STUDIES ON OCULAR SARCOCYSTIS IN BUFFALOES IN ASSIUT GOVERNORATE

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ABSTRACT:

A total of 150 buffalo’s ocular muscle samples were examined for the presence of sarcosporidians in Assiut Governorate. These samples included 80 males less than 2 years old and 70 females more than 5 years old. Sarcocystis spp. were detected in 55 (36.66%) of examined animals. The incidence of infection in males was 15.0% and in females was 61.42%. Two species of Sarcocystis were detected in the present work: S. fusiformis was detected in 9.33% of examined animals, where in males it was 1.25% and in females it was 18.57%. S. levinei was detected in 34.66% of examined animals, in males it was 15.0% and in females was 57.14%. Mixed infection with the two species was detected in 11 (7.33%) of examined animals. The morphological characters and the ultrastructure of each species were described. The present data showed that ocular muscles are a preferable site for Sarcocystis especially S. levinei. It is, therefore recommended that Sarcocystis cysts showed be looked for in ocular muscles of humans complaining of ocular muscle disease.

INTRODUCTION:

Sarcosporidia are protozoal parasites found in a wide variety of animals and man, where more than 50 species are detected in mammals and birds (Gracey and Collins, 1992). Sarcocystis has an obligatory prey/predator (2 hosts) life cycle, asexual stages develop in intermediate host (predator) which is herbivorous and sexual stages develop in definitive host (prey) which is carnivorous, predator may act as a prey at the same time (Levine, 1986). Sarcocystis is found mainly in striated muscles (Skeletal and cardiac muscles) where it causes myositis, degenerative changes and atrophy of muscles around the cysts (Smith et al., 1972; Yassein, 1984 and Sharma and Mazaheri, 1992). Three species of Sarcocystis were detected in buffaloes S. fusiformis, S. levinei and S. buffalonis, each one has a distinct cyst wall structure. S. fusiformis and S. buffalonis form macroscopic sarcocysts and are transmitted to buffaloes via faeces of cats, while S. levinei forms microscopic sarcocysts and is transmitted to buffaloes via faeces of dogs (Dissanike & Kan, 1978 and Huong et al., 1997). Generally,
Macroscopic species of *Sarcocystis* are non pathogenic, whereas some microscopic species may cause anemia, weight loss, abortion and death in their hosts (Dubey *et al*., 1989).

In Egypt two species (*S.fusiformis* and *S.levinei*) were detected in buffaloes in different localities by several authors (El-Afifi *et al*., 1962, Abdel-Rahman, 1975, Ali 1985, Hassanien 1992, El-Saieh, 1998 and Mandour *et al*., 1998). The ultrastructure of *S.fusiformis* was studied by Abdel-Ghaffar *et al*. 1978 and Mandour *et al*. 1998. Ocular musculature appears to be a preferred site for development of *Sarcocystis* in different intermediate hosts (Juyal *et al*., 1982). Therefore, the aim of the present work was to determine the incidence of different species of *Sarcocystis* in ocular muscles of buffaloes in Assiut Governorate, as well as identify the species and describe the morphological characters of each species and their ultrastructure.

**MATERIALS AND METHODS:**

Collection of samples: Ocular muscle samples were collected from 150 buffaloes (80 males under 2 years and 70 females more than 5 years) from different localities of Assiut Governorate.

Parasitological examination:

- Each sample was examined macroscopically for detection of macroscopic cysts.
- Microscopical examination was also done by compressing of small pieces of muscle between two slides (Mowafy, 1993).
- For detection of liberated trophozoites, homogenization of muscle samples was done with physiological salt solution (Ali, 1985).
Histomorphological examination:

-Some muscular specimens from positive cases and macroscopic cysts were fixed in 10% neutral buffered formalin, these samples were dehydrated in alcohols and stained with haematoxylin and eosin then examined microscopically (Bancroft and Stevens, 1993).

-Other selected specimens were fixed in 2.5% buffered glutaraldehyde and processed for electron microscopy (Johannessen, 1978).

-Semithin sections were made and examined by light microscope.

-Ultrathin sections were then prepared, stained with urinyl acetate lead citrate stain and examined by JEOL-EM 100 CXII electron microscope.

RESULTS:

Out of 150 ocular muscle samples of buffaloes, 55(36.66%) were infected with Sarcocystis. Incidence of infection with Sarcocystis in males was 15% and in females was 61.42%. Two species of Sarcocystis were detected in the present work, the first species was S.levinei which was detected in 52 (34.66%) of examined animals. Their incidence in males was 15% and in females was 57.14%. The second species was S.fusiformis which was detected in 14 (9.33%) of examined animals. Their incidence in males was 1.25% and in females was 18.57%. Mixed infection of Sarcocystis species was detected in 11(7.33%) of examined animals. The infection rate in animals more than 5 years was 61.42% while in animals less than 2 years was 15% (Table 1).

Table (1): Incidence of S.levinei and S.fusiformis in ocular muscles of buffaloes.

