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## EMERGENCE OF THIRD-STAGE LARVAE OF *UMINGMAKSTRONGYLUS PALLIKUUKENSIS* FROM THREE GASTROPOD INTERMEDIATE HOST SPECIES

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**ABSTRACT:** We investigated the emergence of third-stage larvae (L3) of *Umingmakstrongylus pallikuukensis* from the slugs *Deroceras laeve*, *Deroceras reticulatum*, and the snail *Catinella* sp. in the laboratory and from *D. laeve* on the tundra. Third-stage larvae emerged from 8 of 8 *D. laeve* and 8 of 8 *D. reticulatum* housed at 20 C in darkness and from 9 of 10 *D. laeve* and 5 of 5 *Catinella* sp. housed at 21 C with 10–12 hr of light/day. Larvae emerged from *D. laeve* and *D. reticulatum* over a wide range of infection intensities (2–179 and 20–65, respectively), and the patterns of emergence were independent of intensity. The majority of the L3 emerged from most of the *Deroceras* spp. by 58 or 60 days postinfection (PI). Lower rates of emergence were observed from *Catinella* sp. Larvae emerged from *D. laeve* on the tundra by 10 wk PI and were recovered from the vegetation in some experimental enclosures the following year. Third-stage larvae survived in tap and distilled water at 0–4 C for 13 mo. Emergence of L3 of *U. pallikuukensis* from the intermediate host may increase the temporal and spatial availability of L3 and enhance their survival and transmission.

*Umingmakstrongylus pallikuukensis*, a protostrongylid lung-worm known only in muskoxen (Hoberg et al., 1995) is present at a high prevalence and intensity in the host population west of Kugluktuk, Nunavut, Canada (Gunn and Wobeser, 1993). Similar to other protostrongylids, it requires a gastropod intermediate host (IH) for development in the environment (Hoberg et al., 1995; Kutz et al., 1999).

Transmission of angiostrongylids and protostrongylids typically occurs when the definitive host (DH) ingests gastropods containing infective third-stage larvae (L3) (Anderson, 1992). For some species within the Angiostrongylidae and the related Protostrongylidae, however, L3 emerge from the live IH and transmission may occur when these emerged L3 are ingested by the DH (Heyneman and Lim, 1967; Boev, 1975). Notably, among the Angiostrongylidae, the L3 of *Angiostrongylus cantonensis* and *Angiostrongylus costaricensis* are known to emerge from their IH (Heyneman and Lim, 1967; Ubelaker et al., 1980), a phenomenon considered important for transmission.

An extensive European and Russian literature on the protostrongylids, summarized by Boev (1975), includes the majority of the reports on L3 emergence. Among the Muelleriinae and Protostrongylinae, emergence of L3 from living gastropods has been reported for *Muellerius capillaris*, *Cystocaulus ocreatus*, *Protostrongylus rufescens*, *Protostrongylus davtyani*, *Protostrongylus tauricus*, and *Protostrongylus pulmonalis* and/or *Protostrongylus kamenskyi* (see Boev, 1975), *Protostrongylus stilesi* (see Monson and Post, 1972), and *Protostrongylus boughtoni* (see Kralka and Samuel, 1984). There have been no reports of L3 emergence among the Elaphostrongylinae (Kontrimavichus et al., 1976; Anderson, 1992), but emergence of L3 of *Parelaphostrongylus* sp. (originating from Dall's sheep [*Ovis dalli*]) from the slug *Deroceras laeve* has been observed in the laboratory (S. J. Kutz, unpubl. obs.).

Experimentally, emerged L3 of *P. boughtoni* and *A. cantonensis* produce patent infections in their respective DH (Hey-

neman and Lim, 1967; Richards and Merritt, 1967; Kralka and Samuel, 1984), and emerged L3 of *U. pallikuukensis* can develop to adults in muskoxen (Kutz et al., 1999). Larval emergence is considered important in the epidemiology of *M. capillaris*, *C. ocreatus*, and *A. cantonensis* (Heyneman and Lim, 1967; Boev, 1975). Other authors, however, question the significance of larval emergence and believe the main route of infection for the DH is by ingestion of infected gastropods (Kralka and Samuel, 1984; Anderson, 1992). Patterns of emergence for larval protostrongylids from IH, factors affecting emergence, and its epidemiological significance have not been investigated in detail.

