HELMINTH PARASITES OF THE BLACK-TAILED JACK RABBIT (Lepus californicus melanotis Mearns) IN SOUTHWESTERN KANSAS

by

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INTRODUCTION AND REVIEW OF LITERATURE

The lagomorphs appear to be one of the most thoroughly investigated groups of wild animals in respect to their parasites. Cottontail Rabbits (*Sylvilagus* spp.), Snowshoe Hares (*Lepus americanus*), and European Rabbits (*Oryctolagus cuniculus*) have been the object of a great many parasitological studies. Numerous studies of this nature can be related to wide distribution, abundance, and relative ease in capturing rabbits and hares.

A review of the literature revealed that the above relatives of the jack rabbit have been studied much more thoroughly for parasites than other members of the lagomorphs. Through scattered reports of jack rabbit parasites over a period of years these hares were found to harbor several of the same parasites as other lagomorphs. However, the need for a complete study of jack rabbits appeared to be a necessity if a more thorough knowledge was to be gained of their parasitic fauna.

In connection with an ecological study, by another worker, an endoparasite study was made on the Black-tailed Jack Rabbit (*Lepus californicus melanotis* Mearns) in southwestern Kansas. The parasite study began in September, 1956 and extended through September, 1957. A drought existed in southwestern Kansas from approximately 1952 to 1957 at which time the jack rabbits were greatly concentrated on the edge of the sand-hills and adjacent cropland. The greater concentration of hares afforded an ideal time to study their endoparasites. Theoretically, close contact with each other should have provided a good chance for them to
become infected with a variety of parasites.

Goeze in 1782, according to Stiles (1896), described *Taenia pectinata* from hares and rabbits. Stiles placed it in synonymy with *Cittotaenia pectinata*. Stiles credited the work of Curtice in 1887, on young stages of rabbit tapeworms, as bringing leporine cestodes into prominence. The reason for this seemingly important discovery was that it was believed that these young rabbit tapeworms were involved in the life cycle of tapeworms of cattle, sheep, and other animals. According to Hall in 1910, as stated by Meyer (1955), Brandegee reported the first authenticated case of *Multiceps serialis*¹ in the United States in 1890, which was found in *Lepus californicus*² from California.

Curtice (1892) reported an unidentified species of *Taenia* from *L. texianus* in Colorado and coenuri from *L. texianus* and *L. californicus* from Texas and California respectively. Stiles (1896) published a revision of the tapeworms of hares and rabbits in which tapeworms were reported from *L. melanotis*. The name he used for these tapeworms was *Davainea salmoni* which is now recognized as *Raillietina stilesiella* by Wardle and McLeod (1952).

Palmer (1897) reported that a tapeworm (*Taenia*), ticks

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¹ *Multiceps serialis*, *Multiceps sp.*, *Taenia serialis* and *Coenurus serialis* are scientific names used by different authors in this thesis for the cystic stage, coenurus, of the adult tapeworm *Multiceps serialis* found in dogs, coyotes and related animals.

² *Lepus californicus* includes the following scientific names in this thesis: *L. californicus*, *L. californicus californicus*, *L. texianus*, *L. californicus texianus*, *L. melanotis*, *L. californicus melanotis*, *L. californicus deserticola*, and *L. californicus merriami*. 

(Ixodes), and fly larvae (Cuterebra) were the most important parasites of the jack rabbit. He also stated that L. californicus and the White-tailed Jack Rabbit (L. campestris\textsuperscript{1}) were hosts for Coenuurus serialis.

Species of Cittotaenia were reported from L. melanotis by Lyman (1902). Ackert (1915) showed that Taenia serialis cysts from L. campestris would not develop to adult tapeworms when fed to fowl. The Anoplocephalidae were monographed by Douthitt (1915). A monograph was published by Hall (1916) on the parasitic nematodes of the orders Rodentia, Lagomorpha and Hydracoida. Two lagomorph nematodes mentioned by him are Dermatoxys veligera from L. c. melanotis and Dirofilaria scapiceps (sym. Filaria scapiceps) from L. campestris.

Meggitt (1924) published on the cestodes of mammals including lagomorph cestodes. Yorke and Maplestone (1926) in a comprehensive work reported all known parasitic nematodes of vertebrates. McCambell (1926) listed some parasites of L. c. melanotis and L. townsendii from eastern Colorado. Multiceps serialis was found in both species of jack rabbits and Cittotaenia spp. was found in nine of ten black-tailed jack rabbits studied.

An extensive monograph on the cestode family, Anoplocephalidae, was written by Baer (1927). One of the early compilations on rabbit parasites and diseases was done by Schwartz and Shook (1928). Occurrence of Trichostrongylus colubriformis, a ruminant parasite,  

\textsuperscript{1} Lepus townsendii includes the following scientific names in this thesis: L. campestris and L. townsendii.
in *L. c. melanotis* was reported from Nebraska by Skidmore (1932). A one-year study on helminth parasites of 420 Snowshoe Hares (*L. americanus*) in Canada, was published by Boughton (1932). Vorhies and Taylor (1933) in a bulletin on the Antelope Jack Rabbit (*L. alleni*) and *L. californicus* ssp. in Arizona reported *Multiceps* sp., *Nematodirus* sp., and *Dermatoxys veligera* in both of these hares as well as a specimen of *Raillietina* sp. from *L. alleni*. Four species of cestodes and three species of nematodes were discovered by Ward (1934) from *L. californicus* collected in Oklahoma. Erickson (1944) reported a ten-year survey of the helminth parasites of *L. americanus* in Minnesota.

The larval stage of *Dermatoxys veligera* was recovered from the cecum of *L. alleni* from Arizona by Dikmans (1931). Two new species of the genus *Nematodirus*, one from *L. c. texianus* and one from *L. alleni*, were named by Dikmans (1937).

Further contribution to the knowledge of lagomorph parasites was made by Arnold (1938) in his work on anoplocephaline cestodes of rabbits of North America. In Wyoming, Scott (1943) recovered lung worms from *L. townsendii*. Olsen (1948) reported *L. c. merriami* as a reservoir host for the common liver fluke, *Fasciola hepatica* in Texas. Wardle and McCleod (1952) in a comprehensive work on cestodes, brought the tapeworm information of the world up to date. Rohrbacher and Ehrenford (1954) created a new genus, *Biogastranema*, from nematodes found in *L. c. californicus* from California. Lechleitner (1955) found one nematode species and four tapeworm species in *L. c. californicus* from California.
A large Multiceps serialis cyst was discovered by Grundmann, Parker, and Stagg (1955) in L. c. deserticola in Utah. Honess and Winter (1956) listed several parasites of jack rabbits in Wyoming. Grundmann (1957) examined L. c. deserticola in Utah and found three species of nematodes.

There were five objectives in the present helminth survey: (1) collect jack rabbits throughout the year to investigate season fluctuations in incidence of parasitism, (2) investigate whether the total number of each species of helminth varied during the year, (3) determine if the jack rabbits in this area acted as reservoirs for cattle or sheep parasites, (4) observe if the endoparasites were associated with mortalities among jack rabbits, and (5) present a clearer picture of jack rabbit helminths in Kansas.

MATERIALS AND METHODS

One hundred and thirty Black-tailed Jack Rabbits (Lepus californicus melanotis Mearns) were examined for endoparasites over a thirteen-month period from September, 1956, through September, 1957. These jack rabbits were all collected within a ten-mile radius of Lakin, Kansas, in Kearny County. Monthly collections were made except for November, 1956, and January, March and May, 1957, thus making a total of nine collections. From ten to twenty jack rabbits were collected in each of the nine collecting periods. Larger collections were made when there was a lapse of one month between collecting periods, than when consecutive
monthly collections were made. These hares examined for endo-
parasites were selected from a larger collection of from approxi-
mately 20 to 60 animals per collection. All jack rabbits were
weighed, measured, aged, and information pertaining to reproduc-
tion was collected by Frank Bronson (Bronson, 1957).