<table>
<thead>
<tr>
<th></th>
<th>Examined animals</th>
<th>Infected animals</th>
<th>Single S.levinei</th>
<th>Single S.fusiformis</th>
<th>Mixed infection</th>
<th>Total S.levinei</th>
<th>S.fusiformis</th>
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<tr>
<td></td>
<td>No.</td>
<td>%</td>
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<tr>
<td>Males less</td>
<td>80</td>
<td>12</td>
<td>15</td>
<td>13.75</td>
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<td>1</td>
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<tr>
<td>than 2 years</td>
<td>70</td>
<td>43</td>
<td>61.42</td>
<td>30</td>
<td>42.85</td>
<td>3</td>
<td>4.28</td>
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<tr>
<td>Females more</td>
<td>150</td>
<td>55</td>
<td>36.66</td>
<td>41</td>
<td>27.33</td>
<td>3</td>
<td>2</td>
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<td>than 5 years</td>
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Morphological description:

1-Sarcocystis fusiformis: It apprears grossly as broad, white cyst not embeded deeply between the ocular muscle fibers, measuring 3.0-15.0 mm long (mean 9.0 mm) and 1-4 mm wide (mean 2.5 mm) (Plate I, Fig.1). Histopathological examination revealed that the cyst wall is relatively thin measuring 2-4 μ and provided with cyst wall-processes (cytophaneres) which are cleared by semithin section as a highly branched villar protrusions like a cauliflower. These villar protrusions are attached to each other giving the wall spongy appearance. Septa arise from the inner surface of cyst wall dividing each cyst into chambers containing groups of crescentic trophozoites. These trophozoites measure 12-15.4μ x 2.5-4.5μ (13.5x3.8μ). Between the inner side of the wall and these chambers there are a number of metrocytes of variable size (Plate I Fig. 2, 3). Ultrastracture examination of the cyst
has shown that the parasitophorous vacuolar membrane (PVM) of the villar protrusions is wavy and possesses small papillary projections. Their ground substance contain numerous microfilaments (MF) and electron dense granules (Ed). Longitudinal section of bradyzoites shows numerous micronemes (Mn) at the apical end. The middle part of the bradyzoite contains polysaccharide granules (Ps), and refractile bodies (Rb) of variable size while the nucleus is located at the posterior end (Plate I 4, 5).

2- *Sarcocystis levinei*: It was detected microscopically. It appears as a small spindle shaped cyst measuring 0.40-0.86 mm long (0.70mm) and 0.10-0.17mm wide (0.14mm). The cyst wall is thick measuring 4-7μ and appeared finely striated by light microscope. Examination of semithin section cleared that these striations are finger-like projections located on well distinct basement membrane. They are not branched and unequal in length. Their distal end is dome-shaped while their anterior end is tapering toward the host cell. No septa were detected in examined cysts and their trophozoites are collected in large number in groups. The trophozoites measuring 9.5-11.8x 2-3.5μ (10.8x3μ). Overcrowding of trophozoites obscure any morphological details inside the cyst except the presence of few number of small rounded metrocytes, adhering to the inner side of the wall (Plate II Fig. 1, 2). Ultrastructure of the cyst has shown that the surface of villar protrusions is smooth and not papillated while their ground substance is finely granulated and lacks microfilaments. The bradyzoite has the same morphological characters of *S.fusiformis* in addition to presence of ductules of rhopty (Dr) at the anterior part. Longitudinal section of metrocytes appears as peculiar cell, where it is characterized by thick and infolding pellicle. It is provided with two refractile bodies (Rb) one is small and anterior to nucleus while the second is so large and posterior to the nucleus. These bodies are osmiphilic, electron dense and homogenous. Nucleus is located at the middle of metrocyte and it has a small round nucleolus. Septa appear as a very thin double membrane behind the metrocyte (Plate II 3, 4).

**DISCUSSION:**

Ocular muscles play an important role in dissemination of *Sarcocystis* infection. Because eye and its surrounds musculature are considered as unedible parts for human consumption. So these parts are thrown away by the butchers and eaten by carnivorous animals which are the definitive hosts of *Sarcocystis*. In the present work *Sarcocystis* infection was detected in 36.66% of examined samples, the incidence of both *S.levinei* and *S.fusiformis* was 34.66% and 9.33% respectively. This result is considered so high than that detected by Mansour (1994) who detected *S.fusiformis* in 0.48% of examined buffaloes. This difference may be attributed to the method of diagnosis, which depended only on macroscopical examination of buffaloes in abattoirs.

The prevalence of infection of both species of *Sarcocystis* in the present work was high in females (61.42%) than males (15.0%). This result agrees with Ali (1985) and El-Saieh (1998). The difference of infection rate between males and females is
attributed to that the prevalence of Sarcocystis infection which is mainly high in aged animals.

