

Original article

# Prevalence of *Sarcocystis* spp. in Sheep Cardiac Tissue at Al Bayda Municipal Abattoirs: A Histological Cross-Sectional Study

Ayiman Aboulqassim\* , Fawzia Mohamed 

Faculty of Veterinary Medicine, Omar Al-Mukhtar University, Al-Bayda, Libya

Email: [eman.saleh@omu.edu.ly](mailto:eman.saleh@omu.edu.ly)

## Abstract

Sarcocystis is an intracellular cyst-forming protozoan parasite belonging to the phylum Apicomplexa. It infects a wide range of vertebrates, including sheep, and has significant veterinary and public health importance. This study aimed to determine the prevalence of *Sarcocystis* spp. infection in cardiac tissues of slaughtered sheep at the municipal abattoirs of Al Bayda, Libya, using histopathological examination, focusing on cases with no visible macroscopic cysts. Fresh cardiac tissue samples were collected from 50 slaughtered sheep (both sexes and various ages) between May 2025 and April 2026 from hearts lacking visible macroscopic sarcocysts. Tissues were fixed in 10% neutral buffered formalin, processed by routine histological methods, sectioned at 5 µm thickness, and stained with Harris's hematoxylin and eosin (H&E) for microscopic examination. Histopathological examination revealed *Sarcocystis* infection in all 50 samples (100% prevalence). Microscopic thin-walled sarcocysts with similar morphology were observed in cardiac tissues, while no macroscopic cysts were detected by naked-eye inspection. No inflammatory reaction was observed around the sarcocysts (absence of pericystic inflammatory infiltrates), and no prominent degenerative or inflammatory changes were noted in the infected cardiac tissue. The study concludes that *Sarcocystis* infection is highly prevalent (100%) in the cardiac tissue of sheep slaughtered in Al Bayda abattoirs, with the absence of macroscopic cysts and lack of pericystic inflammation. Histopathological examination is of great diagnostic value for detecting *Sarcocystis* infection, especially when macroscopic lesions are absent, and gross examination alone is insufficient to estimate the true prevalence of the parasite.

**Keywords.** Sarcocystis, Cardiac tissues, Sheep, Histopathological examination, Al Bayda, Libya.

## Introduction

The *Sarcocystis* (phylum: Apicomplexa) is an intracellular cyst-forming protozoan parasite that infects a wide range of vertebrates, and is the most common protozoan found in animal muscle tissue that is meant for human consumption [1, 2]. Approximately 220 species of *Sarcocystis* spp. have been identified in birds, mammals, and reptiles [3, 4]. Sheep and cattle can contract *Sarcocystis* infections from many species. These parasites cause macrocysts or microsarcocysts to form in the skeletal, cardiac, and central nervous systems [5]. The life cycle of *Sarcocystis* requires two necessary hosts: the definitive host (a carnivorous animal, such as a dog), in which sporogony and gametogony occur in the intestinal lumen, producing intestinal sarcocystosis, and the intermediate host (an herbivorous or omnivorous animal) in which merogony and cyst (sarcocyst) formation develop in the muscles, leading to muscular sarcocystosis [6]. Developing in a two-host, intermediate, and definitive host, humans can serve as an intermediate host and definitive host for different *Sarcocystis* species. Consequently, humans are definitive hosts when they consume tissue cysts with undercooked meat, sarcocysts [5]. Animals that are significantly afflicted by sarcocystosis commonly exhibit clinical signs such as pyrexia, debilitation, cachexia, and loss of body mass, as well as compromised wool quality, alopecia, and diminished lactation output. Furthermore, the meat industry incurs considerable financial detriment due to the rejection of carcasses from heavily infected animals that display discernible *Sarcocystis* cysts [7, 8].