The jack rabbits were divided into two age-classes, adults
and juveniles. Aging was based on size and condition of the
reproductive organs and on epiphyseal closure of the humeri. By
using a combination of the two aging methods, the aging was con-
sidered accurate from July, 1956, through November, 1956. During
December, January and February there may have been some degree of
inaccuracy in aging because of the difficulty in telling the older
juveniles from the adults at that time of year. After March, 1957,
Bronson considered the aging methods to be accurate again.

Collections of the jack rabbits were made with a .22 caliber
rifle and a 16-gauge shotgun. Jack rabbits to be used for the
parasite study, were selected from those showing no apparent shot
damage of the digestive tract. Since a few of the animals in the
first two collections had their digestive tracts damaged slightly
by shotgun pellets, subsequent animals were selected from those
shot with a rifle.

All jack rabbits were examined within a few minutes to approxi-
mately 15 hours post-collection. They were taken to Lakin where
they were weighed, measured, and other data were collected. Then
the ones used for the parasite study were opened by making an
incision along the mid-ventral line of the body. The body cavity
worms and cysts were collected, as were the cysts from other parts of the body. These worms and cysts were then fixed and stored in vials of 70 per cent alcohol.

The heart, lungs, liver, kidneys, and gastro-intestinal tract were removed and wrapped in cheese cloth. After labeling they were placed in 10 per cent formalin, and brought back to Manhattan for examination in the laboratory.

The heart and kidneys were cut open and examined macroscopically for worms. In each case, the liver and lungs were divided into small pieces which were placed in a Petri dish containing water. The pieces then were teased apart and examined for worms with the aid of a dissecting microscope.

Each gastro-intestinal tract was divided into its component parts: stomach, small intestine, large intestine, and cecum. The parts were cut open longitudinally and the contents scraped into separate finger bowls. Running tap water was used to wash the parts of the digestive tract as they were being scraped. Generally a dissecting needle was used to open the tracts, as scissors would frequently damage the tapeworms.

Two methods were tried in separating the worms from the fine particles of chyme and feces. The first method was that of decanting the supernatant fluid after the heavier material and worms had settled to the bottom of a finger bowl. In several cases it was noticed that a few worms floated to the top of the dish, probably facilitated by air bubbles and fat particles. The second method tried was that of pouring the contents into an 80 mesh brass wire gauze and washing away the fine debris under tap water.
This second method seemed to be superior to the decantation method, so it was used for a majority of the work. Although no actual count was made of the worms lost in the two methods, none was found in the screening method after several visual checks. Very immature worms may have passed through the wire gauze but it is reasonably certain that few, if any, of the adults were lost.

Washed contents of the digestive tract were poured, in small quantities, into a Petri dish and examined for worms with the aid of a dissecting microscope. Early in this study every worm was counted; however, when large numbers of worms, principally pin-worms, were found it was impossible to obtain an accurate count. A dilution technique was used for approximately 100 jack rabbits to get an index of the total worm burden of the large intestine and cecum. The dilution method involved putting the appropriate contents into a 1000 ml. Erlenmeyer flask and adding water to a 650 ml. level. The flask was stoppered and shaken by rotation for approximately one minute. Then a 65 ml. jar was filled quickly from the flask and all the worms in this sample were counted. By multiplying the total number of worms found in the 65 ml. sample by ten, an index of the total number of worms was obtained. This count from the dilution method was used when one or more worms were found in the 65 ml. sample. If there were no worms in the 65 ml. sample, all the 650 ml. of material from the flask was examined and the actual number of worms was recorded. However, for the September, 1957, collection, if there were no worms found in
the 65 ml. sample, a record was made that less than ten worms were found. The reason for this change of procedure in the September collection was that after the above counting method was used for a year it seemed unnecessary to count the worms if there were less than ten per jack rabbit.

All worms collected were placed in vials of 70 per cent alcohol for future identification. Glycerine was used to clear all the nematodes. Temporary mounts were made of all the nematodes, except for the pinworms which were identified and stored in vials of glycerine. For temporary mounts, a cleared specimen was placed in a drop of glycerine on a slide. A cover slip, supported on each corner by a small portion of hand exercise material, Theraplast\(^1\), was then placed over the specimen. This type of mount prevented the worms from being crushed and also allowed the worms to be moved around for different views.

Larval cestodes, cysticerci and coenuri, were cleared in lacto-phenol for identification. Only the smaller coenuri and those in which the scoleces could not be seen, were cleared. All the cysticerci were cleared. After being cleared, all the specimens were stored in glycerine.

Adult and immature tapeworms were gradually dehydrated through the higher alcohols and then cleared in beechwood creosote. After identification, they were washed in xylol and gradually, decreasing percentages of alcohol were added until a concentration of 70 per cent alcohol was reached, in which they were stored. The

\(^1\) Thera-plast Co., 154 Nassau St., N. Y. C. 38.
vials were then dipped in hot paraffin to avoid evaporation.

RESULTS AND DISCUSSION

Jack rabbits in this study harbored no cattle or sheep parasites but closely related species of *Nematodirus* were found. No trematodes were found in this study.

No nematodes were recovered from the kidneys, lungs, and heart. A few specimens of *M. brevicauda* were found on the liver; in two instances, these worms were partially embedded in the liver tissue.

Table 1 shows the endoparasites found in the study and the number and per cent of jack rabbits infected.

Class Cestoda

**Family Taeniidae Ludwig, 1886. Taenia pisiformis** Bloch, 1780. There was a seven per cent incidence of larval tapeworms (cysticerci), *T. pisiformis*, in the jack rabbits examined (Table 1). The number of cysts recovered from each hare ranged from one to nineteen. The cysticerci were found on the mesenteries of the stomach, cecum, large intestine, lungs, liver, diaphragm, and adjacent to the spinal cord.

Cysticerci are bladder-type worms about the size of a large pea and having a single inverted holdfast organ. They are surrounded by a membrane, called an adventitious membrane by Wardle and McCleod (1952), which is produced by the host reaction to the parasite.

It was found that the cysts were generally singly attached
Table 1. Helminths recovered from 130 jack rabbits collected in Kansas

<table>
<thead>
<tr>
<th>Helminths</th>
<th>No. Infected</th>
<th>Per cent Infected</th>
</tr>
</thead>
<tbody>
<tr>
<td>Passalurus nonanulatus</td>
<td>97</td>
<td>75</td>
</tr>
<tr>
<td>Raillietina spp. (and Tapeworm spp.)</td>
<td>89</td>
<td>66</td>
</tr>
<tr>
<td>Micipsella brevicauda</td>
<td>65</td>
<td>50</td>
</tr>
<tr>
<td>Multiceps spp. (coenuri)</td>
<td>25</td>
<td>19</td>
</tr>
<tr>
<td>Nematodirus spp.</td>
<td>22</td>
<td>17</td>
</tr>
<tr>
<td>Taenia pisiformis (cysticerci)</td>
<td>9</td>
<td>7</td>
</tr>
<tr>
<td>Dermatoxys veligera</td>
<td>8</td>
<td>6</td>
</tr>
<tr>
<td>Nematodirus arizonensis</td>
<td>6</td>
<td>5</td>
</tr>
<tr>
<td>Physaloptera spp.</td>
<td>3</td>
<td>2</td>
</tr>
<tr>
<td>Nematodirus leporis</td>
<td>2</td>
<td>1.5</td>
</tr>
</tbody>
</table>
but a few were connected together by the host reaction material. This particular parasite has probably been studied by more workers than any other lagomorph parasite.

**Multiceps** spp. (Plate XI, Figs. 4, 5.). According to Erickson (1944), larvae of *M. serialis* and *M. packii*, both occurring in lagomorphs, can not be distinguished from each other. Therefore, the present study refers to these coenuri as **Multiceps** spp.

Each of 25 jack rabbits, twenty-three adults and three juveniles, was infected with from one to thirteen coenuri of this tapeworm species (Table 1). This represented a 19 per cent incidence of infection. The September, 1956, collection had the highest per cent incidence, 40 per cent, of any collecting period.