The objectives of the present study were: (1) to investigate in the laboratory the patterns of L3 emergence of *U. pallikuukensis* from 2 potential natural and 1 laboratory IH; and (2) to determine whether L3 emerge from *D. laeve* under natural conditions in the Arctic. We propose that emergence of L3 of *U. pallikuukensis* is a natural phenomenon that occurs consistently from different IH species, that it is not a function of infection intensity, and that it may be epidemiologically important in the maintenance of this parasite in muskox populations.

### MATERIALS AND METHODS

#### Laboratory experiments

**Infection of gastropods:** First-stage larvae (L1) were obtained from the feces of an experimentally infected muskox (Kutz et al., 1999). *Deroceras laeve* and *Catinella* sp., both potential natural IH, were collected near the Coppermine and Rae rivers near Kugluktuk (67°49'N, 115°08'W), Nunavut, Canada and *Deroceras reticulatum*, a laboratory IH, were obtained from a laboratory colony (Kutz et al., 1999). For *D. laeve* and *D. reticulatum*, only adult specimens were used ( $\geq 15$  mg and 400 mg, respectively), but because few *Catinella* sp. were available, specimens with both 2 and 3 whorls were used. Prior to experimental infection with *U. pallikuukensis*, the feet of all wild-caught gastropods were examined for lesions indicative of existing protostrongylid infections (S. J. Kutz, unpubl. obs.). No such lesions were seen on any of the specimens examined.

Gastropods were exposed in groups to L1 on filter paper in plastic petri dishes over a single 3-hr period (Hoberg et al., 1995; Kutz et al., 1999). Additional gastropods, not exposed to L1, were used as controls in the experiments with *D. laeve* and *Catinella* sp.. Four experiments were established at 2 different levels of L1 exposure: in experiments 1 (*D. laeve*) and 2 (*D. reticulatum*) slugs were each exposed to 1,000 L1/slug; in experiments 3 (*D. laeve*) and 4 (*Catinella* sp.) gastropods were each exposed to 500 L1/gastropod (Table I).

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TABLE I. Emergence of L3 of *Umingmakstrongylus pallikuukensis* from 3 species of gastropods under laboratory conditions.

Experiment	Gastropod species	Number of gastropods infected/control	L1/gastropod*	Temperature (°C)	Days of experiment (PI)	Days between examination	Range of L3/gastropod
1†	<i>Deroceras laeve</i>	8/8	1,000	20 ± 0.5	12–60	2	38–179
2†	<i>Deroceras reticulatum</i>	8/0	1,000	20 ± 0.5	14–58	2–5	20–65
3‡	<i>Deroceras laeve</i>	10/10	500	21 ± 1.0	20–60	2–4	1–19
4‡	<i>Catinnella</i>	5/6	500	21 ± 1.0	24–60	2–8	2–12

\* Number of L1 to which each gastropod was exposed.

† In total darkness in temperature-controlled incubator.

‡ In 12–14 hr light/day at room temperature.

**Gastropod maintenance and examination:** Gastropods were housed initially in Rubbermaid® containers (Kutz et al., 1999). At 12–24 days postinfection (PI) (see Table I), they were placed individually in moistened glass petri dishes (50 mm inside diameter) containing lettuce, carrots, and chalk. Dishes and gastropods were examined for L3 frequently (see Table I). Foot lesions and emerging L3 were detected by examining the foot surface using a dissecting microscope with overhead lighting. Gastropods were then transferred to new sterilized petri dishes with fresh food. The uneaten food was removed from the old dishes, water was added to the lids and bases of the dishes, and these were examined with a dissecting microscope. The food removed was placed in individual 100-ml glass containers with approximately 80 ml of tap water. The water in the containers was agitated vigorously in order to dislodge larvae from the food. After 16–24 hr in the light, at 21 ± 1 C, the sediment was examined for larvae. At the end of each experiment, gastropods were killed, digested (Hoberg et al., 1995), and the remaining larvae counted and classified to stage of development.

