STUDIES ON THE BIONOMICS OF THE POWL GESTODE RAILLIPTINA CESTICILLUS (MOLIN)

by

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PART I. INTERMEDIATE HOSTS OF CHICKEN TAPEWORMS FOUND IN KANSAS

INTRODUCTION

The recent findings of Ackert and Case (1938) on effects of tapeworms on chickens have re-emphasized the importance of these parasites. These authors found that cestode infections of three weeks duration affected the blood of the chickens and that infections of eight weeks duration lowered the hemoglobin and sugar content of the blood and retarded the growth of the chickens. While fowl taeniasis can be checked momentarily by administering taeniacides, no effective control of these parasites can be established without breaking the life cycle at some stage. In order to do this, knowledge must be gained of the available intermediate hosts. To determine these, life history studies of various species of fowl cestodes have been conducted recently in several laboratories; notably at the United States Zoological Division, Washington, at the Veterinary Academy, Mannover, Germany, and at the Kansas State College of Agriculture and Applied Science, Manhattan.

Whereas, a decade ago, garden slugs, earthworms, and flies were thought to be the principal intermediate hosts of chicken tapeworms, the recent findings have shown that ground beetles (Carabidae) are doubtless the principal means of transmitting the more common chicken tapeworms. It was to make known more generally the newer knowledge of intermediate hosts of fowl tapeworms found in Kansas that this study was undertaken.

NORTH AMERICAN CHICKEN TAPEWORMS

Of ten species of tapeworms found in the United States (Cram, 1928), the following six species have been reported from Kansas:

Raillietina cesticillus (Molin, 1858)
Choanotaenia infundibulum (Bloch, 1779)
Raillietina tetragona (Molin, 1858)
Eymenolepis carioca (Magalhaes, 1898)
Raillietina echinobothrida (Megnin, 1880)
Amoebotaenia sphenoides (Railliet, 1892)

The other four species reported from this country are:

<u>Davainea proglottina</u> (Davaine), <u>Diorchis americana</u> Ransom,

<u>Hymenolepis cantaniana</u> (Pol.), and <u>Metroliasthes lucida</u>

(Ransom).

Kansas Fowl Costodes and Their Intermediate Hosts
Raillietina costicillus. The fowl costode that

probably is of the most common occurrence in Kensas is the rather large species, Raillistina cesticillus whose habitat is the duodenum. Ferry (1934) reported it from 62 percent of the chickens he examined. However, Adams and Geiser (1933) found it in only 12 percent of the chickens they exemined in Texas. Although houseflies can probably act as intermediate hosts (Ackert, 1918: Reid and Ackert, 1937) the most important intermediate hosts are various species of beetles (Coleoptera), especially the family Carabidae (ground beetles). Besides several genera of the families Scarabaeidae, Tenebrionidae, and Ostomidae, there are eight genera and 25 species of Carabidae which may serve quite readily as intermediate hosts. The beetles most frequently found at Manhattan (Riley County, Kansas) are of these species. In half an hour in midsummer (July) it has been possible to collect 175 Carabidae on the campus of Kansas State College. The most numerous of these Carabidae are species of the genera Amara and Cratacanthus. After a morning rain, beetles of these genera may be seen by the hundreds. Species of Cratacenthus and Amera have been found to be the best intermediate hosts for the cestode R. cesticillus as determined by experimentation. The ease with which mature cysticercoids can be developed in beetles has led to the use of R. cesticillus for experimental

studies of Ackert and Case, mentioned above, of Ackert and Reid (1957) who demonstrated an age resistance of older chickens to the growth of these cestodes, and of Harwood and Luttermoser (1958) who found that infections of this cestode retarded the growth of chickens.

