Observations on the feeding behaviour of parasitic third-stage hookworm larvae

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SUMMARY

The feeding behaviour of parasitic 3rd-stage larvae (L₃) of the hookworms Ancylostoma caninum, A. ceylanicum and Necator americanus was examined. Less than 11% of A. caninum L₃ recovered from the small intestines of dogs infected orally were feeding at 4—48 h post-infection (p.i.), and none of the A. ceylanicum L₃ recovered from the intestines of orally infected hamsters had resumed feeding. All L₄ of both species recovered at 36 and 48 h p.i. had resumed feeding. On the other hand, approximately 16% of the A. ceylanicum L₃ recovered from the skin of percutaneously infected hamsters at 18 h were feeding, and the percentage feeding increased to nearly 58% at 44 h p.i. Necator americanus L₃ recovered from the skin of percutaneously infected neonatal hamsters resumed feeding at 6-12 h p.i. and reached 90-94% by 18 h. Feeding began to decline at 66 h, and reached 29% at 120 h p.i. This decrease was associated with the migration of larvae from the skin to the lungs. By 192 h p.i. over 95% of the larvae had reached the small intestine, and all had moulted to the L₄.

The results indicate that parasitic L₃ resume feeding in the skin during percutaneous infections, and suggest that feeding by hookworm L₃ correlates with the resumption of development.

Key words: Ancylostoma caninum, Ancylostoma ceylanicum, Necator americanus, hookworms, resumption of development, in vitro feeding.

INTRODUCTION

The behaviour of infective larvae of hookworms following entry into the host has been an area of long-standing research. Most of this research was conducted using the canine hookworm Ancylostoma caninum and, consequently, the migratory behaviour of A. caninum larvae in the dog and abnormal hosts following oral and percutaneous infections is relatively well defined (Scott, 1930; Foster & Cross, 1934; Banerjee, Prakash & Deo, 1970; Miller, 1971; Bhopale & Johri, 1975). More recently, the hookworms that infect man have been adapted to laboratory hosts (Ray and Bhopale, 1972; Sen, 1972; Schad, 1979), enabling studies of hookworm biology in models that, although not the usual definitive host, permit completion of their life-history (Behnke, 1990). Investigations of host–parasite interaction and the behaviour of human hookworm infective larvae during the early parasitic stages are now possible (Rajasekariah et al. 1985; Behnke, Paul & Rajasekariah, 1986a; Behnke & Pritchard, 1987; Gupta, Srivastava & Katiyar, 1987; Garside & Behnke, 1989).

Despite these advances, many aspects of the early parasitic portion of the hookworm life-history remain unknown. For example, the time at which invading larvae resume feeding is poorly understood. Hawdon & Schad (1990) suggested that larval feeding represents an initial event in the transition to the parasitic life-style, and have proposed that the resumption of feeding can be used as a marker for the reactivation of development in A. caninum 3rd-stage larvae. Subsequent experiments indicated that infective larvae of several hookworm species, including A. duodenale and A. ceylanicum, resumed feeding in vitro (Hawdon et al. 1992). The investigations reported herein were undertaken to determine the role of larval feeding in the life-history of hookworms.

MATERIALS AND METHODS

Ancylostoma caninum and A. duodenale were maintained in dogs as described previously (Schad, 1979, 1982). Necator americanus (Behnke, Wells & Brown, 1986b) and A. ceylanicum (Garside & Behnke, 1989) were maintained in hamsters. Infective larvae were harvested from 7 to 14-day coprocultures and stored in nematode handling buffer (BU; 50 mM Na₂HPO₄/22 mM KH₂PO₄/70 mM NaCl, pH 6-8) until needed (1-14 days). Storage under these conditions had no effect on subsequent feeding (Hawdon & Schad, 1991a).

