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Cyst Wall Ultrastructure of Two *Sarcocystis* spp. from European Mouflon (*Ovis ammon musimon*) in Germany Compared with Domestic Sheep

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Abstract: Muscle samples from six wild and two captive European mouflons (*Ovis ammon musimon*) in Germany as well as one domestic sheep from a German zoo were infected with sarcocysts (*Sarcocystis* Sarcocystidae, Apicomplexa). *Sarcocystis tenella* and *S. arieticanis* were identified by light and electron microscopy. Both species are determined for the first time from wild sheep, and this is the first description of *S. arieticanis* from wild sheep.

Key words: *Sarcocystis tenella*, *S. arietianis*, *Ovis ammon musimon*, domestic sheep, zoo, free range, Germany.

Sarcocysts of at least four *Sarcocystis* species occur in domestic sheep (*Ovis ammon* L., hemerotype) (Dubey et al., 1989): *Sarcocystis tenella* (Railliet, 1886), *S. gigantea* (Railliet, 1886), *S. arietianis* Heydorn, 1985, and *S. medusiformis* Collins et al., 1979. The first three species occur globally; *S. medusiformis* has been reported only from Australia and New Zealand (Dubey et al., 1989).

Sarcocysts occasionally were found in free-ranging mouflon (*Ovis ammon* L., agriotype), conspecific with domestic sheep. Most of these reports did not include detailed descriptions. Nigro et al. (1991) first described a *Sarcocystis* species in detail from wild sheep (*O. ammon musimon*) in Italy; the parasite probably was *S. tenella*, based on the fine structure of the cyst wall. In our study, we extend the host range of two *Sarcocystis* species to free-ranging mouflon, and compare them with same species of sarcocysts from a domestic sheep, occurring in the same area.

Muscle samples from eight European mouflons (*Ovis ammon musimon*) were evaluated for the occurrence of sarcocysts in 1993 and 1994. Six of these mouflons originated from a population near Niederfinow, Land Brandenburg, Germany (52°51′N, 13°40′E), founded in 1981 (Briedermann, 1990). The other two mouflons came from two zoos (Tierpark Berlin-Friedrichsfelde and Tierpark Eberswalde, Land Brandenburg, Germany). Sarcocysts from a 7-year-old domestic sheep that died in June 1993 were used for comparison; this sheep as well as its parents were born in the zoo Tierpark Berlin-Friedrichsfelde. Fresh samples of muscle tissue from head, neck, larynx, heart and loin (Musculus psoas major) were tested. Individual sarcocysts found in the musculature were extracted from the muscle fibers under a dissecting microscope for fresh-state examination or prepared for transmission electron microscopical (TEM) investigations. The size of the bradyzoites (cystozoites) was determined in fresh preparations. Their length was taken by measuring the more or less bent median line from pole to pole. Their width was measured at the widest diameter. For TEM investigation the sarcocysts were fixed according to Pospischil and von Bomhard (1979). After repeated washing with 0.1 mol phosphate buffer, they were post-fixed in 2% osmium tetroxide (Serva Feinbiochemica Heidelberg, Germany) solution, dehydrated in ethanol and embedded in Epon® 812 (Serva Feinbiochemica Heidelberg, Germany), and exposed to polymerization for 3 days. The TEM investigations were carried out using an EM 902 A microscope (Zeiss, Oberkochen, Germany).

We found *Sarcocystis tenella* (Railliet, 1886) (synonyms: *S. ovicantis* Heydorn et al., 1975; *Sarcocystis* sp. Nigro et al., 1991) in all eight mouflons and the domestic sheep. The sarcocysts were up to 2.4 mm long and 186 μm wide. The cyst walls were 1.08 to 3.85 μm thick (n = 130) and had...
a palisade-like texture, with finger-shaped villar protrusions that were positioned closely side by side (Figs. 1 and 2). The protrusions were 2.10 to 3.85 μm long and 0.42 to 2.14 μm wide (n = 105). The distance between the protrusions was 0.2 to 0.5 μm at their base; in mouflons the mean ± SD value was 0.32 ± 0.10 μm (n = 15) and in the domestic sheep the mean ± SD value 0.43 ± 0.11 (n = 10). The protrusions had a truncated tip with plaquelike condensations (Fig. 4). Numerous fine and a few large granules lay in the core (Figs. 3 and 4). The diameters of the compartments in the region of the cyst wall ranged from 6.2 to 31.0 μm in the mouflon (x ± SD = 14.8 ± 6.6 μm, n = 50), and 11.0 to 49.5 μm in the domestic sheep (x ± SD = 28.6 ± 14.6 μm, n = 6). The diameters of the compartments in direction of the center were 6.2 to 28.5 μm in the mouflon (x ± SD = 13.2 ± 5.9 μm, n = 20) and 16.5 to 30.2 μm (x ± SD = 24.3 ± 6.4 μm, n = 6) in the domestic sheep.