Concerning the morphological studies of Sarcocystis, the results of the present work agrees with the opinion of Mandour (1974) who mentioned that: the cyst wall and morphological characters of villar protrusions are considered the main critaria to differentiation between species of Sarcocystis. The morphological characters and ultrastructure studies of cyst wall and bradyzoites of S. fusiformis in the present work agrees with the description of the previous authors as Mandour (1974) and Abdel-Ghaffar et al. (1978), but it differs from that described by Mandour et al., (1998) which was known as S. fusiformis aegyptii. This differance may be attributed to the age of the cyst.

The present morphological study of S. levinei is considered the first report in Egypt, especially the description of the cyst wall and their ultrastructure. The villar protrusions of the cyst wall of S. levinei are characteristic and easily differentiate it from those of the cyst wall of S. fusiformis. The morphological characters of S. levinei in the present work agrees with the description of Dissanaike & Kan (1978). Although Sarcocystis species found in livestock are considered host specific, the manner of arrangement of the villar protrusions in S. levinei is closely similar to that of S. cruzi of cattle described by Claveria et al., 2001. The large number of villar protrusions on the surface of the cyst of S. levinei are important for nutritional requirment. This large number increase the surface area which lead to enhance the nutritional absorption.

Plate (I): Morphological characters and ultrastructure of Sarcocystis fusiformis
1- Grossly visible sarcocysts in ocular muscle.
2- Light micrograph of sarcocyst showing septa, metrocyes and aggregation of bradyzoites (H&E X1000).
3- Semithin section of sarcocyst. Note host cell (Hc), villar protrusions (Vp) and septa (S). Toulidine blue stain X1000.
4- Transmission electron micrograph of villar protrusion. Note microfilaments (Mf), Electron dense granules (Ed), wavy parasitophorous vacular membrane (PVM) and cyst wall (Cw) X10000.
5- Transmission electron micrograph of bardyzoites. Note septa (s) micronemes (Mn) at the apical end, polysaccharide granules (Ps), refractile bodies (Rb) and nucleus (N) at the posterior end X5000.
Plate (II): Morphological characters and ultrastructure of *Sarcocystis levinei*.

1-Light micrograph of sarcocyst in ocular muscle. Note fine striation cyst wall (opposing arrows) H&E X1000.

2-Semithin section of sarcocyst. Note finger like projection of the villar protrusions. Toulidine blue stain X 1000.

3-Transmission electron micrograph of the same cyst. Note ductule of rhoptry (Dr) at the anterior end of merozoite, section of metrocyte (M) and their nucleus (N), section of the villar protrusion (Vp) and host cell (Hc) X2700.

4-Transmission electron micrograph of metrocyte. Note refractile body (Rb), nucleus (N), endoplasmic reticulum (Er) ground substance (G) and very thin septa (S). X4000.
The refractile bodies in the metrocytes of *S. levinei* in the present work are so large. These are similar to that of *Eimeria ninakolyakimovae* sporozoite of goat described by Scholtyseck (1979). He added that, the refractile bodies are one of the most prominent fine structure of subphylum Apicomplexa.

According to the previous results we could concluded that:

Ocular muscles are considered a preferable site for *Sarcocystis* especially *S. levinei*. It is, therefore recommended that *Sarcocystis* cysts should be looked for in ocular muscles of humans whenever clinical manifestations devotes muscle disease. Ultrastructure studies of the cyst wall could be used as one of important method to identify different species of *Sarcocystis*.

REFERENCES:


دراسات عن الساركوسستس العيني في الجاموس في محافظة أسيوط

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تم فحص عدد 150 عينة من عضلات العيون في الجاموس لعمل دراسة طفيلية عن تواجد الساركوسستس في محافظة أسيوط. واشتملت تلك العينات على 80 عينة من الذكور أقل من عامين و70 من الإناث أكبر من خمس سنوات. وبلغت نسبة الإصابة الساركوسستس 36.3% حيث تم اكتشافها في 55 حيوان.

بلغت نسبة الإصابة في الذكور 15% وفي الإناث 61.4%. وتم تصنيف الساركوسستس إلى نوعين الأول: ساركوسستس فيوزفورماس حيث كانت نسبة الإصابة به 9.33% فقد بلغت نسبة الإصابة في الذكور 12.35% وفي الإناث 18.57%. والنوع الثاني هو ساركوسستس ليفيني، وقد كانت نسبة الإصابة به 24.66% حيث كانت نسبة الإصابة في الذكور 15% وفي الإناث 57.14%. وقد كانت نسبة العدوى المشتركة من النوعين 3.37%. وقد تم وصف الشكل المورفولوجي والتركيب الدقيق لكل نوع بواسطة الميكروسكوب الإلكتروني.

وقد أثبت التجارب الحالية أن عضلات العين تعتبر مكاناً مفضلاً لطفيل الساركوسستس وخاصة ساركوسستس ليفيني ولذلك يوصي الباحثون بفحص عضلات عيون الإنسان في حالة الشكوى من أعراض مرضية بها.