Histological changes in skin and gill of fresh water EUS infected fish *Channa punctatus*

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Abstract

Histopathology is an important diseases diagnostic tool. Penetration into the host is the first step for a microbial agent to multiply and invade the vital organs of its host. Like all animals fishes suffers from environmental, stress, which is followed by pathogenic attacks and parasitic afflictions causing Epizootic Ulcerative syndrome (EUS). Histological investigations of tissues of EUS infected fish *Channa punctatus* showed an eroded epidermis, necrosis of skeletal muscles and intense granulomatous inflammation, infiltration of mixed inflammatory cells in muscular as well as submuscular layers of the skin. Also a characteristic focal granular granuloma in the gill lamellae was noticed. The electron microscopic observations, of the tissue cells of the fishes showed viruses like Parvovirus even at a red spot stage. Thus, it can be said that Viruses being crystalline in nature have cellular penetrating power & the EUS infection follows the phylogenetic evolution in life in true sense i.e. RNA virus followed by Bacterial cell and then Fungus. Thus, the presence of EUS in *Channa punctatus* is confirmed histologically.

Key words: *Channa*, Epizootic Ulcerative syndrome, Fungus, Histology, Parvovirus.

Introduction

A body of water with low pH has been documented in other outbreaks as a significant predisposing factor for EUS outbreaks (Sammut *et al.*, 1996; Baldock *et al.*, 2005; Choongo *et al.*, 2009). Release of pollutants in the nature are affecting the basic environmental parameters like temperature, salinity, DO and pH affecting the protective mucus layer of the fishes making them vulnerable to the attack of virus followed by bacterial and fungal infection. Histopathological changes would be expected in organ, when the gross clinical and pathological signs of this type of disease occur (Vogelbein *et al.*, 2001). Bacterial colonies observed in tissues are a proof of the presence of bacterial (Lio-Po *et al.*, 1998), fungal (Lio-Po *et al.*, 1998) and viral (Kanchankhan, 1996) infections. Kane *et al.*, (2000) carried out the etiologies, observations and reporting of eustuarine fish fin lesions and histologically observed that the lesion demonstrated a marked chronic inflammatory infiltrate and granulomas in response to fungal hyphae throughout large areas of necrotic muscle tissue and also observed Gram-negative rod-shaped bacteria in the lesions, a common finding in ulcers of aquatic organisms. Fish showing sores and ulcers referable to EUS were reported in the Equateur Province of the Democratic Republic of the Congo in December 2014, with a record of heavy fish mortalities. The present investigation is aimed to the identification of EUS through, histopathological observations of fresh water fish *Channa punctatus*. The critical gross pathology was represented by cutaneous lesions fitting the case definition of EUS – red spots, erosions, ulcers and wounds. Skin ablation is thought to facilitate attraction of and infection by zoospores of *A. invadans* (Songe *et al.*, 2012). The presence of fungal hyphae was demonstrated in the epidermis of some early, stages of infected fish from India (Viswanath *et al.*, 1997). Kanchanakhan *et al.*, (1998) achieved virus isolations from 3 epizootics in Thailand in 1993-1994 epizootic, 9 rhabdoviruses from EUS snakehead (*Channa striata*), three-spot gourami (*Trichogaster trichopterus*). And another 9 rhabdoviruses from 11 tissue extracts from affected snakehead collected in the 1995-1996 outbreak.

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Results of their study suggested that the viruses, especially rhabdovirus, are readily isolated from diseased fishes during the early period of an outbreak, and that they may have a significant role in a complex of etiological agents of EUS. Histopathological changes might be the concurrent effect of both the fungal pathogen (Aphanomyces) and the bacteria A. hydrophila. EUS diseases of many fishes were found to be caused by both the above pathogens by many authors in different fishes (Hasan et al 2008).

**Material and Methods**

The present study was carried out by collecting the suspected diseased fish from near by EUS prone water bodies (Wadali and Malkhed lakes) with the help of fishermen and were brought to the laboratory and maintained in glass aquaria for a week to study the type of infection. The healthy fish weighing about 25 gm and of approximately equal length were brought to the laboratory and kept in aquarium containing 75 litre of water and were acclimatized for 15 days. They were regularly fed on pellet feed and 25% water of the aquarium was also regularly changed at an interval of 7 days to maintain the water quality. 10 fishes were kept in one aquarium each.

