> 1	FOS, MY
<i>Stenotrophomonas maltophilia</i> 2	FR, MY
<i>Stenotrophomonas maltophilia</i> 3	MY
<i>Stenotrophomonas maltophilia</i> 4	AMP, OL, SP, K, NV, FR, AML, F, MY, E, RL

AMP: ampicillin (25 µg); AML: amoxicillin (10 µg), C: chloramphenicol (30 µg), CT: colistin sulphat (25 µg), DO: doxycycline (30 µg), E: erythromycin (15 µg), FFC: florfenicol (30 µg), UB: flumequine (30 µg), FOS: fosfomicin (50 µg), FR: furazolidone (15 µg), K: kanamycin (30 µg), MY: lincomycin (15 µg), NA: nalidixic acid (30 µg), F: nitrofurantoin (50 µg), NV: novobiocin (30 µg), OL: oleandomycin (15 µg), OA: oxolinic acid (2 µg), OT: oxytetracycline (30 µg), SP: spiramycin (100 µg), TE: tetracycline (30 µg), RL: sulphamethoxazole (25 µg) (Oxoid, England).

Table 3. MAR Index for bullfrog bacteria.

Bacteria Species	Multiple Antibiotic Resistance Index
<i>Acinetobacter lwoffii</i> (n=1)	0.14
<i>Aeromonas hydrophila</i> (n=9)	0.19-0.43
<i>Aeromonas caviae</i> (n=1)	0.14
<i>Chryseobacterium indologenes</i> (n=2)	0.05-0.14
<i>Elizabethkingia meningoseptica</i> (n=4)	0.24-0.57
<i>Citrobacter amalonaticus</i> (n=3)	0.10-0.29
<i>Citrobacter freundii</i> (n=17)	0.05-0.71
<i>Escherichia coli</i> (n=4)	0.24-0.71
<i>Escherichia hermanii</i> (n=1)	0.33
<i>Edwardsiella tarda</i> (n=8)	0.19-0.57
<i>Morganella morganii</i> (n=10)	0.24-0.71
<i>Pantoea agglomerans</i> (n=4)	0.10-0.19
<i>Plesiomonas shigelloides</i> (n=3)	0.14-0.29
<i>Pseudomonas aeruginosa</i> (n=3)	0.29-0.62
<i>Serratia liquefaciens</i> (n=2)	0.10-0.24
<i>Shewanella putrefaciens</i> (n=1)	0.05
<i>Stenotrphomonas maltophilia</i> (n=4)	0.05-0.52

Table 4. Incidence of heavy metal tolerance in bacteria from bullfrog farm.

Heavy metal	n	Number of isolates with heavy metal tolerance (µg/ml)														Resistance ^a		
		2.5	5	10	20	40	25	50	100	200	400	150	300	600	1200	2400	n	%
Cadmium	77	-	-	-	-	-	0	5	8	9	30	-	-	-	-	-	69	89.6
Cromium	77	-	-	-	-	-	0	0	0	0	0	-	-	-	-	-	77	100
Copper	77	-	-	-	-	-	-	-	-	-	-	3	7	18	42	77	59	76.6
Mercury	77	0	0	0	0	12	-	-	-	-	-	-	-	-	-	-	77	100

^a Resistance Concentration:: Hg²⁺ (10µg/ml); Cd²⁺ and Cr⁶⁺ (100µg/ml); Cu²⁺ (600µg/ml). n : Number of total isolates; - : Not Tested

Table 5. Expression of antimicrobial activity between antibiotics and heavy metal resistance of bacterial isolates from bullfrogs.

Antibiotic	TNo.	Heavy metal							
		Cd		Cr		Cu		Hg	
		No.	%	No.	%	No.	%	No.	%
AML	39	38	97.4	39	100	33	84.6	39	100
AMP	31	30	96.8	31	100	25	80.6	31	100
K	4	3	75	4	100	1	25	4	100
NV	35	35	100	35	100	32	91.4	35	100
MY	71	65	91.5	71	100	56	78.9	71	100
E	39	37	94.9	39	100	35	89.7	39	100
OL	56	54	96.4	56	100	49	87.5	56	100
SP	31	31	100	31	100	29	93.5	31	100
NA	5	5	100	5	100	5	100	5	100
OA	7	7	100	7	100	6	85.7	7	100
UB	5	5	100	5	100	5	100	5	100
RL	26	24	92.3	26	100	21	80.8	26	100
DO	6	6	100	6	100	6	100	6	100
OT	11	11	100	11	100	8	72.7	11	100
TE	7	6	85.7	7	100	5	71.4	7	100
C	2	2	100	2	100	2	100	2	100
FFC	2	2	100	2	100	2	100	2	100
F	16	16	100	16	100	13	81.3	16	100
FR	55	53	96.4	55	100	46	83.6	55	100
FOS	21	20	95.2	21	100	18	85.7	21	100
CT	15	15	100	15	100	13	86.7	15	100

AML: Amoxicillin; AMP: Ampicillin; K: Kanamycin; NV: Novobiocin; MY: Lincomycin; E: Erythromycin; OL: Oleandomycin; SP: Spiramycin; NA: Nalidixic acid; OA: Oxolinic Acid; UB: Flumequine; RL: Sulphamethoxazole; DO: Doxycycline; OT: Oxytetracycline; TE: Tetracycline; C: Chloramphenicol; FFC: Florfenicol; F: Nitrofurantoin; FR: Furazolidone; FOS: Fosfomicin; CT: Colistin Sulphate; TNo: Number of isolates resistant to particular antibiotic; No: Number of isolates resist to heavy metal and antibiotic; %: Percentage of isolate resistance to antibiotic and heavy metal.