Ovine species exhibit a susceptibility to infection by six distinct species of *Sarcocystis*. Among these parasites, *S. tenella* (*S. ovis*), *S. arietianis*, *S. microps*, and *S. mihouensis* are characterized by canids serving as their definitive hosts. Conversely, the remaining two species, *S. gigantea* (*S. ovifelis*) and *S. medusiformis*, identify felines as their ultimate hosts [9, 10]. It is significant to note that *S. tenella* and *S. arietianis* are categorized as pathogenic and are known to develop microscopic cysts, while *S. gigantea* and *S. medusiformis* are classified as non-pathogenic and are responsible for the formation of macroscopic cysts within ovine hosts [11]. This study aimed to determine the prevalence of *Sarcocystis* in slaughtered sheep at the Al Bayda Municipal Abattoirs in Libya, specifically in cardiac tissues where macroscopic cysts are absent. This process can be performed using histopathological examination.

## Methods

From May 2025 to April 2026, samples were systematically collected and underwent gross (macroscopic) inspection during the slaughter process at the Al Bayda Municipal Abattoirs in Libya. The sampling included sheep of both sexes and various age groups. Each carcass underwent examination for visible macroscopic cysts that indicated the presence of cyst-forming *Sarcocystis* spp. Following this, fresh cardiac tissue specimens were collected specifically from 50 slaughtered sheep that exhibited no macroscopic cysts during

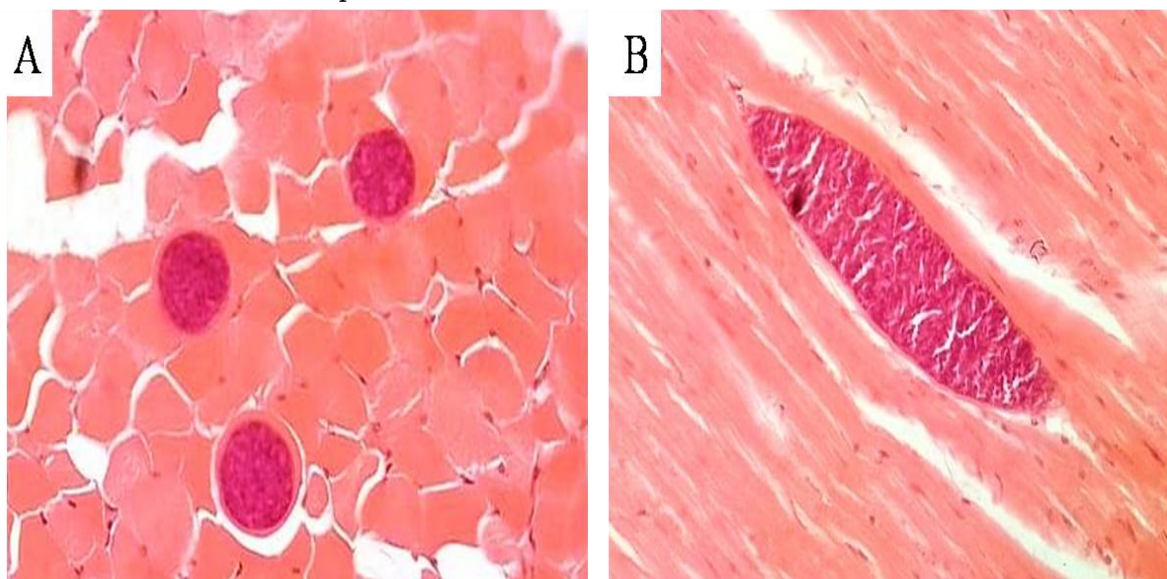
the gross examination. Immediately after collection, the cardiac tissues were fixed in 10% neutral buffered formalin to preserve their cellular and tissue architecture. The fixed samples were then processed using a standard histological workflow that included graded dehydration through increasing concentrations of ethyl alcohol, followed by clearing of tissues and embedding in paraffin wax. Paraffin-embedded blocks were sectioned on a microtome to obtain serial histological sections of 5  $\mu\text{m}$  thickness, which were mounted onto charged glass slides to enhance tissue adherence, and stained for histopathological assessment with Harris's hematoxylin and eosin (H&E) [12].

