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SOME STUDIES ON VIBRIOSIS IN FARMED MUGIL CEPHALUS IN DAKAHLIA GOVERNORATE

(With 5 Tables)

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بعض الدراسات على الإصابة بمرض الفيبريوزس في أسماك البورى المستزرعة في محافظة الدقهلية

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أجريت هذه الدراسة على عدد 80 سمكة من أسماك البورى المستزرعة فى محافظة الدقهلية وبالفحص الظاهرى تبين وجود أنزفة واحتقان على الجلد و تآكل الزعانف وبالفحص البكتيريولوجى للعينات كانت نسبة تواجد ميكروبات الفيبريو المعزولة 33.75% التى صنفت مورفولوجيا وبيوكيميائيا إلى 16 (59.25%) فيبرويو أنجويلارم نوع سي، 8 (29.62%) فيبريو أوردلى، 5(18.51%) فيبريو دامسيلا و 3(11.11%) فيبريو فالنيفيكس وبدراسة مدى ضراوة ميكروبات الفيبريو انجويلارم نوع سى فى أسماك البورى الطبيعيه كانت نسبة النفوق عالية (80%) بين الأسماك التى تم حقنها وبإجراء اختبار الحساسية للمعزولات البكتيرية وجد أنها حساسة لكل من الأنروفلوكساسين ، سلفات الكوليستين، نيتروفيوران، أوكسى تتراسيكلين، تتراسيكلين وسيفوتاكسيم.

SUMMARY

80 naturally infected farmed *Mugil Cephalus* were collected from a private fish farm at Dakahlia Governorate, revealed clinically congestion and haemorrhages on skin and fins rot. The fish were examined bacteriologically for detection of Vibrio species. The obtained results revealed that 27 (33.75%) were positive for Vibrio species which identified morphologically and biochemically to *V. anguillarum* biotype C 16 (59.25%), *V. ordalii* 8(29.62%), *V. damsela* 5 (18.51%) and *V. vulnificus* 3(11.11%). The pathogencity of isolated *V. anguillarum* biotype C strains in Mugil fish revealed high mortality (80%) in experimentally healthy fish. Sensitivity test of the isolated strains showed that Vibrio spp. were sensitive for Enrofloxacin, Colistin sulphate, Nitrofurantoin, Oxytetracycline, Tetracycline and Cefotaxim.

Key words: Mugil cephalus, vibriosis, fish diseases

INTRODUCTION

Fish are regarded as being most popular and more perishable than other high protein foods. The flesh of healthy fish is considered bacteriologically sterile. However, they are sometimes contaminated with bacterial pathogens and thus can inflect heavy losses in fish and causing diseases in man. Vibriosis is an enzootic disease of fish allover the world. It occurs among various fish species predominatly in marine water, brackish water and freshwater fish. (Hacking and Budd, 1971; Kitao *et al.*, 1983).

Losses from vibriosis have attained considerable importance and become a serious threat to fish production especially with the increasing utilization of sea, brackish and inland waters near the coast to cultivated and fatten fish of various species (Schaperclaus *et al.* 1992). Epizootics of vibriosis take place in fish in presence of overcrowding, poor hygiene and organical polluted water (Kitao *et al.*, 1983; Noga, 1995). Vibrio is Gram-negative, non spore forming, facultative anaerobe and rod shaped bacterium, either currved or straight, it is motile by a single polar flagellum. Vibrio species grow in a wide temperature $5 - 35^{\circ}$ C and rarely at 37° C.It grows well on most common laboratory media in the presence of 3 - 10% NaCl. (Cowan *et al.*, 1975).

Family Vibrionaceae including V. anguillarum, V. damsela, V. ordalii, V. vulnificus V. alginolyticus, V. fischeri and V. fluvialis (Oliver and kaper, 1997 and Hurley *et al.* 2006). Vibrio anguillarum is the most common fish pathogen that affecting fresh water as well as marine fishes (Hacking and Budd, 1971); Rock and Nelson, 2006).

Some Vibrios produce hemolysin which may cause anemia and proteases which may cause muscle damage, reducing the keeping quality, marketability of fish and so economically losses (Hjeltnes and Roberts, 1993).

Vibriosis among various fishes cause acute, subacute and chronic infection as well as external signs as erythema at the base of fins, in the mouth and along the grooves of the lower jaw (Bullock, 1987).

This investigation was planned to study the prevalence of vibrio species in *Mugil cephalus* fish, isolation and identification of the recovered Vibrios, pathogenicity to such fish and in vitro sensitivity to antibiograms.