**Analysis:** It was assumed that all L3 produced in each gastropod were recovered by examination of the dishes and vegetation and by digestion. The total number of L3 for each gastropod was determined by adding the number of L3 that emerged during the experiment to the number recovered by digestion at the end of the experiment. The cumulative proportion of L3 that had emerged from an individual gastropod by a given sampling day was the sum of all the L3 that had emerged up to and including that day divided by the total L3 recovered from the gastropod. The mean cumulative proportion of emerged L3 in an experiment was calculated based only on gastropods from which at least 1 L3 had emerged during the experiment.

### Field experiments

As part of an arctic field study of the development of *U. pallikuukensis*, we investigated L3 emergence under natural conditions. Wild-

caught *D. laeve* were experimentally infected with *U. pallikuukensis* (225 L1/slugs) (Kutz et al., 1999) and placed in enclosures (10 slugs/enclosure) made from perforated plastic pails (150 mm high, 200 mm inside diameter) containing tundra and lined and covered with a fine netting. Enclosures were placed into the tundra so that the vegetation in the pails was at the same level as that outside. Experiments were established every 2 wk, from 19 June to 28 August 1997, in a mesic sedge meadow near Kugluktuk (Table II). Multiple control enclosures containing 10 uninfected slugs were established for each experiment.

Three enclosures were removed from each experiment every 2 wk and gastropods were recovered using a water bath technique (Kralka, 1986) and subsequently digested and examined for larvae. Some enclosures were left in the tundra over winter and were examined the following year. After the first mobile L3 was recovered by digestion from a slug in an experiment, the vegetation from some of the enclosures in that experiment was sampled at varying intervals for emerged L3 (Table II). To recover emerged L3, vegetation was clipped to the level of the soil and placed in a glass flask or jar. Initially (vegetation examined 28 August and 11 September 1997), approximately 250 ml of tap water was added to each container that was then shaken vigorously. The sediment was examined for larvae after 24 hr. The vegetation from 11 September 1997 was re-examined on 12 September 1997 by adding 1 drop of Ivory® liquid dish detergent to 5 L of tap water and using 250 ml of this solution for the larval recovery. The sediment was examined after 24 hr. The detergent technique was used for all subsequent samples. After the vegetation was clipped, slugs were recovered from the remaining turf and examined for larvae by digestion.

### Survival of L3

Live L3 that had emerged from *D. reticulatum* in a preliminary laboratory experiment were placed in 50 ml of tap water (80 L3) or distilled water (164 L3) in 125-ml Erlenmeyer flasks with a foam cork and

TABLE II. Emergence of L3 of *U. pallikuukensis* from *D. laeve* under field conditions.

Experiment	Start date	First L3*	Numbers of emerged L3 recovered from the vegetation				
			28 August 1997†	11 September 1997	18 September 1997	15 June 1998	15 July 1998
1	19 June 1997	17 July 1997	1‡ (1/2)	2‡ (1/3)	—§	0 (0/6)	—
2	3 July 1997	31 July 1997	—	27‡ (3/3)	—	3   (1/4)	—
3	17 July 1997	14 August 1997	—	—	0 (0/3)	5# (1/6)	—
4	31 July 1997	15 July 1998	—	—	—	—	0 (0/3)
5	14 August 1997	15 July 1998	—	—	—	—	0 (0/3)
Control	17 July 1997	NA¶	—	—	—	0 (0/3)	—
Control	31 July 1997	NA	—	—	—	—	0 (0/3)

\* Recovered by digestion of slugs.

† First value is the number of emerged L3 recovered; values in parentheses are numbers of enclosures with emerged L3/total enclosures examined.

‡ All live L3.

§ Not examined.

|| One live L3, 1 dead L3, 1 coiled L3.

# Two live L3, 3 dead L3.

¶ No larvae recovered by digestion.

placed at 0–4 C (water froze at least once during this period). Water was added as needed to prevent dehydration. Larvae were examined at room temperature 13 mo later.