Intermediate Hosts of R. costicillus. COLEOPTERA:
Family Cantheridae: Podabrus modestus Sey; Family
Carabidae: Amera sp., Amera besilleris (Say), Amera fellax
Lec., Amera (Curtonotus) laticollis Lec., Amera (Celis)
muscula Say, Amera (Percosia) obesa Say, Anaferonia
(Pterostichus) sp., Anaferonia constricta Say, Anaferonia
near substriata (Lec.), Anisotarsus sp., Anisotarsus agilis
(Dej.), Anisotarsus subvirens Csy., Chalenius tomentosus
Say, Cratacanthus dubius Beauv., Harpalus sp., Harpalus
faumus Say, Harpalus herbivarus Say, Harpalus pennsylvanicus
DeGeer, Pterostichus (Poscilus) chalcites Say,
Pterostichus near constricta, Pterostichus (Abacidus)
permundus Say, Pterostichus near permundus, Pterostichus
(Eumolops) torvus Lec., Selenophorus (pedicularius LeC.),
Triplectrus pusticus Say, (Jones, 1929).

Family Ostomidae: Tenebroides mauritanious (L.) (Case and Ackert, 1938); Family Scarabaeidae: <u>Choeridiva</u> historoides (Web.), <u>Aphodius granarius</u> (L.); Family Tenebrionidae: <u>Tenebrio</u> sp., <u>Tribolium confusum</u> Duval., Tribolium sp. Hymenolepis carioca. Second in the list of tapeworms from Kansas chickens appears to be Hymenolepis carioca, a delicate, thread-like cestode which may be present in large numbers (500 or more) in a single bird, according to Ferry (1934). Twiehaus (unpublished) likewise found large infections of this cestode in ailing chickens in Kansas that were sent to the Kansas State College Poultry Diseases laboratory in April, 1939. Adams and Geiser (1935) found this species to be the most abundant in Texas (37 percent). As to transmission, Cram and Jones (1929) found that various species of Coleopters may act as intermediate hosts of M. carioca. Subsequent studies by Jones indicated that the beetles that could transmit these tapeworms belong to the Scarabacidae and possibly to the Carabidae.

Intermediate Hosts of H. carioca. COLSOPTERA: Pamily Carabidae: Anisotarsus agilis (Dej.); Pamily Scarabaeidae: Aphodius granarius (L.), Choeridium historoides (Web.); Pamily Historidae: Caroinops quatuor-decimstriata Steph.

Raillietina tetragona. The third ranking tapeworm in prevalence in the vicinity of Manhattan appears to be Raillietina tetragona. It was third (38 percent) in Perry's studies in eastern Kansas and second in Texas (adams and Geiser). This large tapeworm attaches itself to the lower portion of the small intestine and may produce

nodules of the intestine which closely resemble lesions of tuberculosis. Horsfall (1938) reported that the common pavement ants Tetramorium casepitum (L.) and Pheidole vinelandica Forel. could serve as intermediate hosts of R. tetragona. Repeated attempts to infect various species of beetles with feedings of gravid R. tetragona proglottids have been unsuccessful, so it is probable that beetles are not natural intermediate hosts of this tapeworm. Studies upon ants as means of transmitting chicken tapeworms from one host to another are in progress at the Kansas Agricultural Experiment Station.

Intermediate Hosts of R. tetragona. HYMEHOPTERA:
Family Formicidae: Tetramorium caespitum (L.) and Pheidole
vinelandica Porel (Jones and Horsfall, 1934).

Choanotaenia infundibulum. A fowl cestode that is of very common occurrence if the vicinity of Manhattan, Kansas, is Choanotaenia infundibulum. Both Ferry (1934) and Adams and Geiser (1935), however, found this species to be the least abundant of the tapeworms they studied. Its habitat is the duodenum.

As to intermediate hosts, Guberlet (1916) found that houseflies (Musca domestics L.) are a means of transmitting Choanotaenia infundibulum from one chicken to another.

Cram (1928), Cram and Jones (1929) and Wetsel (1936)

obtained evidence that pointed to beetles as being important intermediate hosts of this tapeworm. Horsfall and Jones (1937) added six species of beetles and two grasshoppers as new intermediate hosts of <u>G</u>. <u>infundibulum</u>. The writer has found recently that three additional species of beetles may serve as intermediate hosts of this tapeworm. They are: <u>Amera fallex Lec.</u>, <u>Anaferonia constricts</u> Sey, and <u>Tenebroides mauritanious</u> (L.).