MATERIALS and METHODS

Fish:

A total number of 80 diseased fish (*Mugil cephalus*) (170g. \pm body weight) were collected from private fish farms in Dakahlia Governorate. Diseased fish showed wide spread skin and fin haemorrhages, sloughed skin, fin rot and anal congestion.

Fish specimens were transferred to laboratory and bacteriologically examined.

Bacteriological examination:

Specimens of fish gills, skin, liver, spleen and kidneys were taken under complete aseptic precautions for bacteriological examination of vibrio species according to Schaperclaus et al. (1992) and Austin and Austin, (1993). The samples were inoculated into Brain Heart Infustion (BHI) broth and peptone water containing 3% sodium chloride tubes and adjusted at pH 8.5 and incubated aerobically at 25°C over night, loopfuls from the inoculated tubes were streaked on (BHI) agar with 3% sodium chloride. Moreover, Thiosulphate Citrate Bile Salt Sucrose agar (TCBS) was also used as a selective media and incubated at 25°C for 24 hours according to Inglis et al. (1993) and Quinn et al. (1994). The typical colonies were picked up on Trypticase Soya Agar (TSA) slant with 3% NaCl and incubated at 25°C for 24 hours. The isolates were morphologically and biochemically identified by Gram-stain, oxidase and catalase tests, motility, carbohydrate fermentation, TSI slant and other biochemical tests according to Overman et al. (1985) and Elliot et al. (1995).

Experimental infection:

The isolates were grown separately on BHI broth for 24 hours, then 0.2 ml dose (5X10⁵ CFU/ml) was intraperitoneally injected to *Mugil cephalus* fish (5 fish for each isolate).

Furthermore 5 fish were used as a control group. The inoculated fish were observed during 3-weeks for the development of pathological changes.

Reisolation of the inoculated organism from internal organs of freshly dead fish was carried out.

Antibiogram activity:

Bacterial isolates were tested for their susceptibility towards ten antibacterial agents according to Koneman *et al.* (1992) using the following drugs; Amoxycillin, Ampicillin, Enrofloxacin Colistin sulphate, Nitrofuration, Cefotaxime, Oxytetracycline, Erythromycin,

Assiut Vet. Med. J. Vol. 54 No. 118 July 2008

Lincomycin and Tetracycline. The interpretatrion of results was carried according to Bio-Merieux Manual (1986).

RESULTS

Table	1:	Biochemical	properties	of	the	isolated	vibrio	used	for
		identification							

	Biochemical properties											
Vibrio Strain	Oxidase	Catalase	V. Proskauer	Gelatin liquification	H2S production	Arginin decomposition	Lysine decarboxylase	Acid from sucrose	Acid from arabinose	Acid from glucose	Acid from manitol	Acid from lactose
V. anguillarum	+	+	+	+	-	+	-	+	<u>+</u>	+	+	-
V.ordalii	+	+	-	+	-	+	-	+	-	+	+	-
V. damsela	+	+	-	<u>+</u>	-	+	<u>+</u>	-	-	-	-	-
V. vulnificus	+	+	-	-	<u>+</u>	-	+	<u>+</u>	-	-	-	<u>+</u>
+ = positive $- = Negative$ $+ = positive or negative$												

Table 2: Prevalence of Vibrio species in *Mugil cephalus*.

No. of	Positive	samples	Vibrio species						
No. of examined samples	No. %		V. anguillarum biotype C	V. ordalii	V. V.vulnificus damsela group 2				
80	27	33.75	16	8	5	3			

Table 3: Recovery rate of Vibrio species among various organs of naturally infected *Mugil cephalus*

Vibrio	Total isolates		Gills	%	Liver	%	Spleen	%	Kidneys	%
species	No.	%		,.		,.	~	,.		,.
V. anguillarum biotype C	16	59.25	3	18.75	7	43.75	4	25.00	2	12.50
V. ordalii	8	29.62	2	25.00	3	37.50	2	25.00	1	12.50
V. damsela	5	18.51	1	20.00	3	60.00	1	20.00	0	0.00
V. vulnificus group 2	3	11.11	0	0.00	2	66.66	1	33.33	0	0.00
Total	32		6	18.75	15	46.87	8	25.00	3	9.37

	No. of used	No. of	Total No. of	Total mortalities			
Vibrio species	isolates	inoculated fish/isolate	inoculated fish	No.	%		
V. anguillarum biotype C	2	5	10	8	80.00		
V.ordalii	2	5	10	7	70.00		
V. damsela	2	5	10	6	60.00		
V. vulnificus group 2	2	5	10	4	40.00		
total	8		40	25	62.50		

Table 4: Results of I/P experimental infection of Vibrio species in Mugil cephalus.