Discussion

Intensive farming of *Lithobates catesbeianus* are always risked with bacterial infection, which is mostly due to the environmental factors (Mauel *et al.*, 2002). In order to reduce bacterial infection, the farmers utilize antibiotics to control and prevent diseases. This has been reported to result in bacterial resistance to various antibiotics and heavy metals (Miranda and Castillo, 1998; Mauel *et al.*, 2002; Akinbowale *et al.*, 2007). In this study, bullfrog bacteria isolated from a local farm in Johore were tested for their antibiotics susceptibility and heavy metal tolerance patterns. Antibiogram results in this study showed that an impressive abundance of bacteria isolated from the diseased bullfrogs were resistant to antibiotic and heavy metals.

In this study, resistance to lincomycin was found to be in around 92% of total bacteria tested, while a total of 72 isolated bacterial strains were all susceptible to chloramphenicol, florfenicol and flumequine. These findings were in contrast with Akinbowale *et al.* (2007), reporting chloramphenicol and florfenicol resistance in *Pseudomonas* spp. isolates. Studies from Mauel *et al.* (2002) were similar to the present results that *A. hydrophila* isolated from bullfrog were resistant to ampicillin, erythromycin and oxytetracycline. However *E. meningoseptica* and *C. indologenes* from the previous study were resistant to erythromycin, in contrast with our results. The difference could be due to types of antibiotics applied in different farms. The continuous use of antibiotics with a high dosage in the farming areas is highly associated with the occurrence of resistant microorganism, probably by the transferring resistant plasmids or integrons (Kümmerer, 2004).

MAR index value was high (>0.2) for many bacterial strains such as *E. meningoseptica*, *E. coli*, *E. hermannii*, *M. morgani* and *P. aeruginosa*. This indicates that antibiotics were commonly used by bullfrog farm at Johore. Furthermore, MAR index value for *E. coli* in the present study was 0.24 to 0.71. Similarly, 0.25 to 0.69 were achieved for *E. coli* isolated from seawater, sediment and shrimp from the south coast of Turkey where the contamination level of domestic waste was high (Fatih *et al.*, 2008). Nevertheless, multiple antibiotic resistance up to 15 types of antibiotics were of special concern. Many of the antibiotics present in the aquaculture area are extrinsic. It is likely driven to the contamination either by run-off or the off-label used (Akinbowale *et al.*, 2007). This may explain the antibiotic resistance problems in the present study.

A heavy metal resistance patterns of Hg>Cr>Cd>Cu in bacterial isolates was observed in the present study, which is different from the heavy metal resistance pattern as Cd>Cu>Hg>Cr for the isolates in a different pollution level in various freshwater sources reported

by Miranda and Castillo (1998). Resistance pattern of *Pseudomonas* spp. and *Aeromonas* spp. isolated from rainbow trout farms in Australia was Cu>Cr>Cd (Akinbowale *et al.*, 2007). The differences could be due to standard stain *E. coli* K12 used. The MIC for copper and chromium were 200 and 800 µg/ml, respectively, in the study by Akinbowale *et al.* (2007), while it was 600 and 200 µg/ml in the present study. All the bacteria were resistant to copper at the concentration of 200 µg/ml in the study by Akinbowale *et al.* (2007), but all bullfrog bacteria were only resistant at the concentration of 2400 µg/ml. Large amount of copper used in bullfrog farm for the treatment of red leg diseases, of this study, could be a reason leading to the high resistance patterns of the bacterial strains in our study. Furthermore, bacterial isolates which are resistant to heavy metals tend to be also resistant to antibiotics. This may be due to the co-location of resistance determinants where specific plasmids carried the resistance genes as defense mechanisms (Stepanauskas *et al.*, 2006).

Usage of antibiotics and chemicals in prophylaxis and treatment of bullfrog culture is becoming problematic. Multiple antibiotic resistances in microorganism arise mainly due to injudicious use of antibiotics in disease treatments. Besides, high antibiotic resistance in bacteria isolated from aquaculture organisms could pose a risk to human health. Therefore, the antibiograms are important to review and revise the empirical disease management used in the aquaculture farm or as indicator of the dissemination of antibiotic elements.