Further evidence that houseflies may serve as intermediate hosts of <u>C</u>. <u>infundibulum</u> was presented by Wetzel (1936) who reported that only 20 percent of the houseflies in his experiments developed cysticercoids and that the number of cysticercoids was small. Reid and Ackert (1937) found a natural infection of 91 cysticercoids in a housefly at Manhattan, Kansas. Even if only 20 percent of the houseflies developed cysticercoids, the quantities of flies around poultry yards at certain times of the year could easily supply the cysticercoids necessary to produce the heavy tapeworm infections (75 percent) in the naturally infected chickens in an experiment by Reid and Ackert (1937) at Manhattan, Kansas.

Intermediate Hosts of <u>C</u>. <u>infundibulum</u>. <u>COLEOPTERA:</u>
Family Carabidae: <u>Amara fallax</u> Lec., <u>Anaferonia constricta</u>
Say, <u>Cratacanthus dubius</u> Beauv., <u>Stenocellus debilipes</u>
(Say), <u>Stenolophus conjunctis</u> (Sey); <u>Family Ostomidae</u>:

Tenebroides mauritenicus (L.); Family Scarabacidae:

Aphodius granerius (L.), Aphodius sp., Geotrupes sylvaticus

Pans.; Family Staphylinidae: Apocellus sphaericollis (Say);

Family Tenebrionidae: Alphitophagus bifascietus (Say).

DIPTERA: Family Muscidae: Musca domestica (L.).

ORTHOPTERA: Family Locustidae (Acrididae):

Dicromorpha viridia (Scudder); Melanoplus femur-rubrum
(Dadeer).

Raillietina echinobothrida. Among the less abundant tapeworms in Kansas is the largest species Raillietina echinobothrida. This worm resembles R. Tetragona closely and is distinguished from it with difficulty. R. echinobothrida also attaches itself in the posterior part of the small intestine, and has about the same effects upon the host as R. tetragons. Perry (1934) found this cestode in 4 percent of the chickens examined in Douglas County. Kansas, and Adams and Geiser found it in 12 percent of the fowls examined in Texas. According to Horsfall (1938), the common pavement ant, Tetramorium caespitum may serve as an intermediate host of R. echinobothrida. An interesting observation was made by Horsfall (1938) who stated, "The first clue to the intermediate hosts of R. echinobothrida and R. tetragona was discovered while the writer had under observation in the experimental yard several fecal samples

containing R. schinobothrida proglottids. An ant carried one of these segments from the feces to an entrance to a nest and disappeared with it. Ants were then examined from this yard and all were found to be negative until August 16, 1935, at which time 3 T. caespitum were dissected and found to contain 4 cysticercoids the scoleces of which resembled those of R. schinobothrida. Larvae of R. schinobothrida in the naturally infected ant Tetramorium semilaeve from Marseilles, Prance, were reported by Joyeux and Baer (1937).

Intermediate Hosts of R. echinobothrida. HYMEHOPTERA:
Family Pormicidae: Tetramorium caespitum (L.) T.
semilaevo, and Pheidole vinelandica Forel.

Amoebotaenia sphenoides. A tapeworm present in 4 percent of the chickens examined by Ferry (1934) was the small wedge-shaped Amoebotaenia sphenoides. This cestode, which apparently is of rare occurrence in the United States was reported from 8 percent of the chickens examined by Adams and Geiser (1933) in Dallas County, Texas. The intermediate hosts are annelids, according to Mönnig (1927) who, in 14 days, grew the cysticercoids in earthworms

Ocnorodrilus (Ilyogenia) africanus Beddard . Four weeks were required for the cysticercoids to develop into adult tapeworms in chickens. Orassi and Rovelli (1889) and

Meggit (1916) undoubtly secured cysticercoids of this tapeworm by feeding the oncospheres to earthworms (Allolobophors feetida Ric.)