Feeding was determined by the method of Hawdon & Schad (1990), with slight modifications. L₃ recovered from tissues were resuspended in approx-
imimately 0.1 ml of medium and transferred to individual wells of a 96-well tissue culture plate. An equal volume of 5 mg/ml fluorescein isothiocyanate-labelled bovine serum albumin (FITC-BSA) was added, and the larvae incubated at 37 °C, 5% CO₂ for 1–2 h. After incubation, the L₃ were washed in 15 ml of phosphate-buffered saline (PBS), pH 7.2, transferred to a microscope slide, and examined under ultra-violet illumination. Feeding larvae, indicated by the presence of dye in the intestine, were counted and the number expressed as a percentage of the total number of larvae counted.

Experiment 1 was designed to determine when A. caninum L₃ resumed feeding in dogs infected per os. Ten 12-week-old helminth-naive bengals were divided into 5 groups of 2 each, and each pair inoculated orally with 5000 A. caninum L₃ in gelatin capsules. Several of the dogs regurgitated following inoculation. In these cases, the vomitus was recovered, mixed with cottage cheese, and re-fed to the dog. In one case (4 h post-infection), the dogs were sedated and the vomitus given via stomach tube. Subsequent recovery of larvae in these cases was lower than from animals that had not vomited.

After the appropriate time, the dogs were killed by barbiturate overdose, and the small intestine removed (the stomach was also examined in the 4 h time-point). Each intestine was opened longitudinally, and cut into 3–4 sections. The sections were hung for 2–3 h in 21 cylinders containing PBS at 37 °C. After the incubation, the intestines were removed, and the sediments collected and concentrated. Dilution counts of the sediments were done to determine the number of larvae recovered, and 150–200 larvae were individually picked from the sediments, washed 3–4 times in 15 ml of RPMI tissue culture medium and assayed for feeding. The larvae recovered from each animal served as a replicate.

Experiment 2 was a similar experiment using hamsters inoculated orally with A. ceylanicum L₃. Six groups of 5 weanling LVG hamsters (Charles River Labs, Willmington, Massachusetts) were inoculated with approximately 500 A. ceylanicum L₃ in 0.1 ml by gastric gavage using a curved feeding needle attached to a 1 ml syringe. At the appropriate time post-infection (p.i.), the hamsters were killed by chloroform inhalation, and the small intestines removed. The intestinal lumina were rinsed with PBS, and the intestines opened longitudinally, then cut into approximately 1 cm pieces. The lumen wash and intestinal pieces from each hamster were subjected to the Baermann technique in PBS at 37 °C, 5% CO₂ for 1 h. Recovered L₃ were washed by centrifugation (1000 g) 3 times in 15 ml of PBS, followed by 2 washes in RPMI tissue culture medium. The larvae were assayed for feeding as described above.

Because of the expense and difficulty in isolating sufficient numbers of percutaneously infected L₃ from dogs, the feeding behaviour of skin-penetrating larvae was examined using the hamster–A. ceylanicum model. Groups of 2–3 adult hamsters were inoculated percutaneously with 200–2500 L₃ A. ceylanicum by the method of Behnke et al. (1986b) with slight modification. Hamsters were lightly anaesthetized by Halothane inhalation and their abdomen and back shaved. Larvae in 0.1 ml of water were applied to the pad on a 1 inch adhesive bandage (Curad), and the bandage affixed to the abdominal surface of the hamster. The bandage was secured with surgical tape wrapped around the abdomen. At the appropriate time-point, the hamsters were killed and the bandages removed. The infection site skin, underlying muscle, lungs and intestine were removed, minced and incubated in individual Petri dishes at 37 °C, 5% CO₂ for 4 h. After incubation, L₃ were counted and individually picked into microfuge tubes. The larvae were washed in RPMI, resuspended in 0.1 ml of medium, and transferred to a tissue culture plate for the feeding assay. Because of poor recovery at the 18 h time-point, the larvae recovered from 2 hamsters were pooled for feeding, and therefore each replicate at this time-point is derived from a separate experiment.