## RESULTS

### Laboratory experiments

**Larval emergence:** Third-stage larvae of *U. pallikuukensis* emerged from all the infected *D. reticulatum* and *Catinella* sp. and all but 1 infected *D. laevis*. Only L3 emerged and these were found on the sediment from the uneaten food and on the inside surfaces of the lids and bases of the petri dishes. Those found in water in the bases of the dishes or in the sediment from the food were usually motile, whereas those found in dry areas of the dishes were tightly coiled, coated with mucus, and adhered to the dish surface. When dislodged and placed in water, these larvae became motile. Early in the experiments, a few emerged L3 were still within the second-stage cuticular sheaths (CS2), but most sheaths were broken at the anterior end. CS2 were frequently observed free in the dishes. After day 30 PI, emerged L3 were usually free of the CS2. Larvae were not recovered from any control experiments with uninfected gastropods.

L3 were observed emerging during examination of the feet of live *D. laevis*. In experiment 3, on day 28 PI, a partially coiled L3 within the CS2 (broken at the anterior end) was expelled within a cloud of mucus from the foot of 1 *D. laevis*. This occurred from another *D. laevis* on day 40 PI in experiment 1. Larvae were also observed actively emerging from slugs of this experiment on days 40 PI (2 L3 from 1 slug) and 46 PI (5 L3 from 1 slug). The cephalic extremities of the larvae were free from the foot of the slug and moving vigorously, whereas the posterior portions were still within the tissue. Dark pigment was visible in areas of the slug from which the larvae were emerging.

**Emergence patterns:** L3 emerged from *D. laevis* (experiments 1 and 3) were first observed on day 22 PI. Emerged L3 were first observed from *D. reticulatum* (experiment 2) on day 20 PI and from *Catinella* sp. (experiment 4) on day 38 PI. In experiments 3 and 4, however, examinations for L3 emergence were not started until days 20 and 26 PI, respectively, and L3 emerging earlier would not have been detected.

The pattern of emergence of L3 from *D. laevis*, *D. reticulatum*, and *Catinella* sp. differed among the experiments (Figs. 1–4). The total number of foot lesions observed in *D. laevis* in experiments 1 and 3 varied between days and individuals, but there was a trend for the lesion count to decrease as the proportion of emerged L3 increased (Fig. 1).

The total cumulative proportions of L3 that emerged from individual gastropods in each experiment relative to the intensities of infection are shown in Figures 5–8. There was 100% L3 emergence from some *D. laevis* with both high (64 L3) and low (2 L3) intensities of infection (Figs. 5, 7). By the end of experiments with *D. laevis* and *D. reticulatum* the majority of the L3 had emerged from more than half of the slugs (Figs. 5–7). Levels of L3 emergence were lower in *Catinella* sp. with only 1 snail having lost more than half of its L3 by day 60 PI (Fig. 8). The patterns of L3 emergence from 6 individual *D. laevis* (from experiments 1 and 3) with similar cumulative proportions of emergence by the end of the experiments (approx-

imately 0.6) were similar despite different intensities of infection (Fig. 9). Emergence at lower intensities of infection appears irregular because each L3 represents a higher proportion of the total than with higher intensities.

### Field experiments

Varying numbers of slugs (0–10) were recovered from the enclosures at each sampling period. All emerged L3 recovered from the vegetation in 1997 were alive, whereas some in 1998 were dead or coiled and nonmotile (Table II). More emerged L3 were recovered from the vegetation using the detergent technique after the water technique. In experiment 2, on 11 September 1997, 7, 1, and 1 live L3 were recovered from the vegetation of each of 3 enclosures examined using tap water. This vegetation was processed again using the detergent technique, and 3, 1, and 14 additional L3 were recovered from the 3 enclosures. In 1997, infected *D. laevis* were recovered from all enclosures with L3 on the vegetation, but in 1998 slugs were not recovered from the enclosures with L3 on the vegetation.

