## PREVALENCE OF MARSHALLAGIA MARSHALLI (ORLOV, 1933) IN WILD RUMINANTS IN WYOMING'

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Prevalence of Marshallagia marshalli (Orlov, 1933) in wild ruminants in Wyoming was determined by screening abomasal and small intestinal contents after necropsy and by analyses of feces collected from free-ranging ruminant individuals. No M. marshalli were found in 60 moose. Only two elk fecal samples, of over 2,000 collected, were positive (a new host record) but the parasite was not found at necropsy of 60 elk. One of 51 mult deer fecal samples was positive. The nematode species is very common in pronghorn antelope and bighorn sheep.

Marsballagia marsballi, Orlov, 1933, a trichostrongylid, abomasal nematode often found in domestic sheep and goats, is also a common parasite of wild ruminants in the Rocky Mountain region of the western United States.

M. marsballi was originally described (as Ostertagia marsballi) from specimens taken from domestic sheep in Montana (1). In 1933, the genus was changed to Marsballagia (2). M. marsballi is quite different from most species of the genus Ostertagia (3) in adult morphology, size of eggs, larval development and numbers in host animals.

M. marsballi was reported as O. marsballi in 1932 from a bighorn sheep (Ovis canadensis) (4) and from a deer (Odocoileus odocoileus) in Yellowstone National Park, Wyoming (4). In 1945 M. marsballi was reported as a parasite of prongborn antelope, (Antilocarpra americana) (5) and in 1956 the nematode was listed as a parasite of several wild ruminants in Wyoming (6).

M. marsballi may be cosmopolitan in distribution (7) or according to another authority (8), the species is common only in the western U. S.

Gastrointestinal tracts and fecal samples from moose, elk, mule deer, pronghorn antelope and bighorn sheep were gathered mainly during hunting seasons, from 1963-1973. Adult worms were recovered by screening contents of the gastrointestinal tracts (at least 10% of the contents of each tract was screened). The worms were recovered, counted and sexed. Fecal analyses of worm eggs per gram (epg) feces were made using a modification of Lane's (1923) technique. Flotations (with saturated sucrose solutions) and centrifugation @ 2000 rpm were made of eggs with 3 g feces per 27 ml water.

Listed in Table 1 are the ruminant animals which were sampled for *M. marsballi*. Results of necropsy data and fecal examinations for parasite numbers are also shown in the table. In most cases necropsy examinations were on different individual host animals than those checked by fecal analyses.

TABLE	1.	Prevalence	of	Marsballa	gia	marsh	elli
		ruminants			acc	ording	80
neci	ro psi	es and fecal	an	alyses.			

Host species	No. Nec./ª No. pos./ % pos.	No. fecal exams./ No. pos./ % pos.	epg <sup>b</sup> feces
Moose	10/ 0/0	50/ 0/0	0.0
Eik	60/ 0/0	2,280/ 2/.09	2-4
Mule deer	20/ 0/0	51/ 1/2	2
Antelope	30/14/47	50/18/36	2-20
Bighorn sheep	10/ 8/80	70/47/67	2-80

Number of necropsies/number with M. Marshalli/%

b egg number/gram.

Moose may avoid the infective larvae of *M. marshalli* by browsing extensively. However, the number of moose necropsied and sampled by fecal analyses was low.

M. marsballi had not been previously reported from elk and no adult worms were found in 60 elk gastrointestinal tracts ex-

<sup>&</sup>lt;sup>1</sup>Published with approval of the Director, Univ. of Wyo., Agr. Exp. Sta. as Journal Article 686.

amined during this survey. However, the eggs of *M. marshalli* were found in elk fecal samples. Nearly 200 elk fecal samples were examined before the first *M. marshalli* eggs were found and nearly 1,000 samples were checked before the second positive fecal was found.

Apparently, *M. marsballi* is absent in many deer in Albany Co., Wyoming. Deer also ingest much browse which would not carry infective larval stages of the parasite.

M. marsballi is common in pronghorn antelope and bighorn sheep, especially the latter, but numbers of the parasite usually do not exceed 600 female worms per antelope or 2,000-2,400 female worms per bighorn sheep. M. marsballi worm egg numbers in feces of antelope and bighorn sheep were low (Table 1).

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