It is well-known that the use of chemotherapeutics in the treatment of bacterial diseases represents a public health hazard. In particular, heavy metals are easily accumulated in the food chain and remain in the muscle tissue. The judicious use of antibiotics and heavy metals by the adoption of best management practices (BMPs) by aquaculturists is essential to reduce the risk of bacterial resistance (Boyd and Massaut, 1999). Dosage, withdrawal period, proper use, storage, disposal, and other constraints on the chemicals including environmental, human and food safety precautions should be followed stringently in reducing those problems.

Acknowledgement

We would like to thank members from Fish Disease Laboratory, Faculty of Agrotechnology and Food Science for their supports in this project.

References

- Akinbowale, O.L., Peng, H.H., Grant, P. and Barton, M.D. 2007. Antibiotic and heavy metal resistance in motile aeromonads and pseudomonads from rainbow trout (*Oncorhynchus mykiss*) farms in Australia. *Int. J. Antimicrob. Agents* 30, 177-182.

- Baker-Austin, C., Wright, M.S., Stepanauskas, R. and McArthur, J.V. 2006. Co-selection of antibiotic and metal resistance. *Trends Microbiol.* 14 (4), 176-182.
- Boyd, C.E. and Massaut, L. 1999. Risks associated with the use of chemicals in pond aquaculture. *Aquacult. Eng.* 20, 113-132.
- Cappuccino, J.G. and Sherman, N. 2002. *Microbiology: a laboratory manual*. 6th Edition. Benjamin Cummings, San Francisco, CA, pp: 1-195.
- Clinical and Laboratory Standard Institute (CLSI). 2006. Performance standard for antimicrobial susceptibility testing; sixteenth information supplement, M-100-S 16. CLSI: Wayne (PA).
- Fatih, M., Aysenur, K. and Sadik, D. 2008. Antibacterial agents and heavy metal resistance in Gram-negative bacteria isolated from seawater, shrimp and sediment in Iskenderun Bay, Turkey. *Sci. Total Environ.* 407, 279-285.
- Food and Agriculture Organization of United Nation. 2010. Fisheries and aquaculture information and statistics service. <http://www.fao.org/figis/servlet/SQServlet?ds=Aquaculture&k1=SPECIES&k1v=1&k1s=10048&outtype=html>. Accessed 16 January 2010.
- Gillings, M., Boucher, Y., Labbate, M., Holmes, A., Krishnan, S., Holley, M. and Stokes, H.W. 2008. The Evolution of Class 1 Integrons and the Rise of Antibiotic Resistance. *J. Bacteriol.* 190 (14), 5095-5100.
- Krumperman, P.H. 1983. Multiple antibiotic resistance indexing of *E. coli* to identify high-risk sources of fecal contamination of foods. *Appl. Environ. Microbiol.* 46 (1), 165-170.
- Kümmerer, K. 2004. Resistance in the environment. *J. Antimicrob. Chemother.* 54, 311-320.
- Mauel, M.J., Miller, D.L., Frazier, K.S. and Hines II, M.E. 2002. Bacterial pathogens isolated from cultured bullfrogs (*Rana castesbeiana*). *J. Vet. Diagn. Invest.* 14, 431-433.
- McDaniel, T.V., Martin, P.A., Struger, J., Sherry, J., Marvin, C.H., McMaster, M.E., Clarence, S. and Tetreault, G. 2008. Potential endocrine disruption of sexual development in free ranging male northern leopard frogs (*Rana pipiens*) and green frogs (*Rana clamitans*) from areas of intensive row crop agriculture. *Aquat. Toxicol.* 88, 230-242.
- Miranda, C.D. and Castillo, G. 1998. Resistance to antibiotic and heavy metals of motile aeromonads from Chilean freshwater. *Sci. Total Environ.* 224, 167-176.
- Othman, M.S., Khonsue, W., Kitana, J., Thirakhupt, K., Robson, M.G. and Kitana, N. 2009. Cadmium Accumulation in Two Populations of Rice Frogs (*Fejervarya limnocharis*) Naturally Exposed to Different Environmental Cadmium Levels. *Bull. Environ. Contam. Toxicol.* 83, 703-707.
- Stepanauskas, R., Glenn, T.C., Jagoe, C.H., Tuckfield, R.C., Lindell, A.H., King, C. J. and McArthur, J.V. 2006. Coselection for microbial resistance to metals and antibiotics in freshwater microcosms. *Environ. Microbiol.* 8 (9), 1510-1514.
- United Nations Statistics Division. 2008. Trade of Goods, US\$, HS 1992, 02 meat and edible meat offal. http://data.un.org/Data.aspx?q=frog+leg&d=ComTrade&f=_11Code%3a3%3bcmdCode%3a020820. Accessed on 21 July 2011.
- Vogiatzis, A.K. and Loumbourdis, N.S. 1998. Cadmium accumulation in liver and kidneys and hepatic metallothionein and glutathione levels in *Rana ridibunda*, after exposure to CdCl₂. *Arch. Environ. Contam. Toxicol.* 34, 64-68.