Intermediate Hosts of A. sphenoides. OLIGOCHARTA:
Family Lumbricides: Allolobophora foetida (Eisen), and
Ocnerodrilus (Ilyogenia) africanus Boddard.

Davaines proglottins (not reported from Kansas). A small tapeworm (four to nine segments) that is found in many countries, but has not been reported from Kansas is Davaines proglottina. Because of its minute size it may have been overlooked. Various writers have described it and Levine (1938) has studied phases of its biology. Wetzel (1936) reported that the most important intermediate host of D. proglotting in Germany is the garden slug Agriolimax agrestis. Monnig (1938) lista as intermediate hosts of D. proglotting the slugs: Linex cinerus, Arion sp., Cepoa sp., and the small Physa heterostropha (Say). Of these genera, Agriclimax and Physa are found in Eanses; so that it is possible that D. proglotting for which these gastropods may serve as intermediate hosts, is also present in Kansas, especially in the eastern part of the state where the rainfall is heavier.

Intermediate Hosts of <u>D. proglettine</u>. PULHOHATA:
Femily Arionidae: <u>Arion</u> sp.; Family Limecidae: <u>Arriolimex</u>

sp., Copon sp., Limax cinerus; Pamily Physicae: Physa heterostropha (Say).

SUMMARY OF PART I

- 1. Of ten species of fowl tapeworms reported from the United States, the following six species have been found in chickens in Kansas: Raillieting costicillus (Molin, 1858), Choanotaenia infundibulum (Bloch, 1779), Raillieting tetragena (Molin, 1858), Hymenolopis caricca (Magalhaes, 1898), Raillieting schinobothrida (Megnin, 1880), and Amosbotaenia sphenoides (Railliet, 1992).
- 2. The three following species of beetles are here reported for the first time as intermediate hosts of the chicken costode Choanotaenia infundibulum: Amera fallax Lec., Anaferonia constricte Say, and Tenebroides mauritanicus (L.).
- 3. The known species of intermediate hosts of the tapeworms found in Kansas chickens consists of beetles, flies, ants, slugs, snails, and earthworms. They are given according to order, family, and genus for the respective tapeworms.
- 4. Nost numerous of these intermediate hosts are ground beetles (Carabidae) of which eight genera and 25 species have been identified as intermediate hosts of chicken cestodes.

Altogether, 19 genera and 29 species of beetles may act as intermediate hosts for one or more of the important chicken tapeworms found in Kansas.

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PART II. SPINECTS OF THE FOWL TAPEWORM RAILLISTING CENTULIAS (WOLIN) UPON OROWING CHICKERS

INTRODUCTION

Experimental evidence of effects of tapeworms upon their hosts is almost lacking. This doubtless has been due largely to inability to secure at one time large numbers of infective tapeworm larvae and their definitive hosts. Circumstantial evidence has pointed to certain effects of cestodes upon man, but such effects have been attributed to worry from the knowledge of having a tapeworm rather than to the tapeworm itself. While some knowledge of the subject can be gained by comparing the symptoms of an infected animal with its behavior after anthelmintic treatment, more reliable evidence would be obtainable from numbers of animals of approximately the same age and inheritance, one half of which were parasitized and the other half used as controls.

The literature pertinent to the intermediate hosts of fowl tapeworms found in Kansas, which has been reviewed in Part I of this study, showed that of various invertebrates (ants, flies, beetles, slugs, and snails) ground beetles (Carabidae) are the most important intermediate hosts of the fowl tapeworm <u>Railliotina costicilus</u> (Nolin). From

the work of Gram (1928), Gram and Jones (1929), Wetzel (1936), and Reid, Ackert and Case (1938) upon beetles as intermediate hosts and the ease with which they may be infected, experiments were begun to study the effects of the fowl cestode R. cesticilius on growing chickens.