Almost three-fourths of the 40 coenuri recovered were from various locations in the body cavity. The rest of the cysts were recovered from the musculature. The largest cyst measured almost three inches in diameter and was located in the musculature of the right hip of one hare (Plate XI, Fig. 4). In two cases, coenuri were discovered in the heart muscles with one being in the right ventricle (Plate XI, Fig. 5). There was one case of a coenurus in the lungs.

As in the case of cysticerci, a membrane resulting from host reaction is formed around coenuri cysts. Coenuri differ from cysticerci in that each cyst contains several inverted holdfast organs rather than one.

Evidently the thickness of the adventitious membrane around each coenurus varies with the location of the coenurus in the
It was noticed in this study that when a cyst was loosely attached in the body cavity, the membrane was very thin or entirely absent. When a coenurus was located in the musculature the adventitious membrane was prominent. It appeared that the thicker membrane was made up of more layers of tissue than the thinner membrane. No doubt when a coenurus is in contact with the tissue of the host it causes more host reaction to be set up.

None of the hares seemed to be hindered by these larval tapeworms.

Family Davaineidae Fuhrmann, 1907. Raillietina spp. Eighty-five jack rabbits had an infection of only Raillietina spp. (Table 1). One animal had both Raillietina spp. and Tapeworm spp. while three animals harbored only Tapeworm spp. (Table 1). Probably the Tapeworm spp., a total of seven worms, were also Raillietina spp. but they were too deteriorated to make a positive identification. For purposes of convenience, Tapeworm spp. were figured in with Raillietina spp. for calculating per cent incidence and other statistical data.

Raillietina spp. have armed scoleces and single-pored proglottids which distinguish them from species of the genus Citto-taenia which have unarmed scoleces and double-pored proglottids. A combination of checking for armed scoleces and single-pored proglottids was necessary in a few cases, but an examination of the scoleces was generally all that was necessary for identification. Whenever the scolex was missing but the rest of the worm was present, such characteristics as single-pored proglottids,
egg capsules and other characteristics were used for identification. Some of the immature forms with missing scoleces had to be placed in Tapeworm spp.

The four suckers and retractable rostellum on the scolex were armed with tiny hooks, characteristic of the Davaineidae family. Hooks on the rostellum were generally more difficult to see than those on the suckers, because the rostellum was usually retracted. The rostellum was difficult to see unless the characteristic hooks, which marked its presence, were visible. The hooks on the suckers could not be seen in a few cases, probably because they were lost in handling the worms. In the immature worms, the characteristics of the scoleces appeared to be more easily seen than on adult worms.

The number of infected and noninfected adult and juvenile jack rabbits was calculated (Plate I) for each of the nine collecting periods, as was the total per cent incidence (Plate II). For example, in September, 1956, ten animals, six adults and four juveniles, were collected (Plate I). Five of six adults and all of four juveniles were infected with Raillietina spp. In December no adults were infected of eight collected and five juveniles were infected of twelve collected. For the February collection, only four of sixteen adults and two of four juveniles were infected. In April, the number of infected animals increased even though only adults were collected. During the rest of the study, both adults and juveniles were collected and the ratio between infected and noninfected adults and juveniles was fairly constant. No
EXPLANATION OF PLATE I

Number of adult and juvenile jack rabbits that were either infected or noninfected with Raillietina spp. (and TAPEWORM spp.) in each of the nine collecting periods. The black rectangular blocks below the abscissa represent the months when no jack rabbits were collected.
NUMBER OF INFECTED AND NONINFECTED ADULTS AND JUVENILES

RAILLETING spp. and TAPEWORM spp.

- ADULTS INFECTED
- JUVENILES INFECTED
- NONINFECTED

<table>
<thead>
<tr>
<th>Month</th>
<th>Infected</th>
<th>Noninfected</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sep</td>
<td>6</td>
<td>1</td>
</tr>
<tr>
<td>Oct</td>
<td>5</td>
<td>2</td>
</tr>
<tr>
<td>Nov</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td>Dec</td>
<td>8</td>
<td>2</td>
</tr>
<tr>
<td>Jan</td>
<td>7</td>
<td>1</td>
</tr>
<tr>
<td>Feb</td>
<td>6</td>
<td>1</td>
</tr>
<tr>
<td>Mar</td>
<td>5</td>
<td>2</td>
</tr>
<tr>
<td>Apr</td>
<td>13</td>
<td>3</td>
</tr>
<tr>
<td>May</td>
<td>12</td>
<td>1</td>
</tr>
<tr>
<td>Jun</td>
<td>10</td>
<td>2</td>
</tr>
<tr>
<td>Jul</td>
<td>8</td>
<td>2</td>
</tr>
<tr>
<td>Aug</td>
<td>6</td>
<td>1</td>
</tr>
<tr>
<td>Sep</td>
<td>5</td>
<td>2</td>
</tr>
</tbody>
</table>

PLATE I
EXPLANATION OF PLATE II

The total per cent of jack rabbits infected with Raillietina spp. (and TAPEWORM spp.) and Passalurus nonanulatus in each of the nine collecting periods. The black rectangular blocks below the abscissa represent the months when no jack rabbits were collected.
significant difference was found between adults and juveniles, with respect to degree of infection, in a Contingency $X^2$ Test (Table 3). A significant difference was found between infections during the winter months, December and February, and infections during the other months of the study (Table 3). The lowest per cent incidence was during the winter months of December and February (Plate II). All the other collecting periods showed a consistently higher per cent incidence than the winter collections. An attempt to explain why the winter months had low infections will be made later on in this discussion.

Each month's collection was compared as to infected and non-infected animals and a significant difference was found between months (Table 3). That is, the ratio of infected and noninfected animals was not the same in every monthly collecting period. However, the probability may have been different if a larger number of animals had been taken in each collecting period. There was no significant difference between infected males and females (Table 3).

Specific separation of the tapeworms into adult and immature groups was not done. However, several tapeworms were identified as adults by the presence of gravid proglottids, by size, and other characteristics. They were arbitrarily divided into worms with several segments, probably adults because of size when compared with known adult worms, and into worms with few segments, probably immature worms. The above separation of worms is not considered completely accurate, especially as to adult worms, because it was made mainly by size. Also it is not certain if all the worms were
the same species of *Raillietina*. However, the placing of worms with few segments into the immature group is probably accurate, although some of them may have been only pieces of adult worms.

During December, only immature worms were found. During February, six of fourteen jack rabbits were infected with tapeworms and all but one of the infected animals had immature tapeworms. The so-called immature worms may have been partially developed tapeworms or they may have been adult tapeworms that had lost most of their proglottids.

An explanation for the decrease of tapeworms during the winter months is difficult to make. If November and March collections had been made, the sharp decrease in infected animals between October and December, and the sharp rise in infected animals between December and February, may not have been so great. Honess (1935), in his study on the Black Hills Cottontail (*Sylvilagus nuttalli grangeri*) in Wyoming, found young worms of *Raillietina retractilis* in May, June, July, August, September, and December. While his work shows some seasonal differences in comparison with *Raillietina* spp. in this Kansas study, it is interesting to note some of the similarities. He mentions that the highest infection rate was in the summer months and only one infected animal was found from September through May. In other words his work is similar to this study in that there was a decrease in infections in the winter months and a rise in infections in the summer months. He found no infected animals from December through April but he collected fewer animals than in the Kansas study.
It is possible that the intermediate host, which has not been discovered for species of Raillietina in lagomorphs, may not be present during the winter months and that it reappears sometime between February and April in Kansas. More work certainly needs to be done on the life cycle of species of Raillietina in lagomorphs.

The following are the median numbers of worms in infected jack rabbits for each collecting period: September (2), October (2), December (2), February (1), April (3), June (9), July (7), August (3), and September (2). Probably the rise in number of worms in June and July is due to abundance of the intermediate host and to the susceptibility of the juveniles to higher numbers of worms at this time of year.

While the hares were divided into only adults and juveniles, the younger juveniles collected in the summer were more heavily infected with more fully developed tapeworms than were the older animals. Most work on lagomorph cestodes has been done on species of Cittotaenia in which age resistance was found to be a factor in limiting the incidence of this genus among adult animals. With Raillietina spp. this factor is not as evident, since there was no significant difference between adults and juveniles infected. If age resistance to Raillietina spp. is present, it probably is a matter of degree, as infected adult jack rabbits in this study had few mature worms.