**Experimental set up:**

The experiment was carried out on six sets of 10 fishes in each group as below:

- **Group I:** Consisting of control (Healthy) fish (10, *Channa punctatus*).
- **Group II:** Consisting of Naturally EUS affected fish (10, *Channa punctatus*).
- **Group III:** Consisting of artificially infected fishes injected with 0.2 ml homogenate of ulcerated skin tissue from diseased *Channa punctatus* (10, *Channa punctatus*).

**Suspension preparation:**

The tissue suspension of ulcerated skin tissue was prepared in sterile fish saline (0.3%). A tissue (500 mg) homogenate was prepared by taking affected or ulcerated skin in fish saline (5ml).

**Histological studies:**

The tissues like liver, gill, skin and kidney of EUS affected fishes (*Channa punctatus*) were dissected out, washed with saline water to remove adhering patricals and blood stains and then fixed in Bouin’s fixative for 16 to 20 hrs. Then the tissues were washed thoroughly with water, dehydrated with graded series of alcohol and embedded in paraffin wax and sections were cut at 4 to 5 microns. The sections were processed and stained with Haematoxylin-Eosin by standard methods as described by Weissman (1972).

**Electron microscopical studies for detection of viral infection:**

**Specimen Preparation for TEM:**

The tissues liver, kidney, skin and gills were dissected out aseptically in the laboratory, from the EUS affected fishes, trimmed to size 1.0-1.5 mm thick and fixed into a fixative containing 2.5% Glutaraldehyde and 2% paraformaldehyde, made in 0.1 M sodium phosphate buffer (pH 7.4) for 6-12 hrs at 4°C for 24 hrs. After fixation, the specimens were sent for detection of Viral infections, at AIIMS, New Delhi, where the ultra-thin sections were visualized under transmission electron microscope at different magnifications and photographed.

**Results and Discussion**

A general survey of Amravati fish market was carried, which revealed that most of the EUS affected fishes were brought by the fisherman from Wadali and Malkhed lakes. So, they were procurred from Wadali and Malkhed lakes during September to January months of 2004 to 2006.

All EUS affected fishes exhibited three distinct stages during the EUS infection (Table 1).

- **Stage - I:** Red spot is seen
- **Stage - II:** Red ring with a white patch in the centre
- **Stage - III:** The area of red ring became necrosed with puffy appearance with ulcer and haemorrhage. Multiple ulcers were seen on the body in advanced stage.

The healthy fishes (Fig: 1) were administered with the ulcerated tissue from the naturally EUS affected fish (Fig: 2) and were studied after 7, 14 and 21 days of challenge infection. These fishes also showed the three stages of the disease. In artificially EUS infected fishes a red spot was developed after 7 days of challenge inoculation and later on after 12-14 days a red ring was seen with
loss of scales. Then after 20-24 days, the tissue exposed showing puffy appearance (Fig: 3 to 5). within the ring was totally degraded with muscles

Table: 1. External features shown by naturally EUS affected and artificially EUS infected fishes (*Channa punctatus*)

<table>
<thead>
<tr>
<th>Stage I</th>
<th>Stage II</th>
<th>Stage III</th>
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<tr>
<td>Red spots on the body surface were seen. No discolouration as such on the skin surface or on the skin around the red spot was seen.</td>
<td>The Red spots turned into lesions which were around 1 cm in diameter, they appeared to be raised and circular in shape and the body surface area showed discolouration. At this stage the skin was intact and the scales were loosely attached.</td>
<td>The lesions formed showed the advanced stage of necrosis. They appeared to be circular extending deep into the skeletal musculature. Large haemorrhagic and necrotic ulcers were formed along with new emerging red spots seen on various parts of the body. At this stage the epidermis and scales were completely lost along with loss of dermis at the site of ulcer.</td>
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Fig :1, Control healthy *Channa punctatus*