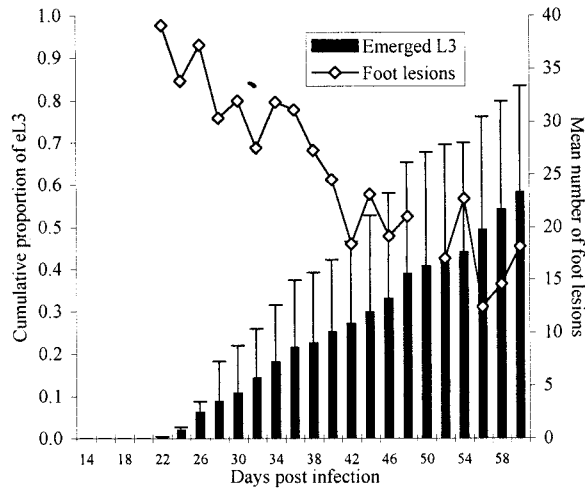
### Survival of L3

Many of the emerged L3 held in water at 0–4 C for over 1 yr were still alive. From the tap water, 21 of the original 80 L3 were recovered, and of these, 1 was dead, 3 were active, and 17 were tightly coiled, the cuticle was intact, and there was no evidence of distortion or cellular deterioration. From the distilled water, 122 of the original 164 L3 were recovered, and of these, 7 were dead, 9 were active, and 106 were tightly coiled and appeared similar to the coiled L3 from tap water.

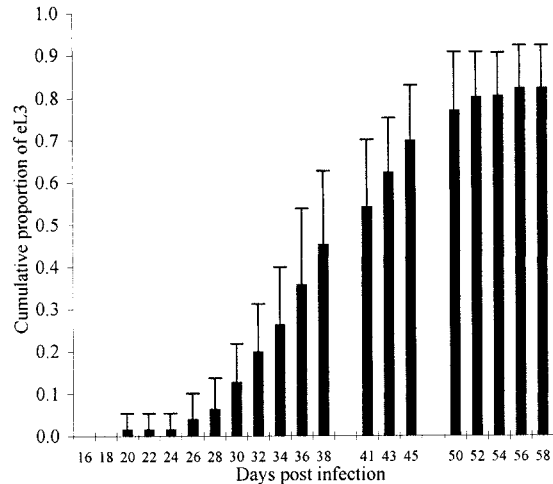
## DISCUSSION

In the laboratory, L3 of *U. pallikuukensis* emerged from 3 species of gastropods, 2 of which occur in the Arctic where naturally infected muskoxen are found. The start of emergence did not coincide with initial appearance of L3 in the gastropods. At 20 C, L1 of *U. pallikuukensis* develop to motile L3 by days 12 and 15 PI in *D. laevis* and *D. reticulatum*, respectively (S. J. Kutz, unpubl. obs.). In the present study, however, L3 did not emerge from these IH until days 20–22 (*D. laevis*) or 18–20 PI (*D. reticulatum*). This suggests that L3 need to reach a particular stage of development before emerging. Morphological and morphometric differences between the early motile L3 in IH and those recovered 3 wk later from the IH or those that had emerged (S. J. Kutz, unpubl. obs.) support this suggestion.

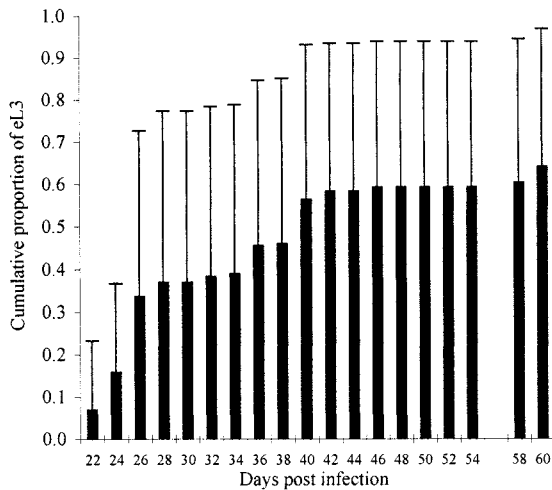
Differences in levels and patterns of emergence were observed among gastropod species. All the L3 had emerged from some *D. laevis* and *Catinella* sp. by day 60 PI, but no *D. reticulatum* had lost all their L3 by day 58 PI (Fig. 6). This difference may be related to the gastropod species. *Deroceras reticulatum*, unlike *D. laevis* and *Catinella* sp., does not occur in the Arctic, nor is it native to North America (Pilsbry, 1948). It is, therefore, an unlikely natural IH for *U. pallikuukensis*. Alternatively, the differences may be an artifact of the small number of gastropods in each experiment. The mean cumulative proportion of L3 emerging from *Catinella* sp. was considerably less than that from either of the slug species. Many of these snails underwent short periods of aestivation during the exper-