MATERIALS AND METEODS

The experiments were conducted in the animal house, a tightly screened, frame structure with concrete floors. To this house the fowls were brought as day-old chicks from a commercial hatchery, and given an adequate diet. The ground beetles (Carabidae) were collected at some distance from any poultry yard to avoid previous infection with tapeworm eggs. As the beetles were known to be cannabalistic, each was placed in a separate container. After a few hours, the beetles readily take the proglettids.

In the experiments, each beetle was fed a motile gravid proglottid of the tapeworm R. costicilius collected from the foces of infected chickens. After returning the beetle to the jar about three inches of moist earth was added, the lid placed on loosely, and the jar set in a dark place.

The beetles were then fed on various animal tissues, such as meat scrap and live beetles. The most satisfactory method was to place in the jar at weekly intervals, a half dosen experimentally reared meal beetles (<u>Tonebrio</u> sp.)
Water was supplied daily and the jars checked for dead
beetles, food, and meisture content. Under such conditions
the beetles lived 20 to 60 days.

In the bodies of the beetles the tapeworm eggs developed into cysticercoids in from 15 to 20 days. Their removal presented problems: (1) the taxonomic structures of the beetles had to be left intact, and (2) the cysticercoids had to be removed alive and infective. Gassing the beetles also killed the cysticercoids. Continued immersion in tap water caused evagination of the cysticercoids which rendered them non-infective.

One of the best methods of removing the cysticercoids was to anesthetise the beetle with ether, pin it dorso-ventrally in a blackened dissecting dish, and float the cysticercoids from the body cavity of the beetle through an opening in the dorsum. To prevent evagination of the cysticercoids, a 0.1 per cent MaCl solution was used.

Measurements of three cystercoids selected at random from each beetle were quickly made as an identification check.

The cysticercoids to be fed to the chickens were transferred from the salt solution to filter paper or were given by means of a small pipette to whose bulb was attached a Hoffman screw compressor. With a slight turn of the screw, the cysticercoids were drawn into the pipette

and held there until fed.

As to methods of blood examination, the hemoglobin tests were made with the New Dare hemoglobinometer. The blood sugar tests were made according to the method of Folin and Wu.

EXPERIMENTAL DATA

In the first experiment, 21 chickens three weeks old were divided into two groups of 17 and four birds, respectively. Each of the 17 chicks was given 10 cysticercoids of R. costicillus per week for three weeks. The other four chickens which received no cysticercoids were kept as controls. Weights of all birds were taken each week during the period the cystercoids were administered. After the last feeding of cystercoids, the chickens were hept index observation for six weeks. During the seventh, eighth, and minth weeks following the final feeding of cysticercoids hemoglobin and blood sugar tests were made (Table 1). From the time the chicks were first parasitized until the blood sugar determinations were made the controls and the parasitized birds were kept in the same pens and had the same feed.

At the beginning of the experiment, the parasitized and control chickens were matched as nearly as possible. For example, parasitized chicken A 690 may be compared with control chicken A 70%. Both weighed 116 gm. at the start of the experiment. In the blood tests, the former had a homoglobin percentage of 52 as compared with 62 for the control. The blood of the parasitized bird contained 168.8 mg. of sugar per 100 cc. of blood as compared with 181.6 mg. for the control chicken. As to growth, the parasitized chicken gained 156 gm. while the control added 284 gm. Chilling was observed in the parasitized chicken but not in the control (Table 1).

The heaviest control chicken (A 667) weighing 132 cm. as compared with the parasitized bird (A 659) which weighed 125 gm. showed less marked differences. There was scarcely any difference in the hemoglobin percentage of the blood. The parasitized chicken (A 659) had 186.8 mg. of sugar as compared with 212 mg. in the control (A 667). In weight the parasitized bird gained 139 gm. and the control 154 gm. Again the parasitized bird manifested chilling while the control failed to do so. As a group, the parasitized chickens averaged 118.5 gm. while the controls averaged 120.5 gm. at the beginning of the experiment. The hemoglobin percentage on the average was practically the same for both groups. That there was a difference in the sugar volume is indicated by an average of 172.7 mg. as compared with an average of 186.7 mg. for the control group. In weights also there was a marked difference.