Class Nematoda

Family Oxyuridae Cobbold, 1864. Passalurus nonanulatus
Skinker, 1931 (Plate IX, Figs. 1 - 3). Skinker in 1931 created a new species, *P. nonanulatus*, which is similar to *P. ambiguus* (Rudolphi, 1819) Dujardin, 1845. Mature female *P. ambiguus* worms have cuticular rings on their tails and males have more than two pairs of cloacal papillae. Female *P. nonanulatus* worms lack cuticular rings on their tails and the males have only two pairs of cloacal papillae. After a thorough study of the type specimens of *P. nonanulatus*, so kindly loaned by Mr. McIntosh, from the United States National Museum Helminthological Collection, the author found that the pinworms from the Kansas jack rabbits fit the descriptions of *P. nonanulatus* more closely than they did *P. ambiguus*.

Approximately 300 adult gravid female worms examined had no cuticular rings on their tails. Six males were examined in ventral cloacal view and only two pairs of cloacal papillae were seen, one pair of large preanal papillae and one pair of large postanal papillae. Also a pair of small "projections" was found just posterior to the cloaca. Immature worms, especially females, were difficult to identify to species level because the specific characteristics are based on the adult worms. As all adult worms examined were classified as *P. nonanulatus*, the immature worms were considered to be the same species.

Boughton (1932), McClure (1934), Ward (1934), Philip (1938), Erickson (1941, 1947), Bravo Hollis (1950), and Grundmann (1957) have all reported *P. nonanulatus* from lagomorphs. Rozycki (1941) found *P. ambiguus* from *Sylvilagus floridanus*. He writes that the
female worms examined had no prominent annulations and male worms had seven perianal papillae. The seven perianal papillae were distributed as follows: one pair of large preanal papillae, one pair of small papillae close together just behind the cloacal opening, one pair of large postanal papillae lateral to the small pair, and a single medium sized papilla just behind the small pair of papillae. Other authors have listed varying numbers of anal papillae in *P. ambiguus* males.

A ventral cloacal view of a male specimen from Skinker's collection was examined and a small pair of "projections" was noticed just posterior to the cloacal opening, besides the two large pairs of preanal and postanal papillae reported. These "projections", not mentioned by Skinker in her description, were also noticed in males in this Kansas study. The drawing that Rozycki made of *P. ambiguus*, a ventral cloacal view of a male, shows the same "projections" but he calls them papillae. So the only difference noted between his reported worms and those of Skinker and the ones in this study, is the presence of a single medium sized papilla behind the small pair of projections or papillae. Rozycki implies that *P. nonanulatus* may be a synonym of *P. ambiguus*. Either the worms he described are almost identical to *P. nonanulatus* or he gave a description of *P. ambiguus* that should cause *P. nonanulatus* to be put in synonymy. More work certainly needs to be done on these worms.

Seventy-five per cent of the jack rabbits collected were infected with this pinworm (Table 1). The cecum and large intestine were the most common locations for this parasite. Occasionally
they were found in the small intestine and stomach, however, these may have been ingested during coprophagy (Erickson, 1944).

As mentioned in the materials and methods, a dilution technique was used for counting most of the pinworms and the number recorded was an index and not an actual count. There was a median of 178 worms and a mean of 2271 worms per hare. One animal harbored approximately 48,000 worms, and another hare had approximately 25,000 worms.

For each of the nine collecting periods, the total per cent incidence (Plate II) and the number of infected and noninfected adults and juveniles were calculated (Plate III). A significant difference was found between the number of infected adult and juvenile jack rabbits (Table 3). Since more adults than juveniles were infected, data were tested to see if there was a higher per cent incidence of infection when the lowest number of juveniles was present in the collections. When collections made in the months of February, April, and June were compared with collections of other months, a significantly higher per cent incidence of *P. nonanulatus* was found when there were low numbers of juvenile and high numbers of adult jack rabbits (Table 3).

Each month's collection was compared as to infected and noninfected animals and a significant difference was found between months (Table 3). That is, the ratio of infected and noninfected animals was not the same in every monthly collecting period. However, the probability may have been different if a larger number of animals had been taken in each collecting period. There was no significant difference between infected males and females (Table 3).
EXPLANATION OF PLATE III

Number of adult and juvenile jack rabbits that were either infected or noninfected with Passalurus nonanulatus in each of the nine collecting periods. The black rectangular blocks below the abscissa represent the months when no jack rabbits were collected.
PLATE III

Passalurus nonanulatus

<table>
<thead>
<tr>
<th></th>
<th>Adults Infected</th>
<th>Juveniles Infected</th>
<th>Noninfected</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sep 1956</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Oct 1956</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Dec</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Feb</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Apr 1957</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Jun</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Jul</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Aug</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sep</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Number of Infected and Noninfected Adults and Juveniles

No Juveniles Collected
As mentioned earlier, it was found that there was a higher per cent incidence of worms when the juvenile population was low and the adult population was high, during February, April, and June. Evans (1940) in his work on *Oryctolagus cuniculus* in Wales found *P. ambiguus*. He reported that when there was an increase of numbers of young rabbits, there was a decrease in infection rate of the population.

Evidently the life cycle of *P. nonanulatus*, which probably is direct, is longer than for *Raillietina* spp. When juvenile jack rabbits increased, the tapeworm incidence increased but the pinworm incidence decreased (Plate II). Evans mentioned an age resistance with *P. ambiguus* but this was not evident with *P. nonanulatus*. No pathological effects were noticed with high numbers of worms in jack rabbits in Kansas.

*Dermatoxys veligera* (Rudolphi, 1819) Schneider, 1866 (Plate IX, Figs. 4 - 5) (Plate X, Figs. 1 - 2). Six per cent of the jack rabbits were infected with *D. veligera* (Table 1). The incidence of infected animals may have been higher since the worms were present in such low numbers that they may not have been found by the dilution method used. Whenever specimens of *D. veligera* were found in the dilution method, all appropriate contents were examined with the aid of a dissecting microscope. This worm was found in the large intestine of seven of eight animals and the cecum of one animal. Only light infections, from one to two worms, were found in the infected animals.

_Family Physalopteridae Leiper, 1908._ *Physaloptera* spp.
(Plate X, Figs. 3 - 4). Morgan and Hawkins (1949) list mink, fox, dog, cat, swine, wolf, raccoon, and bobcat as hosts for species of Physaloptera. Physaloptera spp. have been found by D. E. Worley in the striped skunk (Mephitis mephitis) and by the author in the badger (Taxidea taxus) in the Lakin area.

Three adult jack rabbits, two per cent, harbored eight immature female Physaloptera spp. (Table 1). One animal had six worms and the other two animals had one worm each. The jack rabbit harboring six worms had two of these in the small intestine. All other worms were found in the stomachs of the infected hosts. Rozycki (1941), Bell and Chalgren (1943), and Erickson (1947) reported Physaloptera spp. in Sylvilagus floridanus. Rozycki writes that the worms he found were immature and the worms found in this Kansas study were immature. The other two investigators make no mention of age of the worms. Evidently this parasite is an incidental parasite that does not develop to maturity in lagomorphs.

Family Trichostrongylidae Leiper, 1912. Identification of species of Nematodirus was made mainly by using the characteristics of the bursa and spicules of the male worm. Whenever only female worms were found in an animal, they were classified as Nematodirus spp. In no case were there more than one species of male Nematodirus found in any single animal. So, whenever both male and female worms were found in a jack rabbit, all the female worms were considered to be the same species of Nematodirus as the male worms. Measurements of the female worms are so similar in
species of *Nematodirus* that it is difficult, if not impossible, to separate them when only female worms are present.

Twenty-seven jack rabbits, 21 per cent, were infected with species of *Nematodirus*. All the infected hares were adults except for one infected juvenile animal. Worms were found in every collecting period except for July, August, and September, 1957.