The feeding behaviour of N. americanus L₃ during percutaneous infection was also examined. N. americanus L₃ are obligate skin penetrators, and infection is established by percutaneous inoculation of 2 to 4-day-old hamster pups (Sen & Seth, 1970; Sen, 1972; Behnke, Wells & Brown, 1986). Therefore, the number of animals available for infection depends on litter size and mortality of the pups prior to infection, but a minimum of 2 pups were used per time-point. Hamsters were either bred on site, or purchased as timed-pregnant adults. Neonates were infected with approximately 1000 L₃ by the method of Behnke et al. (1986b). At the appropriate time, the pups were killed by decapitation, and the skin, lungs, and intestines removed to individual Petri dishes. The tissue was minced and incubated in RPMI medium for 2–3 h. L₃ were counted and picked to individual tubes, washed with RPMI, and assayed for feeding. The larvae recovered from each hamster represent 1 replicate in the feeding experiments, and between 40 and 140 total L₃ were examined per replicate, depending on the number of L₃ recovered.

Feeding data (percentages) were transformed to angular values by arcsin transformation. One-way ANOVA was performed on the transformed data, and the means compared using the Bonferroni post-test; P < 0.05 was considered significant. Mean angular values were transformed back to percentages for expression.
RESULTS

Recovery of L₃ *A. caninum* from orally inoculated dogs ranged from 2 to 56%. Recovery was lower from dogs that had vomited the inoculum, ranging from 2 to 28%, whereas recovery ranged from 45 to 56% in dogs that retained the inoculum. All of the larvae recovered were exsheathed.

As seen from the data presented in Table 1, *A. caninum* L₃ were feeding at low levels from 4 to 24 h post-infection. The percentage of L₃ that were feeding reached a maximum of 10-3 ± 5-6% by 24 h, after which time it dropped to 0% at 36 h. At this time, 1-5% (3/200) of the larvae examined for feeding were 4th-stage larvae (L₄), whereas 64-0% (137/214) of larvae examined at 48 h were L₄, indicating that larval development had resumed. The mean increase in feeding at 48 h was caused by the L₄, which had resumed feeding (Table 1).

### Table 1. Feeding by *Ancylostoma caninum* larval stages recovered from orally infected dogs

| Time p.i. (h) | Dog | L₃ feeding* (%) | L₄ feeding (%) | Mean percentage feeding |
|---------------|-----|-----------------|----------------|------------------------|
| 4             | P3  | 4 (3/85)        | N.R.           | 1.0±1.4                |
|               | L6  | 0 (0/79)        | N.R.           |                        |
| 12            | T5  | 0 (0/100)       | N.R.           | 2.8±4.0                |
| 24            | J3  | 4 (3/81)        | N.R.           | 10.3±5.6               |
|               | P2  | 19 (7/37)       | N.R.           |                        |
| 36            | T4  | 0 (0/100)       | N.R.           | 0.8±1.1                |
| 48            | J4  | 0 (0/29)        | 100 (85/85)    | 64.4±11.3              |
|               | P5  | 2 (1/48)        | 100 (52/52)    |                        |

* (Number feeding/total number examined.)
† N.R., None recovered.

Table 2. Feeding by *Ancylostoma ceylanicum* 3rd-stage larvae recovered from percutaneously infected hamsters

| Time p.i. (h) | Replicate | Percentage feeding | Mean percentage feeding ± S.D. |
|---------------|-----------|--------------------|-------------------------------|
| 18            | 1         | 15.8 (15/95)*      | 15.7±0.7                      |
|               | 2         | 15.6 (5/32)*       | 15.7±0.7                      |
| 24            | 1         | 46.1 (41/89)       | 46.3±0.1                      |
|               | 2         | 46.4 (32/69)       | 46.3±0.1                      |
| 44            | 1         | 65.0 (65/100)      | 57.6±7.2                      |
|               | 2         | 50.0 (8/16)        | 57.6±7.2                      |

* Larvae recovered from 2 hamsters were pooled for feeding.

