



Ultrastructural Data on the Spore of *Myxobolus dermatobius* Ishii, 1915 (*Myxosporaea: Myxobolidae*) Infecting Eye of Nile-Tilapia (*Oreochromis niloticus*) in Egypt

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Abstract: A total of 1000 cultured Nile-tilapia (*Oreochromis niloticus*) were collected from different governmental and private fish farms and examined for detection of myxosporean parasites infection. The infected fish showed slight unilateral exophthalmia with whitish cyst in the eye. Numerous white cyst like plasmodia of *Myxobolus dermatobius* were recovered from the eye of the examined fish with low prevalence rate (1%). Small intact cyst was isolated, fixed in 3% glutaraldehyde in 0.1 M sodium cacodylate (pH 7.4) and prepared for electron microscopy examination. Ultrathin sections of *Myxobolus dermatobius* spore revealed pair of capsulogenic cells at the apical pole of the developing spore.

INTRODUCTION

Fishes consider as one of the most important sources of animal protein all over the world. Egypt is a country with numerous lakes, seas and long river having the most diversified fauna of fresh and marine water fishes. Fish eye is considered a very important organ, it is adapted for the vision in air as well as waters either in wild and cultured fish which do display some differences to the mammalian eyes^[1]. The awareness of and experience with parasites that affect fish health, growth and survival are increased with increasing the interest in fish culture and production. Therefore, the contribution to the knowledge of fish parasites is a prerequisite for early and correct diagnosis of the disease agent that can lead to preventive measure which is the best way to reduce outbreaks of disease^[2,3].

Myxosporidea are frequently described in fresh water, brackish and marine fishes and have a great importance in ichthyopathology. Myxosporean parasites

are the most important fish pathogens and more than 2,300 species have been reported from marine and fresh water fish in several geographical areas^[4-7]. Myxosporaea infecting Egyptian fish is a group of parasites which is considered as the major cause of myxosporidiosis and harm of the fish^[8-11]. *Myxobolus Butschli*, 1882, is one of the largest genus of myxosporean groups. Landsberg and Lom^[12] gave a list of 444 *Myxobolus* species followed by Eiras *et al.*^[13, 14] who listed approximately 744 and 856 species of *Myxobolus* parasitizing fish from all over the world respectively. The species of *Myxobolus* infect a diverse set of specific tissues that can include specifically the tegument, eyes, gills, glands, gonads, kidneys, muscle, digestive tract and nervous system^[15, 6]. *Myxobolus dermatobia* recovered from *Oreochromis niloticus* causes petechial to focal haemorrhages in orbit, exophthalmia and unilateral eye opacity, especially in advanced cases^[16] while *Myxobolus dematobia* isolated from *Tilapia zilli* at Giza province causes unilateral eye opacity^[17].

The ultrastructural morphology of myxosporean species have been widely studied^[12, 15], however, few species have been ultrastructurally described in Egypt. So, the present paper give ultrastructural data of *Myxobolus dermatobius*, infecting Nile tilapia, *Oreochromis niloticus* by using transmission electron microscope.

MATERIALS AND METHODS

A thousand of live or recently dead Nile-tilapia; *Oreochromis niloticus* were collected from different governmental and private fish farms in Sharkia Governorate, Egypt. The collected fish were transported to the laboratory and dissected. The different organs were examined macroscopically and microscopically for detection of any visible myxosporean cysts.

Small intact cysts with minimum surrounding tissue were isolated and fixed in 3% glutaraldehyde in 0.1 M sodium cacodylate (pH 7.4) for at least 24 h, washed in the same buffer and post-fixed with 2% OSO_4 in the same buffer. The specimens were dehydrated in series of graded ethanol, transferred to propylene oxide and finally, embedded in araldite. Ultrathin sections were stained with uranyle acetate and lead citrate and examined with a Philips (208) electron microscope at 80-100 kV^[18].

RESULTS AND DISCUSSION

The prevalence of infection with *Myxobolus dermatobius* among the examined fish was 1% (10 out of

1000). Several species of *Myxobolus* were isolated from *Oreochromis niloticus* in Egypt; *Myxobolus nilei* from eyes, skin, gills, kidney, spleen and pancreas^[19], *M. heterosporus* from eye, muscle and kidney^[20], *M. spheroidalis* and *M. ocularis* from eye^[21], *M. heterosporus* from eye and gills^[22], *Myxobolus* sp. from the inner wall of cornea, the base of the gill arch and roof of the mouth^[23], *M. cornealis* from the eye^[24], *M. dermatobia*^[16] and *M. heterosporus*^[25] from eye and cornea. *Myxobolus dermatobia* was isolated from eye of *Tilapia zilli* at Giza province^[17]. The parasite was found in the form of whitish cyst in the eye of tilapia causing slight unilateral slight exophthalmia. The spores of *Myxobolus dermatobius* (Syn. *Myxobolus dermatobia*) were recovered from their original plasmodia found in the infected eye of Nile tilapia, *Oreochromis niloticus* (Fig. 1). Similar lesion of exophthalmia was noticed by Abdel-Aal^[16] and Mohamed *et al.*^[17].