*Nematodirus leporis* Chandler, 1924 (Plate XI, Fig. 3). Two adult jack rabbits were infected with *N. leporis* (Table 1). One hare had one worm and the other had 18 worms. Rozycki (1941) and Bell and Chalgren (1943) reported this worm from species of *Sylvilagus* in Kansas.

*Nematodirus arizonensis* Dikmans, 1937 (Plate XI, Figs. 1, 2). Six adult jack rabbits were infected with *N. arizonensis*; there were one to 52 worms in each animal. Rozycki, and Bell and Chalgren also reported this worm from species of *Sylvilagus* in Kansas. According to Travassos (1937), *N. arizonensis* is a synonym of *N. triangularis* Boughton, 1937.

*Nematodirus* spp. (Plate X, Fig. 5). Immature worms, pieces of worms, and female worms with no accompanying males, were classified as *Nematodirus* spp. Twenty-two jack rabbits, 21 adults and one juvenile, harbored these worms. Probably most of these worms were *N. arizonensis*.

**Family Dipetalonematidae** Wehr, 1935. *Micipsella brevicauda* n. sp. (Plate VI, Figs. 1 - 5; Plate VII, Figs. 1 - 5; Plate VIII, Figs. 1 - 5). Sixty-five jack rabbits, 50 per cent, were infected with from one to fourteen filariid worms (Table 1). These worms,
located in the abdominal cavity, belong in the genus *Micipsella* Seurat, 1921, by using the key to the Filaroidea by Chaubaud and Choquet (1953). This is the first known report of this genus from the United States.

Dipetalonematidae Wehr, 1935; Splendidofilarininae Chaubaud and Choquet, 1953; genus *Micipsella* Seurat, 1921. Body filariform and gradually tapering at both ends which are digitiform and rounded; caudal end of the male often coiled. Mouth on summit of a hemispherical cap which carries a circle of small papillae; there are four submedian papillae a little farther back.\(^\text{1}\) Cuticle thick and smooth, ornamented with small slightly projecting bosses arranged in two zigzag rows along the lateral lines which are broad and very conspicuous. Esophagus of uniform width and not divided. Microfilariae unsheathed (in utero).

**Male**: Total length 22 to 58 mm. (36 mm.\(^\text{2}\)) and maximum width 0.192 to 0.240 mm. (0.216 mm.). Width at anus 0.072 to 0.096 mm. (0.084 mm.), at nerve ring 0.067 to 0.108 mm. (0.084 mm.). Tail length 0.108 to 0.184 mm. (0.144 mm.). Nerve ring 0.156 to 0.228 mm. (0.192 mm.) from anterior end of body. Length of esophagus 0.648 to 0.984 mm. (0.840 mm.). Spicules are subequal and dissimilar. The larger left spicule 0.100 to 0.120 mm. (0.108 mm.) long; the shorter right spicule 0.067 to 0.096 mm. (0.084 mm.)

\(^{1}\) These papillae were not seen in *M. brevicauda* but little work was done on en fossa views.

\(^{2}\) Numbers in parentheses are the median lengths.
long. Both spicules curved and measurements taken approximately through the center of the spicules to include the curvature. Pre-anal papillae present, four to six pairs. Postanal papillae absent. Caudal end generally coiled.

**Female:**
- Total length $\frac{1}{4}$ to 106 mm. (65 mm.) and maximum width 0.26$\frac{1}{4}$ to 0.408 mm. (0.300 mm.).
- Width at anus 0.072 to 0.110 mm. (0.08$\frac{1}{4}$ mm.), at nerve ring 0.076 to 0.120 mm. (0.096 mm.), at vulva 0.096 to 0.192 mm. (0.120 mm.).
- Tail length 0.168 to 0.520 mm. (0.223 mm.).
- Nerve ring 0.108 to 0.240 mm. (0.192 mm.) and vulva 0.32$\frac{1}{4}$ to 1.200 mm. (0.48$\frac{1}{4}$ mm.) from anterior end.
- Length of esophagus 0.720 to 1.032 mm. (0.86$\frac{1}{4}$ mm.).
- Opisthodelphic, viviparous. Microfilariae unsheathed (in utero).

**Host:** *Lepus californicus melanotis* Mearns

**Location:** Abdominal cavity

**Locality:** Kearny County, Kansas

Seurat (1917) described filarial worms from the abdominal cavity of *Lepus pallidior* and *Lepus kabylicus* from Algeria and named them *Filaria numidica*. In 1921 Seurat created a new genus, *Micipsella*, for this species. Kalantarian (1924) described similar worms from *Lepus* spp. in Armenia and created a new genus to receive Seurat's existing species. Kalantarian called this worm *Cercofilaria numidica*, evidently not knowing that Seurat had transferred *numidica* from the genus *Filaria* to the genus *Micipsella*. Yorke and Maplestone (1926) placed *Cercofilaria numidica*

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1 Now considered a subspecies of *L. capensis* (All information on classification of *L. capensis* in this thesis was kindly provided by Dr. H. W. Setzer, Smithsonian Institution, U. S. N. M.).
in synonymy with *Micipsella numidica*. Rao (1938) described filarial worms from a blood clot of the portal vein of *L. nigricollis* in India and named them *Micipsella indica*. Ivashkin (1954) found filarial worms in the portal vein of *L. tolaei* from Mongolia which he classified as *Micipsella numidica*. These worms differ somewhat from Seurat's (1917) original description in dimensions and number of anal papillae. According to Ivashkin (1954), *M. numidica* has also been found in *L. lehmani* and *Lepus* spp. in Russia.

*M. brevicauda* differs mainly from *M. numidica* and *M. indica* in the male characteristics; the tail is much shorter and has no postanal papillae. In one male, an "end on" view of the tail, examined under oil immersion, revealed two tiny papillate structures but this feature was not seen in several other worms examined. The tiny structures mentioned are the only structures that might be called postanal papillae. In other descriptions of postanal papillae in species of *Micipsella*, no mention was made about the size of the preanal papillae. However, the drawings by Seurat, Ivashkin, and Rao (1938) show the postanal papillae to be about the same size as the preanal papillae and none are located on the tip of the tail.

Table 2 shows a comparison of *M. numidica* as described by Seurat and Ivashkin, of *M. indica* by Rao (1938) and of *M. brevicauda* in this thesis. Rao gives only a few measurements for *M. indica* and the characteristics given seem to be very similar to

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1 Now considered as a subspecies of *L. capensis*. 
Table 2. Comparison of species of *Micipsella.*
(all measurements in millimeters)