## Results

Histopathological examination was conducted on cardiac tissue samples collected from 50 slaughtered sheep at Al Bayda Municipal Abattoirs, Libya. Microscopic assessment demonstrated that *Sarcocystis* infection was highly prevalent, with an overall infection rate of 100%. Sarcocysts were detected in all examined animals, confirming that the parasite was widespread among the slaughtered population during the study period. Sarcocysts observed in the cardiac tissues showed similar morphologies across samples. The consistency of the sarcyst type suggests that the tissue-stage infection shared common developmental and morphological characteristics within the examined group. Importantly, the parasites were clearly identifiable at the microscopic level, providing direct histopathological evidence of infection.

A key histopathological observation in the present study was the absence of an inflammatory reaction around the sarcysts within the cardiac muscle. No pericycstic inflammatory infiltrates were detected in association with the cysts, and inflammatory changes were not observed as a consistent feature of the lesions. In addition, most affected samples were reported to be free from prominent degenerative and inflammatory alterations, indicating that although the parasite was present, it was not accompanied by extensive tissue disruption in the cardiac tissue at the microscopic level.

Interestingly, while sarcysts were readily detectable histologically, macroscopic sarcysts were not observed in any of the examined samples. None of the animals showed visible sarcysts in cardiac tissues when assessed by naked-eye inspection. This finding emphasizes that gross examination alone may underestimate or fail to detect *Sarcocystis* lesions. Overall, the study outcomes demonstrate a pattern characterized by widespread microscopic infection without pericycstic inflammatory pathology in cardiac tissue. The results highlight the diagnostic value of histopathological examination for confirming *Sarcocystis* infection in sheep, especially in cases where macroscopic lesions are absent.



**Figure 1. Histopathological findings in *Sarcocystis* spp. , sheep, hematoxylin and eosin. A) A cross-section of myocardium showing: Within multiple myofibers, several oval to round sarcocysts are clearly visible, adapted to the cross-sectional plane. Each sarcocyst is bordered by a thin, smooth, eosinophilic cyst wall that lacks any radial spines or protrusions. Inside the cyst, numerous crescent-shaped bradyzoites (zoites) are packed densely, often arranged peripherally around a granular, basophilic central core. (100X). B) longitudinal section of myocardium showing: thin-walled septate sarcocyst containing a myriad of banana-shaped bradyzoites in the myocardium. The bradyzoites stain deeply basophilic with H&E. (400X)**

## Discussion

The present study demonstrates a high prevalence and significant parasite burden of *Sarcocystis* in the cardiac tissue of slaughtered sheep, as indicated by the presence of numerous sarcocysts. Such findings are important for both animal health and public health. It is significant to note that pathogenic species form microscopic cysts, whereas non-pathogenic species form macroscopic cysts within ovine hosts [11]. Consequently, the detection of abundant sarcysts in the current cardiac tissues not only confirms the occurrence of *Sarcocystis* infection in sheep but also suggests the potential presence of pathogenic

microscopic stages that may remain undetected by routine gross inspection, thereby strengthening the epidemiological and diagnostic significance of the histopathological findings. In intermediate hosts (including sheep), *Sarcocystis* infections are associated with economic and health impacts such as poor growth, reduced productivity, carcass condemnation at meat inspection, and, in some situations, reproductive losses [13]. These effects may be subtle in many animals, but the presence of heavy cyst burdens in edible tissues highlights the need for control measures at slaughter. From a zoonotic perspective, *Sarcocystis* is relevant to human health because people can become infected mainly through the consumption of undercooked meat containing viable sarcysts.

Human intestinal sarcocystosis has been linked to the consumption of raw or undercooked meat containing viable *Sarcocystis* tissue cysts (bradyzoites). After ingestion, the bradyzoites excyst in the small intestine, invade the intestinal mucosa, and develop into sexual stages, with subsequent egg/sporocyst shedding in stool. Therefore, the detection of abundant sarcysts in cardiac tissue supports the importance of meat inspection, hygienic processing, and proper cooking to reduce transmission risk [14]. Concerning organ distribution, although cardiac muscle was clearly infected in the current study, previous investigations have reported that the occurrence/intensity among organs can vary, and macroscopic cysts (when assessed across organs) have been reported as lowest in the heart compared with other examined tissues [15].

