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Junul FOR RESEARCE	Original Research Paper	Environmental Science
Truena informational	HISTOPATHOLOGICAL CHANGES IN THE GILL OF FRESHWATER TELEOST HETEROPNEUSTES FOSSILIS (BLOCH) EXPOSED TO METANIL YELLOW	
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ABSTRACT The chronic toxic effect of Metanil Yellow on the gills of freshwater teleost Heteropneustes fossilis (Bloch) was studied. The fishes were exposed for 45 days treatment in 2a/L of Metanil Yellow. However, after 45		

days treatment, marked pathological changes in the gills were found. Necrosis found in the primary gill lamellae. Shortened and clubbing ends of the secondary gill lamellae. Buldging tip of primary gill filaments and distortion was occurred in the shape of secondary filament. Pycnotic nuclei, vacuolization and degenerative epithelial cells were found.

KEYWORDS : Histopathology, Gill, Metanil Yellow, Heteropneustes fossilis (Bloch)

INTRODUCTION

A non-permitted synthetic food colour Metanil Yellow is principally the monosodium salt of 3-[[4-(Phenyl amino) phenyl] azo] benzene sulfonic acid which is used in different food items namely biriyani, ghugni, different sweet items viz., laddu, jalebi, bundia, amriti etc. The acute and short term toxicity of popular blend of Metanil Yellow and Orange –II in albino rats was recorded by Singh and Khanna (1998). Toxicity of Metanil Yellow and Derma Orange to a fresh water fish Channa punctatus was reported by Goel and Basu (1989). Gupta et al., (2002) reported the tumor promotion by Metanil Yellow and Malachite Green during rat hepatocarcinogenesis. Many authors (Mehrotra et al., 1974; Khanna and Das 1991; Gupta et al., 2002) have investigated the toxic effects of Metanil Yellow. In fish, gills play vital functions like respiration, osmotic regulation and excretion. Gills act as tissue indicator of water quality and also for the assessment of environmental impact. In the present work, an attempt was made to evaluate the chronic toxic effects of Metanil Yellow on the histology of gill tissues of the freshwater teleostean catfish Heteropneustes fossilis (Bloch).

MATERIALS AND METHODS

Teleostean fish, Heteropneustes fossilis (Bloch) of 23.2 \pm 1.5 cm with an average weight of 64.3 \pm 2.0 g were collected from local pond of East Barddhaman, W.B. and after collection the fishes were acclimatized in glass aquaria of dechlorinated water and fed with Tubifex sp. twice daily for two weeks. Water temperature and pH were maintained during experimentation. After acclimatization two sets carrying ten fish specimens in each glass aquaria as control and treated were arranged. One set of the test fishes were fed with Metanil Yellow at a dose of 2g/L for 45 days. On the $46^{\mbox{\tiny th}}$ day of the experiment the desired gill tissues were collected from both control and treated specimens and were subjected to histopathological technique. For this the required tissues were fixed in aqueous Bouin's fluid solution. After decalcification of gills, the paraffin sections of 4-5 micron were cut and stained in Delafield's Haematoxylin and Eosin solution.

RESULTS AND DISCUSSION

In fish gills are the main respiratory organs and they provide maximum oxygen for intracellular oxidation. The gills are situated in the pharynx. Each teleostean gill consists of a longer lower limb supported by ceretobranchial, and a shorter upper limb supported by the epibranchial. Each gill arch bears gill rakers towards the inner (buccal) side and gill filaments towards the outer (opercular) side. Each gill arch bears two hemibranchs, which are formed by endodermal evaginations in the form of leaf like folds, called gill filaments or primary gill lamellae. The primary gill lamellae (PGL) of

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Heteropneustes fossilis consist of a central rod like supporting axis and a row of secondary gill lamellae (SGL) also termed as respiratory lamellae are present on each side of the primary gill lamellae (PGL) whereas the secondary lamellae (SGL) are highly vascularised and are covered with a thin layer of squamous epithelial cells (Fig. 1). Single nucleus is found in each blood cell of secondary lamellae.