Fig :2, Naturally EUS affected *Channa punctatus.*
Histopathological changes in skin:
Non-infected healthy control fishes exhibited an intact epidermis comprising stratified squamous epithelium 10-15 cells thick (Fig: 6). In contrast, infected *Channa* consistently exhibited widespread partial to complete loss of the epidermal layer of the skin (Fig: 7). After 21 days of infection, the fishes showed eroded epidermis over much of the infected regions (Fig: 8). Histologically the lesions are characterized by the presence of highly invasive, aseptate fungal hyphae penetrating deeply into the body muscles. Necrosis of skeletal muscles.
and intense granulomatous inflammation directed at the fungus were the prominent features of lesions seen in the V.S. of skin (Fig: 8).

**Histopathological changes in liver:**
The hepatic parenchyma is made of hepatocytes spread out as anastomotic cords arranged in two cellular layers and surrounded by sinusoids (Fig: 9). The bile ducts are usually found near the portal vein and they are lined by simple cuboidal epithelium. In EUS infected fishes, focal necrosis was seen in the liver with vacuolation of the hepatocytes; congestion and haemorrhages (Fig:10). The hepatic cells appeared shrinked with fungal infiltration (Fig: 11).

**Histopathological changes in gills:**
In Comparison to the gill lamellae of healthy *Channa punctatus* (Fig:12), The gill lamellae of the EUS infected Channa punctatus showed a well developed encapsulated granulomas around necrotic area is observed, which appeared nodular with characteristic focal (tubercles) granulomas. These are composed of a diffuse distribution of reticuloendothelial cells and macrophages with large caseous necrotic areas (Fig:13 and 14).

**Histopathological changes in kidney:**
In Comparison to the normal structure of T.S. kidney of healthy *Channa punctatus* (Fig: 15). The kidney of EUS infected fishes exhibited tubular necrosis, interstitial lymphocyte infiltration, haemorrhage (Fig:16). In EUS affected channa, shrinkage of the glomeruli with a large space between the Bowman’s capsule and the glomerulus was also seen (Fig:17).

**Electron Microscopic alterations in EUS affected fishes:** Electron Microscopic study was carried out in naturally EUS affected fishes, showing heavy Parvovirus (DNA virus) infection at an area of red spot in *Channa punctatus* (Fig :18)
The EM structure of hepatic cell of naturally affected *Channa punctatus* showed damage and breakdown of mitochondria and distortion of endoplasmic reticulum (ER), (Fig:19). Chloride cell of gill of EUS affected *Channa punctatus* showed infection of Paroviruses (DNA virus) distorted nucleus and disintegration of mitochondria (Fig: 20).
Fig: 10. T.S. of liver of healthy *Channa punctatus* showing focal necrosis and distention of hepatocytes.
FN - Focal necrosis, DH - Distention of hepatocytes.

Fig: 11. T.S. of liver of EUS affected *Channa punctatus* showing shrunked hepatocytes with fungal infiltration.
SH - Shrunked hepatocytes, FI - Fungal infiltration.

Fig: 12. Gill lamellae of healthy *Channa punctatus*

Fig: 13. Tubercle granuloma seen in gills of EUS infected *Channa punctatus*.
TG - Tubercle granuloma

Fig: 14. Tubercle granuloma (TG) in gills of EUS infected *Channa punctatus* (Magnified).

Fig: 15. T.S. of kidney of healthy *Channa punctatus* showing normal structure.
G - Glomerulus
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Fig: 16. T.S. of kidney of EUS affected Channa punctatus showing hydropic degeneration, and damage to tubular epithelium. HD-Hydropic degeneration.

Fig: 17. T.S. of kidney of EUS affected Channa punctatus showing lymphocyte infiltration, granuloma, increased number of inter renal cells and necrotic tubular epithelial cells. G-Granuloma L-Lymphocyte infiltration NEC-Necrotic tubular epithelial cells.

Fig: 18. EM showing heavy viral infection in the skin of Channa punctatus at an area near red spot showing Paroviruses (DNA virus). PV-Parovirus.