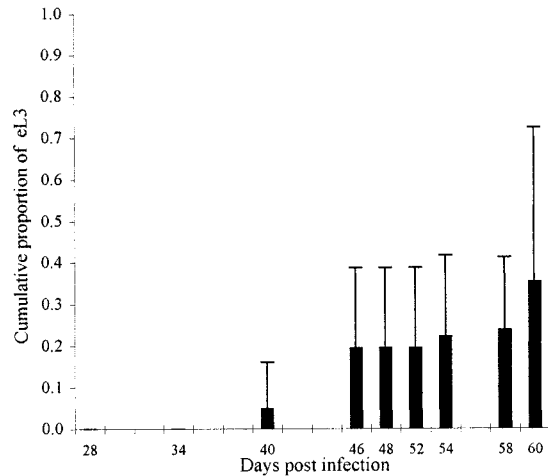
1



2



3



4

FIGURES 1–4. The mean cumulative proportions of third-stage larvae (eL3) that emerged from gastropods during each experiment. Each bar represents the mean cumulative proportion of L3 that had emerged up to that day relative to the total L3 (total L3 = L3 emerged by end of experiment + L3 recovered from the gastropod by digestion at the end of experiment). Error bars = 1 SD of the mean of all gastropods (from which larvae emerged) within an experiment. **1.** Experiment 1 with *Deroceras laeve*. Slugs were examined every second day. Also shown are the mean number of lesions in the feet of 8 slugs (not examined on day 50 PI). **2–4.** Gastropods were examined only on the days indicated by numbers. **2.** Experiment 2 with *Deroceras reticulatum*. **3.** Experiment 3 with *D. laeve*. **4.** Experiment 4 with *Catinella* sp.

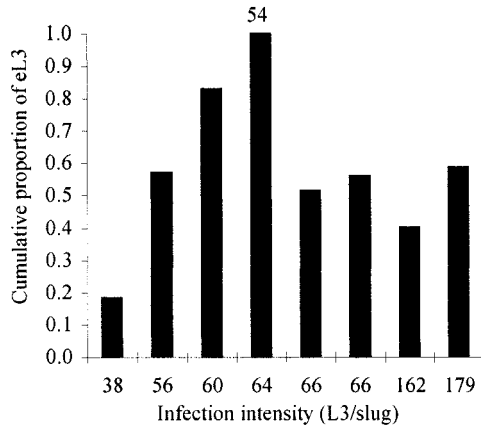
iment, and because IH activity may be necessary for L3 emergence (Boev, 1975), this aestivation may have inhibited emergence. In addition, aestivation can delay development of *Muellerius cf. capillaris* (Solomon et al., 1996) and may have been responsible for the asynchronous development of *U. pallikuukensis* in *Catinella* sp. (at day 60 PI second-stage larvae [L2] as well as L3 were recovered). Fewer larvae at the L3 stage in the snails, compared to *D. laeve*, could result in a lower level of emergence.

Some authors suggest that L3 emergence may be a result of unnaturally high intensities in laboratory infections (Kralka and Samuel, 1984). Gastropods naturally infected with protostrongylids typically have low intensities of infection, but occasionally higher intensities do occur (for example, 75 and 97 larvae of *P. tenuis* in individual *D. laeve*; Lankester and Anderson,

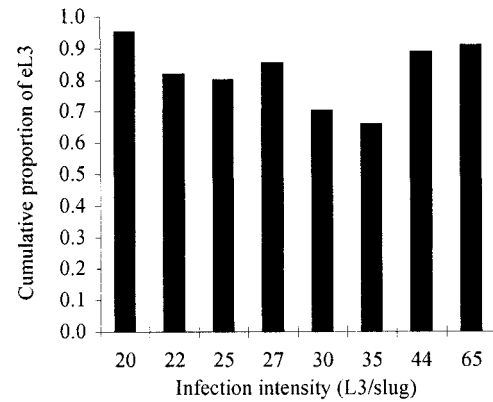
1968; Maze and Johnstone, 1986). Data are not available for the intensity of natural infections of IH with *U. pallikuukensis*; however, in the present study, L3 emerged over a wide range of intensities of infection and there was no evidence that the different intensities affected the cumulative proportion or rate of emergence (Figs. 5–7, 9).