Most of the Egyptian *Myxobolus* species have been described based on light microscopy descriptions and diagrammatic drawings even fewer have been described using ultrastructural observations. The ultrastructural characteristics of *Myxobolus dermatobius* revealed that each spore develops from five cells; a pair of capsulogenic cells, two peripherally arranged valvogenic cells and one sporoplasm cell. Capsulogenic cells are found at the apical pole of the developing spore and together with the sporoplasm, forms a central core that is ensheathed by valvogenic cells. These cells give rise to the two shell valves surrounding each spore and the sutural ridge joining the valves. The differentiation of the capsulogenic cells starts with appearance of a club-shaped structure “capsular primordium” (Fig. 2-4).



Fig. 1: Eye of Nile-tilapia *Oreochromis niloticus*; showing white plasmodia (arrows) of *Myxobolus dermatobius* infection

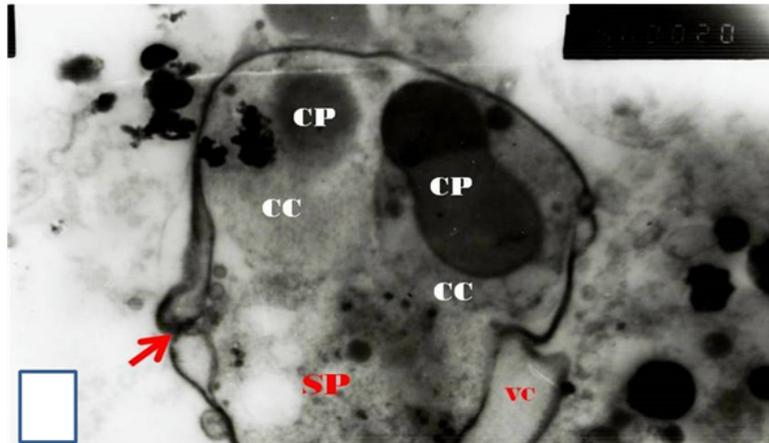


Fig. 2: *Myxobolus dermatobius* premature spore showing two Capsulogenic Cells (CC) containing Capsular Primordium (CP); Sporoplasm (Sp); Valvogenic Cell (VC) and suture valve (arrow). X5000

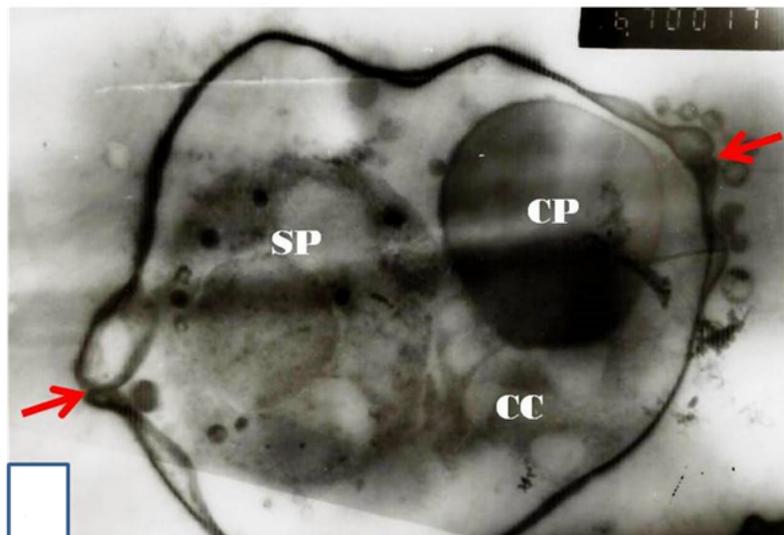


Fig. 3: *Myxobolus dermatobius* premature spore showing two Capsulogenic Cells (CC) containing Capsular Primordium (CP); Sporoplasm (Sp). X5000

This come in accordance to *M. dermatobia* described by Abdel-Aal^[16] and *Myxobolus* sp.^[26]. The capsulogenic cell of the present species showed capsular primordial as described in *M. stomum* and *Myxobolus* sp. by Ali *et al.*^[27] and Abdel-Ghaffar *et al.*^[26], respectively. The valvogenic cells gave rise to shell valve surrounding each spore and sutural ridge joining the valves were similar to *M. dermatobia* described by Abdel-Aal^[16].