Fig: 19. Electron Microscopic image of a hepatic cell of EUS affected Channa punctatus showing mitochondrial damage and swelling and distortion of endoplasmic reticulum. SDR-Swelled and distorted endoplasmic reticulum, MD-Mitochondrial damage.

Fig: 20. EM showing heavy viral infection in the chloride cell of gill of EUS affected Channa punctatus showing Paroviruses (DNA virus), distorted nucleus, and disintegration of mitochondria. DN-Distorted nucleus, DM-Disintegration of mitochondria, PV-Paroviruses.
Ulcers are characterized histomorphologically by the consistent presence of deeply penetrating fungal hyphae, extensive myodegeneration and a special type of host cellular defensive response called granulomatous inflammation directed by the fungal agent (Dykstra et al., 1989; Blazer et al., 1999; Vogelbein et al., 2001). This type of response is typically elicited in vertebrates by infections that are difficult for a host to kill and degrade and is composed predominantly of a mononuclear phagocyte called the macrophage (Cheville, 1983). These cells accumulate within the infected tissues and organize into discrete structures called granulomas that effectively encapsulate the foreign invader. This special type of inflammatory response developed over an attempt by the host to bring to bear a suite of toxic and cytolytic substances produced by the macrophages and designed to inactivate, kill and degrade the infectious agent. Histopathological findings of the present study strongly agree with the definition and diagnostic features of EUS as proposed in the ODA regional seminar on EUS (Roberts et al., 1994). Histopathological results clearly and consistently show that a specific type of broad, non-septate fungus involved in all EUS affected fish species. Chronic inflammatory tissue surrounding the fungal hyphae to form massive epitheloid granulomas was particularly evident in either gills (Fig:13 and 14) or skin (Fig: 8) of the fish specimens studied.

**Conclusion**

Histological studies of the skin of EUS infected channa showed initial epidermal loss exposing the dermis, degeneration of muscles with infiltration of mixed inflammatory cells with deep penetrating of fungal hyphae into the body muscles. As well as in the liver cells. In advanced stage of the disease, the hepatic cells appeared shrunk with fungal infiltration. In EUS infected *Channa punctatus*, a well developed encapsulated granuloma in the gill lamellae was developed which appeared nodular with characteristic focal granular granulomas. These were composed of a diffuse distribution of reticular endothelial cells and macrophages. The kidney of EUS affected fishes exhibited tubular necrosis, interstitial lymphocyte infiltration, haemorrhage and well developed inter-renal tissue. Glomeruli appeared shrunk.

In all the electron microscopic observations, the tissue cells of EUS affected fishes showed viruses like *Parvovirus* even at a red spot stage. Distortion of endoplasmic reticulum and breakdown of mitochondrial cristae were the common observations. In EUS infected *Channa punctatus*, a well developed encapsulated granuloma in the gill lamellae and presence of viruses like *Parvovirus* even at a red spot stage of EUS affected fishes was seen, thus, from the histological observations, it can be said that release of pollutants in the nature are affecting the basic environmental parameters like temperature, salinity, DO and pH affecting the protective mucus layer of the fishes making them vulnerable to the attack of virus being crystalline in nature and have cellular penetrating power, followed by bacterial and fungal infection causing more damage to fish tissue leading to ulcers (EUS) ultimately death of the fish (Economic loss) a Loss to aquatic biodiversity, irreversible ecological damage.

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**References**


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The probable path of EUS

Post rainy season

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Water parameters fluctuate

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Stressed fish

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VIRUS

Immunosuppression

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Loss of mucus and scales

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Red spot appear on the body at the point of infection

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Scale loss, degeneration of epidermis with exposure of dermis, inflammation and appearance of ring

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Bacteria and fungi invade deeper in fish tissue

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Tissue necrosis, Muscle necrosis, infiltration of Lymphocytes, Granulomatous growth, Antigen - Antibody reactions, Proliferation of lymphocytes leads to more damage to fish tissue leading to ulcers (EUS)

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Death of the fish (Economic loss)

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Loss to aquatic biodiversity

irreversible ecological damage