Table	5:	Sensitivity	of	the	isolated	Vibrio	species	to	different
antibiograms									

Antibiograme	Disc concentration	V. anguillarum biotype C	V. ordalii	V. damsela	V. vulnificus group 2
Amoxycillin	10 ug	R	R	S	S
Ampicillin	10 ug	R	R	R	R
Enrofloxacin	10 ug	S	S	S	S
Colistin sulphate	10 ug	S	S	S	S
Nitrofurantion	300 ug	S	S	S	S
Cefotaxime	30 ug	S	S	S	S
Oxytetracycline	30 ug	S	S	S	S
Erythromycin	15 ug	R	S	S	S
Lincomycin	2 ug	R	R	R	R
Tetracycline	30 ug	S	S	S	S

S: sensitivity of the studies isolated of each vibrio species to antibiograme was > 50%. **R**: Sensitivity of the studies isolates of each vibrio species to antibiogram was < 50%.

DISCUSSION

Vibrio spp. is a natural inhabitant of the fish. These organisms are considered food borne pathogens able to contaminate the fish causing world health problems and economic loss in fish industry. Not all strains of vibrio are considered pathogenic strains except that produce thermostable direct hemolysin (Bag *et al.*, 1999).

The data presented in Table (2) indicated that the prevalence of vibrio species in *Mugil cephalus* was 33.75%. The obtained prevalence are nearly similar with reported by Abd El-Gaber *et al.*, (1997) who isolated vibrio species 37.50% from *Mugil cephalus*. In this study, recovered vibrio species were *V. anguillarum* biotype C 16 (59.25%), *V. ordalii* 8(29.62%), *V. damsela* 5 (18.51%) and *V. vulnificus* groups 2, 3 (11.11%). On the other hand, Abd El-Gaber *et al.*, (1997) isolated *V. anguillarum* biotype C. (34.37%), *V. ordalii* (28.12%), *V. damsela* (21.87%) and *V. vulnificus* group 2 (15.62%) from *Mugil cephalus* fish.

In the present study *V. anguillarum* which constituted the highest prevalence rate 16(59.25%) was recovered from *Mugil cephalus* as shown in Tables (2) and (3). This is in nearly agreement with most other studies Muroga and Egusa 1988; Rock and Nelson 2006 and Chai-Yingmei *et al.*, 2006). Such results were high as compared with those reported by Abd El-Gaber *et al.*, 1997) who isolated *V. anguillarum* biotype C 5 (15.62%) from *Mugil cephalus* while high results were recorded by Moustafa *et al.*, (1990) who isolated *V. anguillarum* type A from 74.00% of Mullet fish (*Mugil cephalus*).

The highest isolation rate could be attributed to environemental stresses particularly high water temperature, organically polluted water, high salinity and poor hygiene and handling resulting in depression of one or several defensive mechanisms (Ellis, 1981). This finding was supported by Moustafa *et al.*, (1990) who recorded that water pollution and high salinity were the major stress factors for occurrence of vibriosis among fishes.

Concerning the site of isolation from *Mugil cephalus* fish, vibrios were high from liver 15 isolates (46.87%) followed by spleen 8 isolates (25.00%), gills 6 isolates (18.75%) and kidneys 3 isolates (9.37%) Table (3) .On the other hand, Abd El Gaber *et al.*, (1997) isolated Vibrios from liver, spleen, kidneys and gills of *Mugil cephalus* fish with percentage of 35.00, 37.50, 30.00 and 22.50% respectively.

Regarding to the experimental infection of *Mugil cephalus* with different vibrio species Table (4), exhibited a septicaemic picture within one week post intraperitoneal injection where 62.50% of inoculated fish were dead. Mortality rate ranged from 40.00% with *V. vulnificus* gp. 2 to 80.00% with *V. anguillarum* biotype C. while *V. ordalii* produced 70.00% mortalities and *V. damsela* 60.00% These findings were supported by Abd El-Gaber *et al.*, (1997) who recorded 70% mortalities

among *O. niloticus* with different Vibrio species. El-Bouhy *et al.*, (1990) found 80% mortalities was reported after interperitoneal infection of Nile catfish with *V. anguillarum* and Amany *et al.*, (2000) recorded 90% mortalities among C. lazera post. interaperitoneal infection of *V. anguillarum*.

Moreover, clinical signs and gross lesions induced by Vibrios in the present sudy were nearly similar to thoses observed in Abd El Gaber *et al.*, (1997).Nearly similar observations were also recorded in other fish species which had Vibriosis Austin and Austin (1989); Lavilla pitogo *et al.*, (1992) and Schaperclaus *et al.*, (1992).