Fig.1: Photomicrographs of the gill tissue of control (C) *Heteropneustes fossilis* [H-E] which showing normal aspect of primary filament (PF), primary gill lamellae (PGL), secondary lamellae (SGL) and mucous cell (MC) (normal arrows). (C) [10 X 20]



Fig. 2: Photomicrographs of the gill tissue of Metanil Yellow (MY) treated *Heteropneustes fossilis* [H-E] which showing shortened and clubbing ends of the secondary gill lamellae (SGL) and damaged mucous cell (MC) (broken arrows). (MY, 45d) [10 X 40]

Fig. 3: Photomicrographs of the gill tissue of Metanil Yellow (MY) treated *Heteropneustes fossilis* [H-E] which showing distortion and damages in the primary gill filament (PF), secondary gill lamellae (SGL) and mucous cell (MC) (broken

arrows). (MY, 45d) [10 X 40]

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After an exposure of 45 days treatment with Metanil Yellow huge changes have been found in the gill tissues. Gill rakers and filaments were distorted and damaged. Buldging tips were found in primary gill filaments (PF) and distortion occurred in secondary filament (Fig. 2). Necrosis was shown in the primary gill lamellae (PGL). Shortened and clubbed ends of secondary gill lamellae (SGL) and hyperplasia and hypertrophy of nuclei were found after toxicosis of Metanil Yellow (Fig. 3).

Histopathological study has revealed that Malachite Green causes detrimental effects in liver, gill, kidney, intestine, gonads and pituitary gonadotropic cells. It has been found to causes sinusoidal congestion and focal necrosis in liver, damages mitochondria and also causes nuclear alterations (Gerundo et al., 1991). Hypertrophy and vacuolisation followed by necrosis and cirrhosis have been observed in hepatocytes of *Heteropneustes fossilis* following treatment with Malachite Green (Srivastava et al., 1998a). Exposure to this dye has also found to cause severe damage to gills, resulting in necrosis of lamellar cells and gill epithelium, and leucocyte infiltration in rainbow trout (Gerundo et al., 1991) and *Heteropneustes fossilis* (Srivastava et al., 1998b).

Gill lesions in zinc treated Heteropneustes fossilis was reported by Hemalatha and Banerjee (1977). So many researchers (Daoust et al., 1984; Karlson-Norgren et al., 1985; Tophon et al ., 2003; Gupta and Kumar 2006; Al-Attar, 2007; Kaoud and El-Dahshan, 2010; Patel and Bahadur, 2010; Khoshnood et al., 2011; Patnaik et al., 2011) observed the pathological lesions in fish gills exposed to heavy metals. Changes in the primary and secondary gill lamellae of Heteropneustes fossilis were observed after an exposure of Malathion for 15 days (Yadav et al., 2018). The chronic toxicity of non-permitted food colour Metanil Yellow (MY) for an exposure of 45 days to teleostean catfish Heteropneustes fossilis has shown histopathological changes in stomach, intestine, liver and kidney of H. fossilis (Sarkar and Ghosh, 2010). Degenerations in the seminiferous tubules and spermatocytes and vacuolations in the sertoli cells of albino rat (Rattus norvegicus) were shown when exposed to Metanil Yellow for 30 and 45 days (Sarkar and Ghosh, 2012a). Histopathological and ultrstructural changes in the stomach, intestine, liver and kidney of albino rat (Rattus norvegicus) were observed (Sarkar and Ghosh, 2012). Sarkar and Ghosh (2017) reported the histopathological lesions in spleen of albino rat (Rattus norvegicus) were found after fed with Metanil Yellow for 30 and 45 days. But maximum distortion and degeneration occurred in both the white pulp and red pulp regions after 45 days treatment. The present observations in Heteropneustes fossilis clearly demonstrate the gill tissues affected most severely in response to Metanil Yellow intoxication.

CONCLUSION

The damages shown in gill tissues vary not only with its concentration but with the time of exposure period. During the tenure of 45 days treatment with synthetic food colour Metanil Yellow, maximum damages were shown in primary and secondary gill lamellae.

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