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Table 1. Short

Showing comparative results of blood tests and gains in weight of parasitized chickens with those of unper-sitized fouls. All chickens were 14 weeks of sge. The parasitized ones becare infected, Pariod of parasitizen I weeks.

		Pa	Parasitized			
DATCK	beginning	M 70 -	(cm.)	globin (%)	BLOOD Sugar (mg.)	United
A 654	10.5	226		67	149.0	Yes
A 659	125	264		64	186.8	Yes
A 5555	125	264		99	150.4	Yes
A 663	121	212		533	220.0	Yes
A 677	120	320	200	20	181.0	Kes
A 683	120	286		70	188.0	Yes
A 692	120	304		79	174.6	Yes
A 699	120	25.56		80	145.0	Yes
A 700	125	280		59	198,0	Tes
A 661	117	268		68	102.0	Yes
A 662	114	260		99	178.6	Yos
A 665	112	203		80	207.2	Yes
A 666	115	284		555	186.8	Yes
A 668	114	284		55	175.4	Yes
A 673	114	241		99	158.3	Yes
A 686	115	246		55	155.8	Yes
A 690	116			200	168.8	Yes
Average	110.5	2/5.8	60	62.3	172.7	Yes
		Ш	Controls			
A GES	118		178	54	0.882	No
A 667	132		154	100	212.0	No
A 691	116		152	ස ද	131.6	OH
A 706	116	200	284	62	181.6	No
Average	120.5	10	192	63	186.7	2
						0

The parasitized birds made an average gain of 151.7 gm. as compared with a 192 gm. average for the control group. All of the parasitized chickens manifested chilling on cool or rainy days, whereas no such behavior was observed in the controls which usually remained at the feed hoppers. From these results, it appears that infections with the tapeworm R. costicillus cause reduction in the blood sugar and in the growth of chickens and interfere with normal behavior of the fowls (Table 1).

In the second experiment, 13 chicks 9 days old were divided into groups of 10 and three, respectively. Each of the 10 chicks was given 10 cysticercoids of E. cesticillus per week for eight weeks. The other three chickens which were kept as controls received no cysticercoids. Weights of all birds were taken each week during the period the cysticercoids were administered. After the last feeding the chickens were kept under observation for six weeks. During the seventh and eighth weeks following the final feeding of cysticercoids the hemoglobin and blood sugar tests were made as in Experiment I. From the time the chicke were first parasitized until the blood sugar determinations were made the controls and the parasitized birds were kept in the same pens and had the same feed.

At the beginning of experiment II, the young chickens

were matched as nearly as possible according to weight.
Comparing individuals of the two groups, it may be seen
that the parasitized chick (A 500) which at the beginning
of the experiment weighed 31 gm. and its control (A 606),
53 gm. differ but little in the hemoglobin percentage:
57 for the parasitized chick and 56 percent for the control.
A marked difference occurred in the sugar; parasitized
chicken (A 500) had 136.8 mg. as compared with 165.2 mg.
of sugar for the control. In growth, there was also a
difference. The parasitized bird gained 909 gm. as compared
with 1197 for the control. The chilling noted in Experiment
I recurred also in Experiment II in the parasitized group,
whereas no such behavior occurred among the controls.

A parasitized chick (A 604) and a control chick
(A 585) which weighed the same amount at the beginning of
the experiment showed some variation. The hemoglobin
percentage remained somewhat higher in the parasitized bird
(67 per cent) than in the control (60 per cent), but the
blood sugar in the parasitized bird which had five large
cestodes was only 144.4 mg. of sugar per 100 cc. of blood
as compared with 172 mg. in the control. In growth, there
was also a marked difference. The parasitized bird gained
946 gm. to 1521 gm. by the control.