As shown in Table (5), Vibrio species were sensitive to Enrofloxacin, Colistin sulphate, Nitrofuratoin, Oxytetra-cycline, Tetracycline and Cefotaxime. They resistance to Lincomycin and Erythromycin. These results nearly agreed with that recorded by different authors (Balsgaard and Bjerregaard 1991; Austin and Austin 1993; Shaahan *et al.* 1995; Yonis *et al.*, 1997; Abd El Gaber *et al.*, 1997; Zeinab Soliman 1999 and Amany *et al.*, 2000) who found that the isolated *V. anguillarum* strain was sensitive to Ampicillin and Chlormphenicol, while it was resistant to Erythromycin Joklik *et al.* (1992) and Stephens *et al.* (2006) recorded that most isolates of *V. damsela* were sensitive to Tetracycline.

It was concluded that Vibrio species existed in the examined fishes in varying percentages. *V. anguillarum* is the most serious pathogen of freshwater fish (*M. cephalus*) and it was highly pathogenic when injected in healthy ones and prevent the infections by Good handling, hygienic measurement and prevention the source of water pollution, also administration of the effective drug to fish should be carefully controlled on Vibriosis.

REFERENCES

- Abdel-Gaber, G.; Naguib, M. and Abdel-Aziz, E.S. (1997): Vibrio species infections to Oreochromis niloticus and Mugil cephalus; Sodium chloride tolerance, pathogenicity, serological relatedness and antibiogram sensitivity of recovered vibrios. Vet. Med. J., Giza. 45 (1): 87-99.
- Amany, A. abbass; Shaheen, A.A; Abd El-Azizi, A. Mosaad and Mona, M. Sobhy (2000): Clinicopathological and laboratory investigations on vibriosisi in some fishes. Zag. Vet. J. (ISSN. 1110-1458) Vol. 28, No. 3, PP. 115 – 124.

- Austin, B. and Austin, D.A. (1989): Methods for microbiological examination of fish and shellfish. Ellis Harwood Limited ,Chichester, England,p.317.
- Austin, B. and Austin, D.A. (1993): Bacterial Fish Pathogens. Disease in Farmed and wild Fish. 2nd. ed, Printed and bound in Great Britain by Hartnolls,. Bodmin.
- Bag, P.; Nandi, S.; Bhardra, R.; Ramamurthy, T; Bhattacharya, S; Nishibuchi, M.; Hamabota, T.; Yamasaki, S.; Takeda, Y. and Nair,G. (1999); Clonal diversity among recently emerged strains of Vibrio parahaemolyticus O3:K6 associted with pandemic spread. J. Clic. Microbiol. 37: 2345-2357.
- Balsgaard, I. and Bjerregaard, J. (1991): Enrofloxacin an antibiotic in fish. Acta Vetrinaria Scendinavica Supplementum. 87, 300-301.
- *Bio-merieux (1986):* Laboratory reagents and products Bacteriology Barcy-L. Etoile 69260 charbon-mieres Le-Bains, France.
- Bullock, G. (1987): Vibriosis in fish. United State Department of the Interior, Fish and Wildlife Service, Division of Fisheries and Wetlands Research, Washington, D.C. 20240
- Chai- Yingmei; Huang-XiaoHang; Cong-Bailin; Liu-Sheng Hao; Chenkui; Li-Guang you and Gaisano, H.Y. (2006): Involvement of Vamp- 2 in exocytosis of IL- I beta in turbot (Scophthalmus maximus) Leukocytes after Vibrio anguillarum. Biochemical and Biophysical Research communication. 342(2): 509 – 513.
- Cowan, S.T.; Holt, J.G.; Liston, J.; Murray, R.G.E.; Niven; C.F.; Ravin, A.W. and Stanier, R.W. (1975): In Buchanen, R.E. and N.E. Gibbons. Eds. Bergey's manual of determinative bacteriology 8th Ed. Williams & Wilkins Co., Baltimore, MD, 1, 268p.
- El-Bouhy, Z.M.; Abdel-Monem, AA.; Mohamed, E. and Moustafa, M.B. (1990): Preliminary studies on vibrios in some freshwater fishes. Zagazig Vet. J. 18 (5): 68-86.
- Elliot, E.; Kaysner, C.; Jackson, L. and Tamplin, M. (1995): Vibrio Cholera, V. parahaemolyticus, V. vulnificus and other Vibrio spp. pp. 9.01-9.27. In FDA Bacteriological Analytical Manual, 8th Ed. AOAC International, Gaithersburg, MD.
- *Ellis, A.E. (1981):* Stresses and modulation of the immune response in fish. In stress and fish, Academic press, London. PP. 147-169.
- Hacking, M.A. and Budd, J. (1971): Vibrio infection in tropical fish in a freshwater aquarium. J. Wildlife Dis. 7: 273-280.