The L3 found on the vegetation in the enclosures of the field experiments may have emerged from live or dead slugs; fewer than the original 10 slugs were recovered from the enclosures, suggesting some mortality. Based on the high proportion of L3 that emerged from live IH in the laboratory, it is likely that at least some of the L3 emerged from live slugs in the field. Whether L3 on the vegetation in the spring of 1998 emerged the previous fall or that spring could not be determined. The apparent absence of L3 emergence in field experiments 4 and

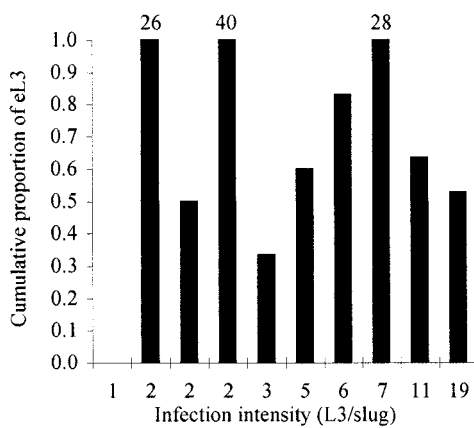




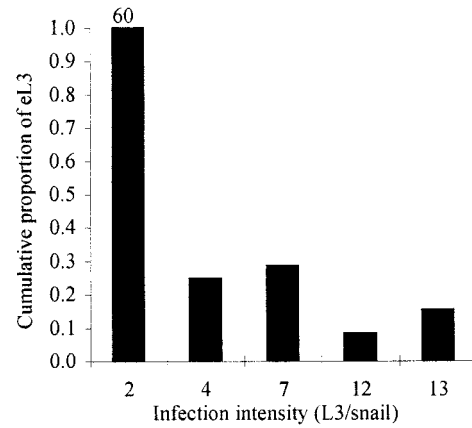
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6



7



8

FIGURES 5–8. Proportion of third-stage larvae (eL3) that emerged relative to intensity of infection. Each bar represents the proportion of the total L3 within an individual gastropod that emerged during the experiment. The total number of L3 (defined in Figs. 1–4) recovered from each individual gastropod is shown on the x-axis. Where numbers are present above the bars they indicate the number of days PI by which 100% of the L3 had emerged from that gastropod. Experiments 1, 3, and 4 ended on day 60 PI; experiment 2 on day 58 PI. 5. Experiment 1 with *D. laeve*. 6. Experiment 2 with *D. reticulatum*. 7. Experiment 3 with *D. laeve*. 8. Experiment 4 with *Catinella* sp.

5 may have been because these larvae had recently developed to L3 (L2 were found in some digested slugs) and were not sufficiently developed to emerge.

The number of L3 that emerged from the gastropods in the field may have been underestimated by our sampling techniques. First, only the vegetation was examined, not the sides of the enclosures or the soil within the enclosures where slugs were frequently observed. Second, the detergent technique seemed to improve larval recovery. It is likely, therefore, that not all L3 were recovered from the vegetation on 28 August 1997, when plain water was used.

Two different mechanisms for emergence of *U. pallikuukensis* from *D. laeve* were observed in the present study, i.e., expulsion of L3 with mucus of the slug and vigorous movements of the L3. These 2 mechanisms may be associated with the presence or absence of the CS2. Expulsion of L3 of *P. boughtoni* contained within the CS2 may be associated with contractions of the gastropod's foot muscle or an active immune response by the IH (Kralka and Samuel, 1984). In contrast,

protostrongylid L3 released from CS2 may have increased motility (Gerichter, 1948; Rose, 1957; Beresford-Jones, 1966) that may facilitate active larval emergence. There appear to be differences among the Protostrongylidae in whether they emerge with or without the CS2 (Monson and Post, 1972; Boev, 1975; Kralka and Samuel, 1984). This and other aspects of L3 emergence may be phylogenetically linked traits that should be investigated within the context of recent phylogenetic studies for the Protostrongylidae (Carreno and Hoberg, 1999).