<table>
<thead>
<tr>
<th></th>
<th><em>M. numidica</em></th>
<th></th>
<th><em>M. indica</em></th>
<th></th>
<th><em>M. brevicauda</em></th>
</tr>
</thead>
<tbody>
<tr>
<td>Male: Female</td>
<td>Male: Female</td>
<td>Male: Female</td>
<td>Male: Female</td>
<td></td>
<td>Male: Female: No. Meas'd</td>
</tr>
<tr>
<td>Total length</td>
<td>76</td>
<td>130</td>
<td>55-61</td>
<td>60-93</td>
<td>70-100: 120-140 (22-59)</td>
</tr>
<tr>
<td></td>
<td>10</td>
<td>(26.1-40.8)</td>
<td>58</td>
<td>(720-1,032)</td>
<td>71</td>
</tr>
<tr>
<td>Maximum width</td>
<td>0.420</td>
<td>0.540</td>
<td>0.736</td>
<td>0.699</td>
<td>0.480</td>
</tr>
<tr>
<td></td>
<td>(0.192-0.240)</td>
<td>(0.216-0.240)</td>
<td>(0.648-0.904)</td>
<td></td>
<td>(0.480-0.500)</td>
</tr>
<tr>
<td>Esophagus</td>
<td>0.970</td>
<td>0.700</td>
<td>0.499</td>
<td>0.745</td>
<td>0.480</td>
</tr>
<tr>
<td></td>
<td>(0.648-0.904)</td>
<td></td>
<td></td>
<td></td>
<td>(0.480-0.500)</td>
</tr>
<tr>
<td>Vulva from</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>anterior end</td>
<td>--</td>
<td>0.720</td>
<td>--</td>
<td>0.782</td>
<td>(0.324-1.200)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(0.648-0.904)</td>
<td></td>
<td></td>
<td>(1.64-2.240)</td>
</tr>
<tr>
<td>Tail length</td>
<td>0.420</td>
<td>0.480</td>
<td>0.299</td>
<td>0.234</td>
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</tr>
<tr>
<td></td>
<td>(0.392**-0.392**)</td>
<td></td>
<td></td>
<td></td>
<td>(0.108-0.184)</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>(0.223-0.223)</td>
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<tr>
<td>Spicules</td>
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<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Left</td>
<td>0.105</td>
<td>--</td>
<td>0.135</td>
<td>--</td>
<td></td>
</tr>
<tr>
<td></td>
<td>(0.100-0.120)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Right</td>
<td>0.085</td>
<td>--</td>
<td>0.095</td>
<td>--</td>
<td></td>
</tr>
<tr>
<td></td>
<td>(0.085-0.095)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Width at vulva</td>
<td>--</td>
<td>--</td>
<td>--</td>
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<td></td>
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<tr>
<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Width at anus</td>
<td>--</td>
<td>--</td>
<td>--</td>
<td>--</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nerve ring</td>
<td>0.240</td>
<td>0.240</td>
<td>--</td>
<td>--</td>
<td></td>
</tr>
<tr>
<td>from anterior</td>
<td>(0.156-0.228)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>end</td>
<td>(0.108-0.240)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>width at</td>
<td>--</td>
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<tr>
<td></td>
<td></td>
<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Preanal papillae</td>
<td>5-7 pr.</td>
<td>--</td>
<td>8 pr.</td>
<td>--</td>
<td></td>
</tr>
<tr>
<td></td>
<td>(6-7 pr.)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Postanal papillae</td>
<td>2 pr.</td>
<td>--</td>
<td>4 pr.</td>
<td>--</td>
<td></td>
</tr>
<tr>
<td></td>
<td>(3 pr.)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Location in host</td>
<td>abdominal cavity</td>
<td>portal vein of liver</td>
<td>portal vein of liver</td>
<td>abdomenal cavity</td>
<td></td>
</tr>
<tr>
<td>Host</td>
<td><em>L. pallidior</em></td>
<td><em>L. tola</em></td>
<td><em>L. nigricollis</em></td>
<td><em>L. californicus melanotis</em></td>
<td></td>
</tr>
<tr>
<td>Geographic</td>
<td>Algeria</td>
<td>Mongolia</td>
<td>India</td>
<td>U. S. A. (Kansas)</td>
<td></td>
</tr>
<tr>
<td>distribution</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* Now considered subspecies of *L. capensis.*

** Calculated from his scaled drawing.

*** Range in parentheses, median recorded below range.
**M. numidica** except that the spicules may be a little different in shape. Ivashkin shows that there is some variation in **M. numidica**, especially in the tail length of both sexes and in the numbers of preanal and postanal papillae of the male. The male of **M. brevicauda** has a tail length of 0.108 to 0.184 mm. (0.144 mm.) with four to six pairs of preanal papillae and no postanal papillae. Ivashkin gives the male tail length for **M. numidica** as 0.299 mm. with five to seven pairs of preanal papillae and two pairs of postanal papillae. Seurat gives it as 0.420 mm. with eight pairs of preanal papillae and four pairs of postanal papillae. Rao gives no male tail measurements for **M. indica** but by measuring his scaled drawing it is approximately 0.392 mm. long and has six to seven pairs of preanal papillae and three pairs of postanal papillae.

The shape of the spicules of males of **M. brevicauda** is more easily seen than described (Plate VI, Fig. 2; Plate VII, Fig. 2). The shorter spicule appears to be bluntly pointed at its tip. In some specimens the tip is more pointed than in others. The tip of the longer spicules has a bifurcated appearance. According to Hyman (1951) the left spicule is generally longer than the shorter right spicule in nematodes. However, in several of the male worms examined in this study the longer spicule seemed to be on the right side and the short spicule on the left side. A little more than half the worms had the long spicule on the left side and the short one on the right side. Since most of the worms were flattened out on the slide to allow examination with an oil immersion objective, the spicules may have been displaced from their
normal positions. Other authors in describing species of Micipsella mentioned that the left spicule was longer than the right one. It is assumed in this present study that the left one is longer also.

*M. brevicauda* females have measurements that vary slightly from the other described species of the genus. The distance of the vulva from the anterior end is quite variable but the median distance is approximately half way between the anterior end of the body and the junction of the esophagus and the intestine. Both sexes appear to be narrower in width than the other species. This measurement probably should not be used in describing these worms as the width depended upon how flattened out they were by the pressing down of the cover slip. Total lengths of the worms, especially females, showed quite a range in length, 44 to 106 mm. The smallest female having microfilariae in utero was 44 mm. long. Only females with microfilariae in utero were used for measurements. Several female worms longer than 44 mm. and lacking microfilariae were found but were not measured although they probably were adults. The smallest female found was 18 mm. long and it was definitely immature. Wide ranges in total length seem to be common in filarids.

Distinguishing adult male from immature male worms was more difficult than the arbitrary standard of using the presence of microfilariae in females as a means for classifying them as adults. All the males found were considered adult, as even the smallest one, 22 mm., had measurements similar to larger males and also had
germinal material in the reproductive system.

In both sexes the cephalic, hemispheric cap was visible (Plate VIII, Fig. 1). A regression of this cap was evident in some of the larger worms, particularly in the females. That is, the smaller worms had a cap that was generally easily seen, but it was hardly visible in some of the larger worms.

Bosses in two zigzag rows on the lateral lines, characteristic of this genus, are found in *M. brevicauda*. The name for these bosses or minute projections seem to vary with different authors. Rao (1938) calls them both "papillae" and "bosses" in his paper. Seurat (1917) uses "verrues" translated to mean "warts" to describe these structures. Yorke and Maplestone (1926) call them "bosses" in the description of *M. numidica*. *M. brevicauda* has two zigzag rows of very minute projections along the wide lateral lines which no doubt are "bosses" (Plate VII, Figs. 4, 5; Plate VIII, Fig. 2). Seurat mentions that these same "bosses" are on the dorsal and ventral surfaces of the male tail of *M. numidica*; these are not present in *M. brevicauda*.

There seems to be no predisposition of this filarial worm for a particular location in the abdominal cavity of the hare. Worms were recovered from the body cavity walls and from almost all of the mesenteries. They were also found on the surface of the small intestine, large intestine, cecum, and liver. Worms were found partially embedded in the liver in two different instances. In several cases, adult worms were found partially encased in yellowish, fatty-like cysts. In two or three cases, worms were completely enveloped by the cysts. Many yellow cysts
were dissected apart in which no worms were found. It may be that these cysts, which were attached by small stalks in several cases, are involved in the life cycle.

As with the tapeworms and pinworms, a Contingency $X^2$ Test was made on statistical data collected. For each of the nine collecting periods the total per cent incidence (Plate IV) and number of infected and noninfected adults and juveniles was calculated (Plate V). The number of infected animals may have been higher because the body cavity and external surface of the digestive tract were not examined closely for worms during September and October, 1956.

A significant difference was found between infected adults and juveniles (Table 3). During June and July no juveniles were infected and there was a gradual increase of infected juveniles from August through September (Plate IV). Probably the low number of infected juveniles during these months was due to the young age of the juveniles. Evidently the filariid worms take several weeks to mature.

Each month's collection was compared as to infected and noninfected animals and a significant difference was found between months (Table 3). That is, the ratio of infected and noninfected animals was not the same in every monthly collecting period. However, the probability may have been different if a larger number of animals had been taken in each collecting period.