All eight mouflons and the domestic sheep also had Sarcocystis arieticanis Heydorn, 1985 (synonym: Sarcocystis sp. Boch et al., 1979). The sarcocysts were up to 1.8 mm long and 286 μm wide. The cyst walls were thin (0.21 to 0.62 μm, n = 35). They had unstable hairlike, 5.7 to 11.8 μm (n = 100) long villar protrusions (Figs. 5 and 6), which contained numerous fine granules in the core (Figs. 7 and 8). The diameters of the compartments in the region of the cyst wall ranged from 20.0 to 59.9 μm (x ± SD = 35.7 ± 7.9 μm, n = 50). The diameters of the compartments in direction of the center ranged from 46.6 to 73.3 μm (x ± SD = 57.9 ± 10.3 μm, n = 10).

The four Sarcocystis species known from domestic sheep are unequivocally defined morphologically and easily well distinguished from each other (Dubey et al., 1988, 1989). They can be distinguished clearly from S. ferox Dubey, 1983 (with flattened mushroom-like protrusions) described from the bighorn sheep (Ovis canadensis), as well as from Sarcocystis sp. Foreyt, 1989 from the American mountain goat (Oreamnos americanus) in North America (Dubey et al., 1989), and Sarco-
FIGURE 3. *Sarcocystis tenella* from a mouflon; transmission electron micrograph of a longitudinal section of three villar protrusions from a 0.6 mm long sarcocyst from the diaphragm. Note the two large granules in the core of two protrusions. Bar = 0.4 μm.

FIGURE 4. *Sarcocystis tenella* from a domestic sheep; transmission electron micrographs of the cyst wall of a 0.6 mm long mature sarcocyst from Musculus psoas major. Longitudinal section of the villar protrusions; note the electron-dense plaques on the top (arrowhead). Inset: cross-section of the protrusions. G = Ground substance. Bar = 0.4 μm.

FIGURE 5. *Sarcocystis arieticanis* from a mouflon; fresh state view of a 1.2 mm long mature sarcocyst from neck, with hairlike villar protrusions. Bar = 10 μm.

FIGURE 6. *Sarcocystis arieticanis* from a domestic sheep; fresh state view of a 1.1 mm long mature sarcocyst from Musculus psoas major, showing the hairlike villar protrusions on the top. Bar = 10 μm.
Sarcocystis sp. (2) Cornaglia et al., 1980 from the chamois (Rupicapra rupicapra) in Europe. Both sarcocyst forms from the Rupicaprinae, Rupicapra sp. and Oreamnos sp., appear similar if one compares fig. 10 by Cornaglia et al. (1980) with fig. 3 by Foreyt (1989); the protrusions are club-shaped, with two longitudinal grooves and a bundle of microtubules reaching into the ground substance. Sarcocystis tenella and S. arieticanis in sheep and the morphologically very similar species S. capracanis Fischer, 1979 and S. hircicanis Heydorn and Unterholzner, 1983 in goats are different species separated by intermediate host specificity, as shown by unsuccessful transmitting experiments with S. tenella and S. arieticanis to goats (Heydorn, 1985; Dubey et al., 1989) and with S. capracanis and S. hircicanis to sheep (Balbo et al., 1988; Dubey et al., 1989). On the other hand, cross transmission experiments by Balbo et al. (1988) provide evidence for a less marked intermediate host specificity. Most Sarcocystis spp. described from Caprinae and Rupicaprinae can be distinguished according to the ultrastructure of the cyst wall and the species of the intermediate host.

The species composition of Sarcocystis spp. in domestic sheep is well established (Dubey et al., 1988, 1989), especially for S. tenella and S. arieticanis which use canids as definitive hosts. The mature sarcocysts of S. tenella and S. arieticanis can be distinguished microscopically in the fresh state (Boch et al., 1979; Figs. 1 and 2 compared with Figs. 5 and 6) or also in standard histological preparations (Savini et al., 1993). The fine structure of the cyst wall of S. tenella was identical in our domestic sheep and the mouflon (Figs. 3 and 4) and was similar to that described previously for this parasite from domestic sheep (Bergmann and Kinder, 1975; Vlemmas et al., 1989) and from the mouflon (Nigro et al., 1991). Also in S. arieticanis, the fine structure of the cyst wall was identical in both our domestic sheep and the mouflon (Figs. 7 and 8). The TEM pictures from domestic sheep in the literature (Heydorn and Mehlhorn, 1987)
correspond well with ours. *Sarcocystis arleticanis* seems to occur less frequently than *S. tenella* (Schmidtová, 1992), but may be more frequent in western Australia (Savini et al., 1993). The present description is the first report from wild sheep.

There are different data on the size of the sarcocysts of both *Sarcocystis* species described here. *Sarcocystis arleticanis* reaches a length of 1.0 mm according to Boch et al. (1979), of 0.9 mm after Dubey et al. (1989), of 2.92 mm according to Schmidtová (1992), of 1.3 mm after Savini et al. (1993), and 1.8 mm in our material. *Sarcocystis tenella* is up to 0.58 mm long after Boch et al. (1979), seldom over 1.0 mm at 105 days post-infection after Heydorn (1985), up to 0.7 mm according to Dubey et al. (1989), up to 1.3 mm according to Savini et al. (1993), and 2.4 mm in our samples.

**LITERATURE CITED**


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