As groups, the parasitized birds averaged 53.8 gm. at

the beginning of the experiment while the controls averaged 54 gm. The hemoglobin percentage averaged only slightly less in the parasitized group (55.1 per cent as compared with 56.3 per cent in the controls). However, in blood sugar, the parasitized birds averaged 135.8 as compared with 167.06 mg. The average gains in weight likewise showed a marked difference. The parasitized birds gained an average of 971.4 gm. while the controls gained an average of 1217.7 gm. The chilling which was characteristic of the parasitized chickens in the first experiment occurred also in the second group of parasitized chickens. It was entirely absent from the control group.

The results of Experiment II as shown in Table 2 indicate that an infection of fowl tapeworms over a 15 week period failed to produce much effect upon the homoglobin percentage but reduced markedly the amount of sugar in the blood and the rate of growth of the chickens. That the parasites were the cause of these differences was indicated in part by the chilling of the chickens that were infected.

DISCUSSION

Up to the present, the results indicate that growing chickens that have received 30 or more cysticercoids when infected for two to three months show reduced amount of

Showing comparative results of blood tests and gains in weight of parasitized chickens with those of unparasitized fowls. All chickens were 15 weeks of age. The parasitized ones became infected. The pariod of parasitism was 15 weeks.

				Parasitized	zed			
Deginning at end weight globin (gm.)	Chiek	Weight at	Weight	Gain in	Нешо-	Blood	Tape-	Chilling
(gm.) (gm.) (gm.) (f) 51 1050 (gm.) (f) 49 49 95 57 49 61 1175 1114 45 54 1000 1446 65 62 950 1846 67 63 955 864 45 647 747 698 55 47 1555 1308 55 50 1025.2 971.4 55.1 55.8 1375 1197 56 54 1377.66 1317.7 56 55 1270 1137 56 56 1277.66 1317.7 56 56 1277.66 1317.7 56 56 1277.66 1317.7 56 56 1277.66 1317.7 56 57 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.66 1377.7 56 58 1377.7 57 58 1377.7 57 58 1377.7 57 58 1377.7 57 58 1377.7 57 58 1377.7 57 58 1377.7 57		ginni	at end	weight	globin	sugar	WOLMS	
\$1 1050 999 57 \$4 1000 946 65 \$4 1000 946 67 \$5 1000 946 67 \$5 1000 946 67 \$5 900 880 45 \$5 900 880 55 \$5 1255 1308 86 \$5 1025.2 971.4 55.1 \$5 1255 1391 56.1 \$5 1275 67 1317 \$5 1275 67 1317 \$5 1275 13190 1135 \$5 1275 67 1317 \$5 12		(gm.)	(800)	(gm.)	(%)	(mg.)		
49 950 901 55 61 1175 1114 45 54 1000 1446 65 62 950 884 45 62 950 884 45 64 747 696 55 47 747 696 55 50 1025-2 971-4 55-1 55 1250 1137 56 54 1375 1381 55 1250 1137 56 54 1375 1381	A 590	51	1050	666	57	133.8	4 large	
61 1175 1114 45 54 1000 946 67 54 1000 1146 65 62 950 884 45 69 950 888 47 47 1255 1308 55 55 900 850 57 55 1255 1714 551 55 1715 1715 55 54 1775 1715 55 55 1275 1715 55	A 592	49	950	106	55	216.0	0	
54 1000 946 67 61 1200 1146 65 62 950 888 47 49 747 698 55 50 1055-2 1308 55 50 1055-2 971-4 55-1 53 1250 1137 58 54 1375 1321 55 1276 1137 58 54 1275 1321 55 1276 1137 58 56 1277-66 1375 58	A 586	19	1175	1114	45	177.4	100	Yes
54 1200 1146 65 61 925 864 45 62 925 864 45 49 747 698 55 47 1255 1308 55 50 900 850 57 55.8 1025.2 971.4 55.1 55.8 1375 1397 55 54 1375 1317 56 54 127.66 127.67 56 54 127.66 127.67	A 604	54	1000	946	29	144.4	5 large	
61 925 864 45 62 956 888 47 69 747 698 55 50 900 858 55 53*8 1025*2 971*4 55*1 53 1250 1137 55*1 54 1277 66 1321 55 1190 1135 54 1277 66 1377	A 598	54	1200	1146	65	148.0	0	
62 950 888 47 47 747 696 55 47 1355 1306 55 50 900 850 850 55 1250 1037 55 54 1375 1321 55 1276 1137 56 54 1277 66 1277, 66 3	A 580	19	925	864	45	160.0	25 large	
49 747 698 55 47 1355 1308 58 50 900 850 57 55,8 1025,2 971,4 55,1 53 1250 197 55 54 1375 1321 60 55 120 1135 51 55 120 1377 56 56 1271,66 1317, 56 3	A 588	62	950	888	47	149.2	6 large	
47 1255 1308 58 50 900 850 57 53.8 1025.2 971.4 55.1 53 1250 1197 56 55 1275 1321 60 54 1277.66 13175 66 54 1277.66 13175 66	A 607#	49	747	869	55		12 large	
50 900 850 55.8 1025.2 971.4 55.1 53 1250 1097 58 54 1375 1321 55 127.66 1277.66 53	A 584	47	1355	1308	58	133.2	27 large	
55.8 1025.2 971.4 55.1 Controls 55 1250 1197 56 54 1275 1321 60 55 127 56 127.66 1217.07 56.3	A 594	20	006	850	57	85.0	22 large	Yes
53 1250 1137 58 54 1375 1321 60 55 1190 1135 54 1271.66 1217.07 56.3	Average		1025.2	971.4	55.1	135.8	10.4	
54 1250 1197 58 54 1375 1321 60 55 1190 1135 51 54 1277 66 3				Control	83			
54 1375 1321 60 55 1190 1135 51 54 1271.66 1217.07 56.3	A 606	53	1250	1197	58	165,2	0	No
54 1271.66 1217.07 56.3	A 585.	54	1375	1321	09	172.0	0	No
54 1271.66 1217.07 56.3	A 595	55	1190	1135	51	185.0	0	No
1 0000	Average		1271,66	1217.07	56.3	167.06	0	No