The animals collected from September, 1956, through April, 1957, had the highest per cent incidence, December being the
EXPLANATION OF PLATE IV

The total per cent of jack rabbits infected with *Micipsella brevicauda* in each of the nine collecting periods. The black rectangular blocks below the abscissa represent the months when no jack rabbits were collected.
EXPLANATION OF PLATE V

Number of adult and juvenile jack rabbits that were either infected or noninfected with *Micipsella brevicauda* in each of the nine collecting periods. The black rectangular blocks below the abscissa represent the months when no jack rabbits were collected.
NUMBER OF INFECTED AND NONINFECTED ADULTS AND JUVENILES

PLATE V

M. brevicauda
- ADULTS INFECTED
- JUVENILES INFECTED
- NONINFECTED

NO JUVENILES COLLECTED

MONTHS

1956

1957

W. theobaldi

ADULTS INFECTED

JUVENILES INFECTED

NONINFECTED
highest with 80 per cent (Plate IV). From June through August there was a gradual decrease in per cent incidence and a rise occurred again in September. The absence of infected adults during August is not readily explainable but it may have been due to sampling error. No significant difference was found between the number of infected males and females (Table 3).

With such a high incidence of this worm it seems unusual that it has not been reported before in this country. The intermediate host must be common in southwestern Kansas and work on the life cycle should be an interesting challenge. Evidently the life cycle has not been worked out for the other species of Micipsella.

It may be that the variations of M. brevicauda are only host differences and that this worm is the same as an already named species of Micipsella. The ideal procedure would be to compare the anatomical structures of all the known species of Micipsella and then to experimentally transmit them to several lagomorph species and closely related animals to see if variations can be influenced by the hosts. Of course a more practical method would be to experiment with just M. brevicauda and see if there are any variations in the worms. With the present basis for classification, it appears that M. brevicauda must be considered a new species.

Ecology. From September, 1956, through April, 1957, most of the jack rabbits used in this study were collected southeast of Lakin, between the Arkansas River and the sand-hills area. This
was a drought period and food being scarce in the sand-hills, the animals concentrated on the periphery of the sand-hills to feed on adjacent cropland at night (Bronson, 1957). According to Bronson, most of the animals dispersed into the sand-hills after spring rains in 1957, helped to restore the vegetation there.

During July through September, 1957, most of the jack rabbits were collected north of Lakin. These animals concentrated in the draws in the daytime and spread out at night to feed on cropland. Irrigation water was used on cropland both north and south of Lakin.

To correlate the habitat of the host with that of its parasite burden is rather difficult. We have the problem of seasonal fluctuations of the parasite along with the drought period for the first part of the study and the wetter period in the latter part of the study.

No doubt, parasite fluctuations were caused by presence or absence of intermediate hosts, age resistance, and other factors such as concentration of jack rabbits during the drought and dispersion of them when wetter conditions prevailed. The animals normally concentrate to some extent in the winter months but during the drought period they also concentrated in the summer months.

It is probable that intermediate hosts are required for Raillietina spp., Physaloptera spp., and Micipsella brevicauda. Assuming these intermediate hosts to be arthropods, it is interesting to speculate on the greater chances of the concentrated jack rabbits becoming infected during the drought period.
If the intermediate hosts for the endoparasites are wingless, e.g. ticks, mites or fleas, their powers of actively seeking out a jack rabbit are limited. When the hares are concentrated there is more chance for these nonflying arthropods to contact their hosts. El-Rawi (1957) examined 64 jack rabbits from the Lakin area and found 456 ticks (two species) and 24 fleas (two species).

Winged arthropods, that can aggressively seek out an animal, could be involved in the life cycles. For example, mosquitoes may be the intermediate hosts for *M. brevicauda*. The chance for infection may have been higher when animals were concentrated because of the greater chance of the mosquitoes coming in contact with infected animals and then transmitting this filariid to non-infected animals.

The life cycle for the pinworms, *D. veligera* and *P. nonanulatus*, is unknown but is probably direct. Because of the coprophagous habit of lagomorphs (Erickson, 1944), a concentration of hares could increase the chances for infection with pinworms. Once these infections are picked up, retroinfection, if present, would serve to increase the numbers of worms in the hosts.

Species of *Nematodirus* are known to require moisture for completion of their life cycle (Boughton, 1932). During the drought, irrigation water could have supplied the moisture necessary for the eggs and infective stage larvae to develop. However, the only months that can be compared are September, 1956, and September, 1957. Infected jack rabbits were found in September, 1956, but none was found in September, 1957. It is challenging to explain
why no worms were found from July through September, 1957, when a wetter period existed.

Aside from drought conditions and concentration of the jack rabbits, other factors may determine the existence of the endoparasites. Winter temperatures may be a limiting factor for intermediate hosts. During cold weather the intermediate hosts may be completely absent or dormant. Thus, certain parasites requiring intermediate hosts may be present in low numbers or completely absent from their hosts during the winter months.

The age of the host may determine the presence or absence of a parasite. Specimens of *Raillietina* spp. that were more completely developed, were found in younger animals oftener than in older animals. However, the older hares seemed to have more immature worms indicating a possible degree of age resistance rather than a complete resistance to the tapeworms.

*P. nonanulatus, M. brevicauda,* and species of *Nematodirus* were found in more adult than juvenile animals. The length of time needed for maturity of these worms may be a more important factor than their needing older animals for the completion of their life cycle.

For the larval tapeworms, *Taenia* spp. and *Multiceps* spp., the availability of the definitive hosts, dogs and coyotes, is a limiting factor.

The internal and external environmental factors the endoparasite must contend with are complex. Many attempts have been made to explain why fluctuations of endoparasites occur. No
doubt there is no single limiting factor but several factors that
determine when a parasite can survive in a particular host.

SUMMARY

One hundred and thirty Black-tailed Jack Rabbits (Lepus
californicus melanotis Mearns) were examined for helminth para-
sites. The hares were collected near Lakin in Kearny County,
Kansas, during nine collecting periods from September, 1956,
through September, 1957. The heart, lungs, liver, kidneys, gastro-
intestinal tract, body cavity, and musculature of the animals were
examined for endoparasites.

Purposes of the study were: (1) collect jack rabbits through-
out the year to investigate seasonal fluctuations in incidence of
parasitism, (2) investigate whether the total number of each spe-
cies of helminth varied during the year, (3) determine if the
jack rabbits in this area acted as reservoirs for cattle or sheep
parasites, (4) observe if the endoparasites were associated with
mortalities among jack rabbits, and (5) present a clearer picture
of jack rabbit helminths in Kansas.

Following is the per cent incidence of each parasite found
in this study: Passalurus nonanulatus 75 per cent, Raillietina
spp. (and Tapeworm spp.) 66 per cent, Micipsella brevicauda 50
per cent, Multiceps spp. 19 per cent, Nematodirus spp. 17 per cent,
Taenia spp. 7 per cent, Dermatoxys veligera 6 per cent, Nematodirus
arizonensis 5 per cent, Physaloptera spp. 2 per cent, and Nematodi-
rus leporis 1.5 per cent.
P. nonanulatus and M. brevicauda showed the highest per cent incidence in the winter months. Raillietina spp. (and Tapeworm spp.) showed the highest per cent incidence in the summer months.

M. brevicauda n. sp., found in the abdominal cavity, is considered a new species mainly on two characteristics of the caudal end of the male worm. The length of the male tail is more than .100 mm. shorter than closely related species and no postanal papillae were found. Other species of Micipsella reportedly have longer tails in the male worms with two to four pairs of postanal papillae. Probably this is a new record of the genus Micipsella in the United States.

The characteristics of P. nonanulatus are so similar to those of P. ambiguus that separation of the two species is difficult. More work on these worms may reveal that P. nonanulatus is a synonym of P. ambiguus.

Age resistance, lack of intermediate hosts, and concentration of the hosts during a drought period and dispersion of the hosts during a wet period may have caused parasite fluctuations in jack rabbits in this study.

No cattle or sheep parasites were found. Also no pathological effects were noticed that might cause death in the hares. Under stress conditions, high numbers of worms may contribute to mortalities.
ACKNOWLEDGMENTS

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LITERATURE CITED

Ackert, J. E.

Arnold, J. G., Jr.

Baer, J. G.

Bell, J. F. and W. S. Chalgren.