Sporoplasm fills nearly all the space beneath the polar capsules. It contains single mono-nucleated, small vesicles and sometimes exhibited dense matrices known as sporoplasmosomes. A small area of sporoplasm is

occupied by a glycogen body (Fig. 4-5). Corresponding finding of nucleus was observed in *Myxobolus dermatobia* described by Abdel-Aal^[16] while binucleated sporoplasm was described in other species; *Myxobolus stomum*^[27]; *Myxobolus braziliensis*^[28] and *Myxobolus* sp.^[29,30]. The sporoplasmosomes of the present species complied with a similar dense body found in *Myxobolus cotti* reported by EI-Matbouli *et al.*^[31]; *M. dermatobia*^[16]; *M. stomum*^[27] and *Myxobolus* sp.^[26]. The glycogen body noticed in the sporoplasm is essential in the myxosporean spore which could provide the energy necessary for further developmental stages in the life

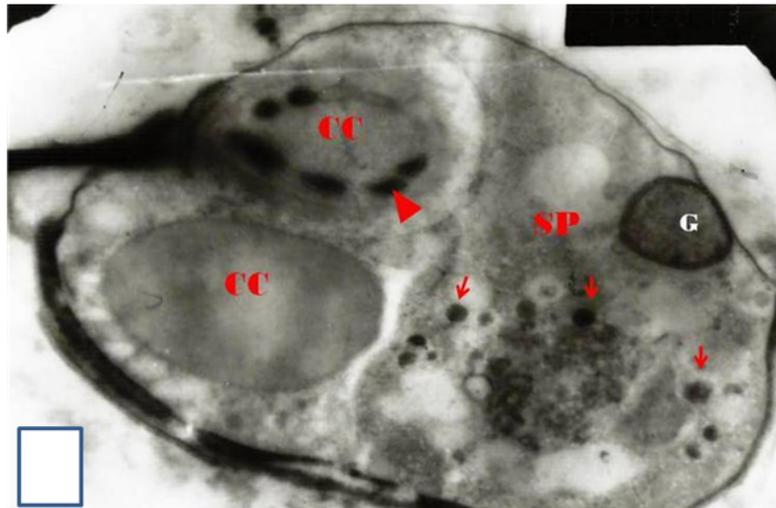


Fig. 4: *Myxobolus dermatobius* nearly mature spore containing two Capsulogenic Cells (CC), primordia of polar filaments (arrow head); Sporoplasm (Sp) with sporoplasmosomes (arrow) and Glycogen body (G). x5000

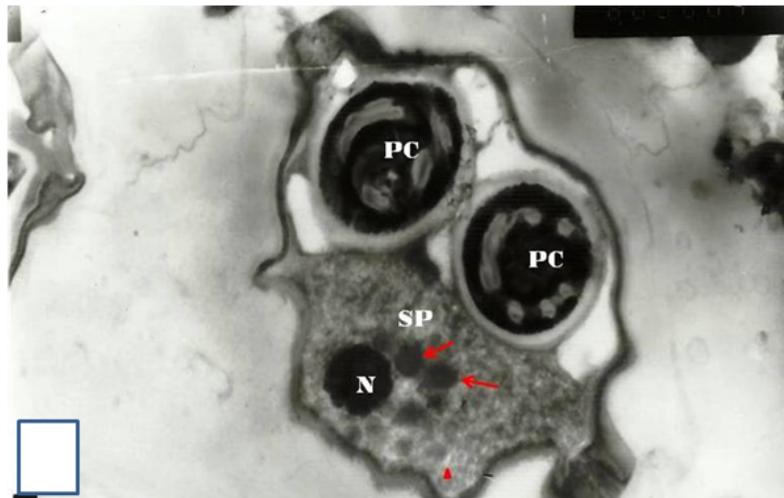


Fig. 5: Transverse section through *Myxobolus dermatobius* mature spore, showing Polar Capsules (PC); Sporoplasm (Sp) containing sporoplasmosomes (arrows); Nucleus (N) and small vesicles (arrow head). X5000

cycle. It was similar to that reported in *M. cotti*^[31]; *Myxobolus* sp. (Abdel-Ghaffar *et al.*, 1994) and *M. dermatobia*^[16].

Two polar capsules are pyriform in shape, equal size, located side by side at the same level and occupy approximately half of the total spore length (Fig. 6). Each polar capsule has a homogenous core of medium electron-density containing polar filaments, surrounded by an electron-lucent layer and an outer layer of medium density (Fig. 7). Similar findings were reported in many species of *Myxobolus*; *M. cotti*^[31], *M. dermatobia*^[16], *M. stomum*^[27], *Myxobolus* sp.^[26] and *Myxobolus sciades*^[32]. The number of polar filament coils is probably

4 turns in each capsule. The apical portion of each polar capsule has cap-like cover plugged the apex of each mature polar capsule (Fig. 8). These was identical to that of *Myxobolus dermatobia* described by Abdel-Aal^[16]. The same number of polar filament coils was reported in *M. heterosporus* by El Mansy^[25] while different numbers of polar filament turns were mentioned by Casal *et al.*^[28], Ali *et al.*^[27], Abdel-Ghaffar *et al.*^[26], Azevedo *et al.*^[32], Kaur and Singh^[33] and Abdel-Baki *et al.*^[34] in *M. maculatus* (14-15) *M. stomum* (5-6) *Myxobolus* sp. (5); *M. sclerii* (4-5), *M. sciades* (9-10) and *M. brachysporus* (6-7), respectively.