- Hjeltnes B. and Robots, R.J. (1993): Bacterial diseases of fish, New York, Halsted Press, PP. 109-121.
- Hurley, C.; Quirke, A.; Reen, F. and Boyd, E. (2006): Four genomic islands that mark post-1975 pandemic Vibrio parahaemolyticus isolates. BMC Genomic, 3 (7): 104.
- Inglis, V.; Oberts, R.J. and Bromage, N.R. (1993): Bacterial diseases of fish. Oxford Blackwell Scientific Publications, London, Edinburgh, Bosten, Mellbournce. Paris, Berlin, Vienna.
- Joklik, W.K.; Willett, H.P.; Amos, D.B. and Wilfert, C.M. (1992): Zinsser Microbiology. 20th Edition. Appleton & Lange. Norwalk, Connecticut/San Mateo, California.
- Kitao, T.; Roberts, R.G. and Bromage, N.R. (1983): Serotyping of Vibrio anguillarum isolated from fresh water fish in Japan. J. Fish Dis., 6: 175 – 181.
- Koneman, E.W.; Allen, S.D.; Janda, W.M.; Schreckenberger, P.C. and Winn, W.C.Tr. (1992): Colour Atlas and Textbook of Diagnostic Microbiology. 4th Ed., L[pincott company, Philadelphia USA.
- Lavilla-Pitogo, C.R.; Castillo, A.R. and De La Cruz, M.C. (1992): Occurrence of Vibrio species infection in Grouper, Epinephelus suillus. J. Appl. Ichthyol., 8 (1/4), 175-179.
- Moustafa, M.; Eissa, I.A.M. and Hanafi, M.S. (1990): Vibriosis in marine fishes of Qarun lake. Zagazig Vet. J., 18 (195):94-105.
- Muroga, K. and Egusa, S. (1988): Vibriosis of ayu. J. Appl. Biolog. Sci., Hiroshima University, 27 (1): 1-17.
- *Noga, E.J. (1995):* Fish disease: Diagnosis and treatments. 1st ed., Wals worth Publishing Co., USA. PP. 149.
- Oliver, J. and Kaper, J. (1997): Vibrio species. In M. P. Doyle, L. R. Beuchat and T.J. Montville, Eds. Food Microbiology: Fundamentals and Frontiers, p 228-264. Washington, D.C., ASM Press.
- Overman, T.; Kessler, J. and Seabolt, J. (1985): Comparison of API20E, API Rapid E and API. Rapid NFT for identification of members of the family Vibrionaceae. J. Clin. Microbiol. 22: 778 – 781.
- Quinn, P.J.; Carter, M.E.; Markey, B.K. and Carter, G.R. (1994): Clinical Veterianry Microbiology. Published by welfe Publishing.

- Rock, J.L. and Nelson, D.R. (2006): Identification and characterization of a hemolysin gene cluster in Vibrio anguillarum. Infection and immunity 74(5): 2777-2786.
- Schaperclaus, W.; Kulow, H. and Schreckenbach, K. (1992): Fish Diseases. Vol. 1, 5th corrected, revised and substantially enlarged edition. A.A. Balkema. Roterdam.
- Shaaban, A.I.; Easa El-S.M. and Diab, S.A. (1995): Characterization of V. anguillarum isolated from wild fish eels (Anguilla japanica) in Egypt. J. Egypt. Vet. Med. Ass., 55 (1, 2): 141-145.
- Stephens, F.J.; Raidal, S.R.; Buller, and Jones, B. (2006): Infection with Photobacterium damselae subspecies damsela and Vibrio harveyi in snapper, Pagrus auratus with bloat. Australian – Vet. J. 84 (5): 173 – 177.
- Yonis A.A.; Amer, M.M.; Azizi Naguib M. and Abdel–Azizi E.S. (1997): Pharmacokinetics, efficacy and tissue residues of enrofloxacin in Oreochromis niloticus infected with Vibrio anguillarum. Vet. Med. J. Giza 45(1). 75 – 85.
- Zeinab, I. soliman (1999): Antibiogram of some bacteria contaminating Tilapia fish at El- Manzala lake in Port–Said Governorate. Vet. Med. J., Giza. 47 (1): 19 – 27