Table 3. Total and tissue-specific recovery of *Necator americanus* larvae from percutaneously infected neonatal hamsters

| Time p.i. of dose recovered* (h) | Percentage of dose recovered* | Mean percentage of recovered larvae from: |
|---------------------------------|------------------------------|-----------------------------------------|
| 6                               | 30.8±4.4                     | Skin 100  N.D.  N.D.                     |
| 12                              | 48.5±0.6                     | Skin 100  N.D.  N.D.                     |
| 18                              | 49.3±1.5                     | Skin 100  0  0                         |
| 24                              | 39.0±4.8                     | Skin 100  0  0                         |
| 48                              | 43.3±1.1                     | Skin 92.6  24  0                       |
| 66                              | 40.3±12.8†                   | Skin 59.4  40.6  0                     |
| 72                              | 43.3±0.5                     | Skin 44.8  38.2  0                     |
| 90                              | 47.4±2.7                     | Skin 11.2  88.8  0                     |
| 120                             | 53.7±5.7†                    | Skin 1.2  98.8  0                      |
| 192                             | 12.6±4.1                     | Skin 0  4.1  95.9                      |

* Mean %, n = 3; n = 2 for 18, 66 and 120 h; n = 4 for 192 h.
† A portion of inoculum was lost during infection.
N.D. Not determined.

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Fig. 1. Feeding behaviour and recovery of *Necator americanus* larvae from percutaneously infected neonatal hamsters. (A) Feeding. Each point represents the mean feeding ± S.D. of the larvae recovered from a minimum of 2 hamsters. (B) Graphical representation of tissue-specific recovery data presented in Table 3.
None of the L₃ recovered at any time-point from hamsters infected with *A. ceylanicum* per os had resumed feeding, whereas all of the L₄ recovered were feeding. At 36 h, 1% of the larvae examined were L₄, whereas by 48 h, the number of L₄ increased to 9% of the examined larvae.

The feeding behaviour of L₄ from a percutaneously initiated infection was examined using *A. ceylanicum* L₂. As shown in Table 2, 15.7 ± 0.7% of the L₃ isolated from the skin of percutaneously infected hamsters had resumed feeding by 18 h. The percentage feeding increased to 57.6 ± 7.2% at 44 h. No larvae were recovered from any of the tissues examined other than those of the infection site skin and underlying muscle. Recovery of larvae was variable, ranging from 2 to 23%. The decreased efficiency of infection by the percutaneous route (Gupta et al. 1987) undoubtedly accounts for the low recovery rates. The L₃ had exsheathed, but there was no morphological evidence of further development, although percutaneously infected control hamsters developed patent infections within 3 weeks p.i., confirming that skin penetration is a viable route of entry for *A. ceylanicum* L₃.

*N. americanus* L₃ recovered from percutaneously infected neonatal hamsters resumed feeding 6–12 h p.i., and reached a maximum percentage feeding, at 90–94%, by 18 h (Fig. 1A). Feeding remained at this high level until 66 h, when the percentage feeding began to decline. This decline coincided with decreased recovery of larvae from the skin, and increased recovery from the lungs (compare Fig. 1A and B). Feeding decreased to 29.0 ± 1.6% at 120 h, when 98.8 ± 4.2% of the recovered L₄ were from the lungs. At 192 h p.i., all of the recovered larvae were L₄ and had resumed feeding. The overall recovery had dropped from approximately 50% at 120 h to 12.6% at 192 h, 95.9 ± 10.2% of which were recovered from the small intestine (Table 3). This decreased recovery was seen in previous studies, and the overall migratory pattern was similar to published reports (Rajasekariah et al. 1985; Behnke et al. 1986b).