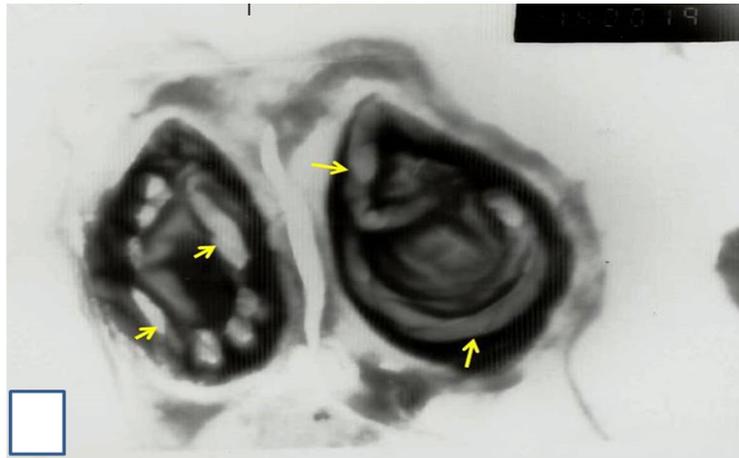


Fig. 6: Longitudinal section through anterior portion *Myxobolus dermatobius* mature spore showing two synchronously developed Polar Capsules (PC) with polar filament coils (arrows) x5000

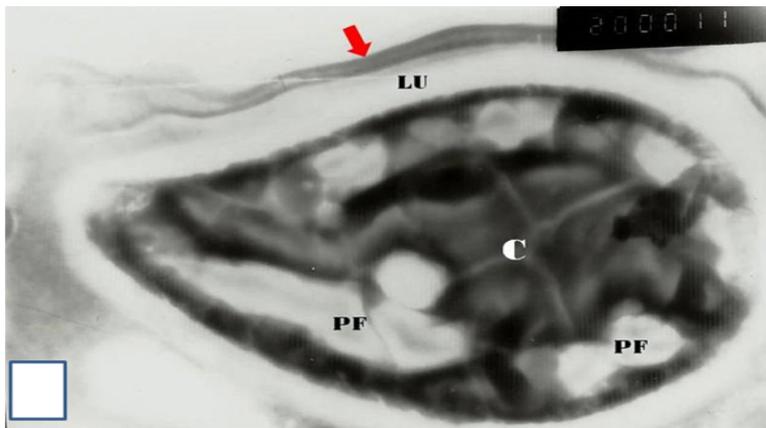


Fig. 7: Longitudinal section through *Myxobolus dermatobius* well developed polar capsule showing an electron-dense outer layer (arrow); a central translucent layer (LU) and inner dense core (C) with polar filament coils (PF). X7600

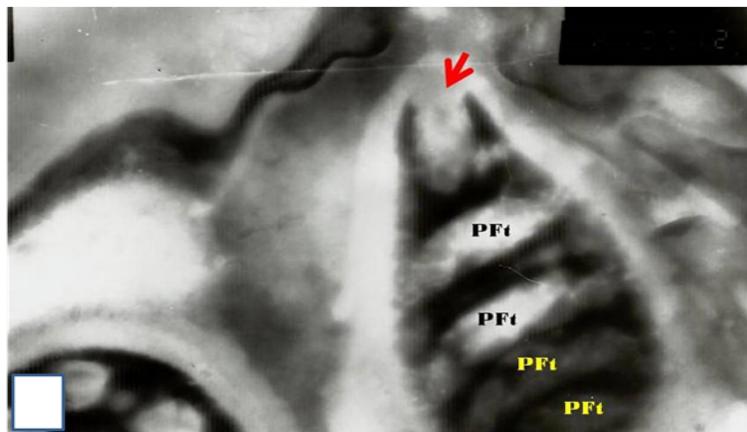


Fig. 8: Longitudinal section through *Myxobolus dermatobius* well developed polar capsule showing four turn of Polar Filament (PFt) and apical cap (arrow). X14000

CONCLUSION

Single sporoplasm containing a nucleus and sporoplasmosomes fills nearly all the space beneath the polar capsules. The later were pyriform in shape, each one had homogenous dense core and 4 turns of polar filaments. Ultrastructural characteristics of the present spore were described and discussed in detail.

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