Similarly, studies evaluating the distribution pattern of *Sarcocystis* across different tissues in sheep show that the parasite preferentially localizes in particular sites rather than uniformly across all organs, supporting the idea that heart infection rates may be lower than other muscles or tissues in some settings [16]. Importantly, these studies were based on macroscopic (gross) examination of cardiac muscle, which may explain differences in detection rates and observed cyst burden compared with microscopic histology. This finding emphasizes that gross examination alone may underestimate or fail to detect *Sarcocystis* lesions, particularly when cysts are thin-walled and may not produce conspicuous macroscopic changes.

## Conclusion

In conclusion, the research presents compelling evidence indicating that *Sarcocystis* infection among slaughtered ovine specimens within the examined abattoir region exhibits an exceptionally high prevalence rate of 100%, characterized by sarcysts displaying a consistently thin-walled morphology, and an absence of any discernible pericyclic inflammatory response within cardiac tissues. The detection of the parasites was effectively accomplished through histological methods; conversely, macroscopic lesions were not identified, thereby underscoring the critical significance of histopathological evaluation for the precise diagnosis and comprehensive assessment of *Sarcocystis* infection in sheep.

## Conflicts of Interest

The authors declare no conflict of interest.

## References

1. Heckerroth AR, Tenter AM. Sarcocystiosis. In: Protozoal abortion in farm ruminants: guidelines for diagnosis and control. 2007. p. 172.
2. Mandal S. Veterinary protozoology. In: Textbook of veterinary parasitology. Springer; 2025. p. 473-571.
3. Dubey JP, Speer C, Fayer R. Sarcocystosis of animals and man. 1989.
4. Fayer R. *Sarcocystis* spp. in human infections. Clin Microbiol Rev. 2004;17(4):894-902.
5. Mohammed RG, Manal HH, Dragh MA. Histopathological diagnosis for *Sarcocystis* spp. in slaughtered sheep and goats in Misan governorate, Iraq. Int J Health Sci. 2022;6(S8):2119-31.
6. Hussein SN, Ibrahim AA, Shukur MS. Histopathology and molecular identification of *Sarcocystis* species forming macrocysts in slaughtered sheep and goats of Duhok, Iraq. Vet Res Forum. 2023.
7. Abdel-Hamied AM, et al. Prevalence, morphology, and genetic relationship of *Sarcocystis* species in naturally infected sheep. Egypt J Vet Sci. 2025:1-16.
8. Pinto MS, et al. Sarcocystosis in farm animals in Brazil: a One-Health approach. Vet Sci. 2025;12(9):842.
9. Marandykina-Prakienė A, et al. Molecular identification of *Sarcocystis* species in sheep from Lithuania. Animals. 2022;12(16):2048.
10. Dong H, et al. Sarcocystis species in wild and domestic sheep (*Ovis ammon* and *Ovis aries*) from China. BMC Vet Res. 2018;14(1):377.
11. Gerab RA, et al. Prevalence and distribution of *Sarcocystis* in buffaloes and sheep in Egypt. J Adv Vet Res. 2022;12(3):302-7.
12. Luna LG. Manual of histologic staining methods of the Armed Forces Institute of Pathology. 1968. p. xii, 258.
13. Fayer R, Esposito DH, Dubey JP. Human infections with *Sarcocystis* species. Clin Microbiol Rev. 2015;28(2):295-311.
14. Harris V, et al. Human extraintestinal sarcocystosis: what we know, and what we don't know. Curr Infect Dis Rep. 2015;17(8):42.
15. Latif B, et al. Prevalence of *Sarcocystis* spp. in meat-producing animals in Iraq. Vet Parasitol. 1999;84(1-2):85-90.