The tissue-specific feeding levels of larvae recovered at 66, 72, and 90 h p.i. were compared (Fig. 2). There was no significant difference between the percentage of feeding larvae recovered from the skin when compared to larvae recovered from the lungs when compared to larvae recovered from the skin, indicating that the decline in feeding resulted from an overall decrease in feeding, and not from decreased feeding by a particular tissue-associated developmental group.

**DISCUSSION**

Although the biology of the free-living and adult stages of hookworms has been extensively studied, little is known about the biology of the parasitic larval stages (Hoagland & Schad, 1978). Most investigations of this portion of hookworm life-history have concentrated on the migratory behaviour of larvae within the host. However, physiological, biochemical and molecular aspects of infecting and developing stages remain poorly understood, due largely to the technical difficulties in recovering sufficient numbers for such studies, and the inability to culture hookworm larvae beyond the L₂ in vivo. For example, the host signal that re-activates development in the invading L₂ is poorly characterized. In order to investigate the early events of the infectious process, including the mechanism of developmental re-activation, Hawdon & Schad (1990) proposed the resumption of feeding as a marker for the re-activation of development, and developed an in vitro assay to measure feeding by *A. caninum* L₂. Subsequent experiments indicated that albumin (Hawdon & Schad, 1991b) and glutathione (Hawdon & Schad, 1992) stimulated feeding, and that in vitro feeding occurred in most species of hookworms examined, including *A. duodenale* and *A. ceylanicum* (Hawdon et al. 1992). The experiments presented here were conducted to determine the role of feeding by parasite L₂ in the life-history of the hookworm.

*A. caninum* L₃ recovered from the small intestine of orally infected dogs were feeding at very low levels. Furthermore, orally inoculated *A. ceylanicum* larvae failed to feed until after moulting to the L₄. One possible explanation for the lack of feeding larvae is that only non-feeding larvae were sufficiently motile to be recovered by the techniques employed. However, in most cases sufficient numbers of larvae were recovered to ensure against selection for non-feeding larvae. Another possibility is that feeding was depressed by removal of the...
Feeding behaviour of hookworm larvae

larvae from the host for extended times during the isolation and feeding procedure. However, all incubations during recovery were under host-like conditions (37 °C, 5% CO₂), which are employed for in vitro feeding studies (Hawdon & Schad, 1990). Once L₃ have begun to feed in vitro, they continue to feed for a minimum of 3 h after removal from the incubation conditions. It is unlikely that further incubation after removal from the host would decrease feeding. Furthermore, larvae that were feeding, such as A. ceylanicum and N. americanus L₃ isolated from skin, were subjected to similar conditions. It is more likely that few of the Ancylostoma spp. L₃ feed after infection per os.

Despite the primacy of the oral route for A. caninum (Wilson, 1982) and A. ceylanicum infection (Gupta et al. 1987), L₃ of these species can also infect their hosts via the percutaneous route (Yoshida, Okamoto & Chiu, 1971; Behnke, 1990). Therefore, feeding behaviour of L₃ isolated from the skin was examined. The A. ceylanicum–hamster model was utilized because of the expense and logistical difficulties of recovering larvae from dogs. The time-dependent increase in feeding by L₃ recovered from the skin of hamsters indicated that feeding occurred in A. ceylanicum parasitic L₃ when the larvae entered via the skin.

Apparently, the facultative skin-penetrating hookworms, such as A. ceylanicum and A. caninum, employ different feeding behaviours depending on their route of entry. One possible explanation for this difference in feeding behaviour is that feeding provides nutrients to meet the energetic demands of migration. In order to reach the intestine, L₃ that enter through the skin must undergo tissue migration, which requires both time and energy expenditure. Indeed, larval development was slower in percutaneous infections than in oral infections, as A. ceylanicum L₃ remained in the skin for 44 h without migration or obvious morphological development. Skin-penetrating L₃ would be exposed to GSH and the phagostimulant(s) present in serum during this skin phase and the subsequent migration, whereas L₃ that enter orally would not be exposed to stimulatory levels of the phagostimulants.