The blood of this bird would not give a reading on either blood sample.

blood sugar and decreased gains in weight (Ackert and Case, 1938). Harwood and Luttermoser (1958) also found that chickens infected with the tapeworm R. cesticillus showed reduced rates of growth. In the present experiments little difference occurred between the hemoglobin percentages of the parasitized group and the controls.

Concerning sugar, Table 2 shows that one parasitized bird failed to give a blood sugar test which would indicate a very low sugar content of this bird's blood. This chicken, A 607, had 12 very large tapeworms, gained much less weight than the average of the parasitized group, and had a hemoglobin percentage of 55, slightly less than the group average. The parasitized chicken in Experiment II, A 592, had no tapeworms at postmortem and this may explain the high blood sugar test for this bird (Table 2). The data revealed that in both experiments, the parasitized birds gained less then the controls, and their blood sugar percentage was much lower, both, as individuals, and as groups. A very noticeable difference between the parasitised and control chickens was evident on cool mornings and on cool days following rains when the parasitized birds would ruffle their feathers, chill, and seek the warmth of the brooder heat unit while the controls in the same pen were apparently normal and remained at the feed hoppers, and never went near the heat units.

SUMMARY OF PART II

- 1. Nethods of culturing bettles for the purpose of obtaining cysticercoids of <u>R</u>. <u>cesticillus</u> are described. Pint Mason fruit jars containing a few inches of soil served as culture receptacles.
- 2. A successful technique for obtaining cysticercoids from beetles (Carabidae) and for peresitizing chickens with them was developed.
- 3. Data from two experiments indicated that chickens parasitized during a period of 11 to 15 weeks showed lowered blood sugar and reduced gains in weight.

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