Boughton, R. V.

Bravo Hollis, M.

Bronson, F. H.

Chabaud, A. G. and M. T. Choquet.

Chandler, A. C.

Curtice, C.

Dikmans, G.
Dikmans, G.

Douthitt, H.

El-Rawi, B. M.

Erickson, A. B.

Erickson, A. B.

Evans, W. M. R.
Observations on the incidence of some nematode parasites of the common rabbit, Oryctolagus cuniculus. Parasitol. 32: 67-77. 1940.

Grundmann, A., D. Parker and G. Stagg.

Grundmann, A. W.

Hall, M. C.

Honess, R. F.

Honess, R. F. and K. B. Winter.

Hyman, L. H.
Ivashkin, V. M.

Kalantarian, E. V.

Lechleitner, R. R.

Lyman, R. A.

McCandless, S. C.

McClure, G. W.

McGhee, F. J.

Meyer, M. C.

Morgan, B. B. and A. B. Hawkins.
Veterinary helminthology. Burgess: Minneapolis. 1949.

Olsen, C. W.
Wild rabbits as reservoir hosts of the common liver fluke, Fasciola hepatica, in southern Texas. J. Parasitol. 34(2): 119-123. 1948.

Palmer, T. S.

Philip, C. B.
Rao, M. A. N. 
*Micipsella indica*, n. sp. Indian J. Vet. Sc. and Animal 
Husb. 8(3):251-253. 1938.

Rohrbacher, G. H., Jr. and F. A. Ehrenford. 
*Bioastranema*, new genus (Nematoda: Trichostrongyliidae) from 
the California Jackrabbit, *Lepus californicus californicus* 

Rozycki, A. T. 
Studies on the intestinal nematodes of the cottontail rabbit. 

Schwartz, E. and W. E. Shook. 
1928.

Scott, J. W. 
A new lungworm from the Leporidae *Prostrongylus sylvilagii*, 

Seurat, L. B. 
Une nouvelle filaire peritoneale des rongeurs. Compt. Rend. 

Seurat, L. G. 
12:31-37. 1921.

Skidmore, L. V. 
*Trichostrongylus colubriformis (=T. instabilis)* in the jack-
Assoc. 80, n.s.v. 33(5):800-801. 1932.

Skinker, M. S. 

Stiles, C. W. 
A revision of the adult tapeworms of hares and rabbits. 

Travassos, L. 
Revisao da familia Trichostrongylidae Leiper, 1912. Monogr. 

The life histories and ecology of jackrabbits, *Lepus allenii* 
and *Lepus californicus* ssp., in relation to grazing in Arizona. 
553. 1933.
Ward, J. W.
A study of some parasites of rabbits of central Oklahoma.

Wardle, R. A. and J. A. McLeod.

Yorke, W. and P. A. Maplestone.
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<th>Table 3. Contingency $X^2$ Test for tapeworm, pinworm and filariid parasite burden in jack rabbits.</th>
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<tbody>
<tr>
<td><strong>Raillietina spp. (and Tapeworm spp.)</strong></td>
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<tr>
<td>Infected and noninfected adults vs. infected and noninfected juveniles</td>
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<tr>
<td>Winter months (Dec., Feb.) of infection vs. nonwinter months (other months) of infection</td>
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<tr>
<td>Infected animals vs. noninfected animals between months</td>
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<td>Infected and noninfected males vs. infected and noninfected females</td>
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<tr>
<td><strong>Passalurus nonanulatus</strong></td>
</tr>
<tr>
<td>Infected and noninfected adults vs. infected and noninfected juveniles</td>
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<tr>
<td>Months (Feb., Apr., Jun.) of low number of juveniles vs. months (other months) of higher number of juveniles</td>
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<tr>
<td><strong>Micipsella breviceuda</strong></td>
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EXPLANATION OF PLATE VI

Micipsella brevicauda

All drawings were made with the aid of a camera lucida.
All measurements are in millimeters and each scale shown equals 0.1 mm. The following abbreviations were used:

A ............... anus
B ............... bosses
E ............... esophagus
I ............... intestine
LS ............... left spicule
NR ............... nerve ring
PP ............... preanal papillae
RS ............... right spicule
T ............... testis
U ............... uterus
V ............... vulva
VA ............... vagina

Fig. 1. Anterior end of male
Fig. 2. Posterior end of male
Fig. 3. Lateral line of female showing bosses
Fig. 4. Posterior end of female
Fig. 5. Anterior end of female
EXPLANATION OF PLATE VII

**Micipsella brevicauda**

**Fig. 1.** Posterior end of male showing tail shape (approximately 100X)

**Fig. 2.** Posterior end of male showing spicules (approximately 500X)

**Fig. 3.** Posterior end of male showing spicules (approximately 100X)

**Fig. 4.** Portion of female showing the bosses on the lateral line (approximately 100X)

**Fig. 5.** Portion of female showing the bosses on the lateral line (approximately 970X)
PLATE VII
EXPLANATION OF PLATE VIII

Micipsella brevicauda

Fig. 1. Anterior end of female showing the hemispheric cap at the very end (approximately 450X)

Fig. 2. Portion of female showing bosses; the width of the lateral line can be seen also (approximately 970X)

Fig. 3. Posterior end of female (approximately 100X)

Fig. 4. Posterior end of female (approximately 450X)

Fig. 5. Anterior end of female; vulvar opening and vagina are at the point where the worm begins to taper towards the anterior end (approximately 100X)
EXPLANATION OF PLATE IX

Fig. 1. *P. nonanulatus*, posterior end of female (approximately 100X)

Fig. 2. *P. nonanulatus*, anterior end of female (approximately 100X)

Fig. 3. *P. nonanulatus*, posterior end of male (approximately 75X)

Fig. 4. *D. veligera*, anterior end of female (approximately 100X)

Fig. 5. *D. veligera*, showing posterior part of esophagus which is enlarged to form a bulb where it joins the intestine (approximately 100X)
EXPLANATION OF PLATE X

Fig. 1. *D. veligera*, posterior end of female; eggs can be seen in utero (approximately 100X)

Fig. 2. *D. veligera*, posterior end of male (approximately 100X)

Fig. 3. *Physaloptera* spp., anterior end of female (immature) (approximately 100X)

Fig. 4. *Physaloptera* spp., posterior end of female (immature) (approximately 100X)

Fig. 5. *Nematodirus* spp., portion of immature female showing the double ovejectors (approximately 100X)
EXPLANATION OF PLATE XI

Fig. 1. N. arizonensis, focused to show the triangular shape of the bursa and the distribution of the bursal rays of the male (approximately 100X)

Fig. 2. N. arizonensis, focused to show the typical shape of the spicule tip of the male (approximately 100X)

Fig. 3. N. leporis, focused to show the "hooked" appearance of the spicule tip of the male (approximately 100X)

Fig. 4. Multiceps spp., coenurus from the right hip of a Jack rabbit.

Fig. 5. Multiceps spp., heart with coenurus in the right ventricle. (Note arrow). The white "knobs" are scoleces.
HELMINTH PARASITES OF THE BLACK-TAILED JACK RABBIT (Lepus californicus melanotis Mearns) IN SOUTHWESTERN KANSAS

by

EUGENE THOMAS LYONS

B. S., South Dakota State College of Agriculture and Mechanic Arts, 1956

AN ABSTRACT OF A THESIS

submitted in partial fulfillment of the requirements for the degree

MASTER OF SCIENCE

Department of Zoology

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1958
A review of the literature revealed that several lagomorphs had been studied quite extensively for endoparasites. While several scattered reports of jack rabbit helminths had been made, a more complete study of endoparasites of jack rabbits seemed necessary in order to get a clearer picture of their parasitic fauna.

One hundred and thirty Black-tailed Jack Rabbits (Lepus californicus melanotis Mearns) were examined for helminth parasites. The hares were collected near Lakin in Kearny County, Kansas, during nine collecting periods from September, 1956, through September, 1957. The heart, lungs, liver, kidneys, gastro-intestinal tract, body cavity, and musculature of the animals were examined for endoparasites.

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