However, feeding is not required for development to resume, as larvae that entered orally resumed development without feeding as L₃. This suggests that the feeding stimulus is independent from the stimulus that re-initiates development. The developmental stimulus is present in the host's intestine, but if this signal is specific to the predilection site, then migrating larvae, although feeding, would not be developing. Once they reach the small intestine, they would be exposed to the second stimulus, and development would resume. Alternatively, if the developmental stimulus was present throughout the host, feeding would indicate that the larvae had resumed development. However, if the feeding stimulus is restricted to parenteral locations, L₃ entering orally would not feed, despite having resumed development. Feeding, therefore, is not mandatory for development, but larvae that have resumed feeding are probably also developing. Considerably more work is required to clarify this interesting situation, which apparently has not been investigated previously by parasitologists and others interested in nematode development.

Although the role of feeding in the life-history of the Ancylostoma spp. is uncertain, the results of in vitro experiments indicate that feeding by parasitic L₃ is an important facet of the life-history of N. americanus. Feeding resumed shortly after entry into the host, and continued while the larvae were in the skin, but decreased as the larvae migrated to the lungs from 48–90 h p.i. This decrease in feeding was a programmed response, and not merely the result of changing locations, as there was no significant difference (P > 0.05) in feeding between L₃ isolated from the skin and lungs at any time-point of the migration. By 120 h, over 98% of the larvae had reached the lungs, and feeding had decreased to approximately 30%. Feeding resumed following the moult to the L₄.

N. americanus is an obligate skin-penetrating hookworm, and larvae cannot establish in the intestine when inoculated orally (Komiya & Yasu- raoka, 1966; Hoagland & Schad, 1978; Behnke et al. 1986a), unless they succeed in penetrating the oral mucosa (Nagahana et al. 1962). Traditionally, this has been interpreted as indicating the requirement of an obligatory lung migration for development to continue (Komiya & Yasuraoka, 1966; Hoagland & Schad, 1978), and morphological development during the long period of residence in the lungs (Sen & Seth, 1979; Behnke et al. 1986a) further supports an obligatory lung phase. However, N. americanus L₃ remained in the skin for 48–72 h p.i. (Behnke et al. 1986a), during which time their motility decreased (Ishikawa, 1966). This period in the skin may be required to re-activate developmental pathways and initiate expression of genes required for the parasitic stages (Rogers & Petronjievic, 1982). Indeed, the results presented indicate that one event occurring during this skin phase is the re-activation of larval feeding. Therefore, the host stimulus that induces feeding is present in the skin, and encountered by the larvae during infection. However, it is possible that there is a second signal, present in the lungs, that actually leads to the re-activation of developmental pathways, and that feeding is not invariably linked to development. Although direct evidence is absent, results argue against this suggestion. The decline in N. americanus L₃ feeding between 48 and 120 h probably represents the onset of the lethargus prior to the L₃–L₄ moult. The onset of a programmed decrease in feeding, or lethargus, prior to arrival in the lungs suggests that devel-
opmental pathways were re-activated in the skin and, therefore, feeding by L₃ indicates that development has resumed.

Although feeding by parasitic L₃ is an important event during *N. americanus* infections, attempts to induce similar feeding *in vitro* have been unsuccessful, indicating that the *in vitro* feeding system lacks the host signal that initiates feeding and development of *N. americanus* L₃ *in vivo*. The nature of this signal remains unknown. Salafsky et al. (1990) found that linoleic acid stimulated penetration of artificial membranes and triggered the production of eicosanoids, but linoleic acid failed to stimulate feeding *in vitro* (Hawdon et al. 1992). The requirement for separate signals to initiate penetration and feeding supports the hypothesis (Matthews, 1977) that the infectious process is composed of distinct steps, including penetration, exsheathment and the re-activation of development and feeding, that are initiated by independent factors encountered in the host, and required in sequence for successful development to the adult stage.

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