L3 examined after 13 mo in water at 0–4 C were coiled, showed no signs of degeneration, and were thought to be alive; however, the infectivity was not determined. Survival of L3 in cold water and subsequent infectivity is of significance because the potential terrestrial IH in the Arctic (*D. laeve* and *Catinella* sp.) live in moist habitats, often in mesic or wet sedge meadows, lake shores, or riverbanks (S. J. Kutz, unpubl. obs.). *Deroceas laeve* in particular is well adapted to wet habitats, hatching and remaining mobile when submerged in water (Rollo and Shibata, 1991). In addition, *U. pallikuukensis* can develop in at

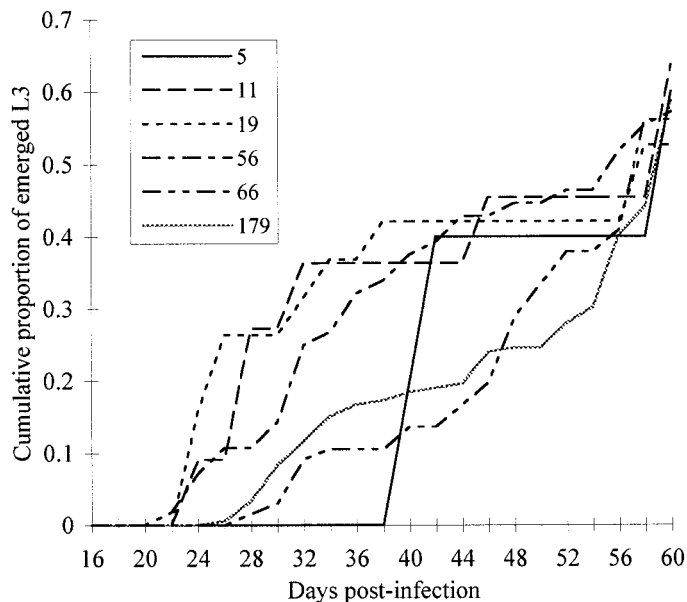


FIGURE 9. Patterns of emergence of L3 from individual *D. laeve* with different intensities of infection (5–179 L3) (slugs from experiments 1 and 3) but similar proportions of emerged L3 by day 60 PI.

least 1 freshwater arctic species, the snail *Aplexa hypnorum* (S. J. Kutz, unpubl. obs.). Larval emergence with subsequent survival in water, perhaps facilitated by the ability to feed when the CS2 is absent, suggest that L3 free in the environment may be a source of infection for muskoxen. Larvae emerging on to dry surfaces may use a different strategy for survival. The thick coating observed on these L3, possibly mucus produced by the gastropods, may be instrumental in protecting the L3 from desiccation and extending its survival in the environment.

Survival and mobility of free-living L3 of other species of protostrongylids have been only superficially investigated. Varying degrees of resistance to desiccation but not to sunlight (*M. capillaris* and *C. ocreatus*), survival after repeated cycles of freezing and thawing associated with overwintering (*C. ocreatus*) (see Boev, 1975), survival in water, and after the death of the IH (*Elaphostrongylus rangiferi*) (see Kontrimavichus et al., 1976) have been reported. In addition, horizontal, but no vertical, mobility of emerged L3 has been reported for *E. rangiferi* (see Kontrimavichus et al., 1976). Understanding the survival, mobility, and subsequent infectivity of emerged *U. pallikuukensis* L3 is important to better assess the significance of larval emergence.

Transmission of *U. pallikuukensis* requires temporal and spatial overlap of L1 with gastropods and, according to the current understanding of protostrongylid life cycles, of gastropods containing infective L3 with the DH (Anderson, 1992). In the Arctic, such transmission of *U. pallikuukensis* and other nematodes to the DH may be limited to seasonally defined periods (Hoberg et al., 1999). The short arctic summers may limit when gastropods are active. Emerged L3 that survive ambient conditions on the vegetation may be available when the gastropods are not and may be more widely distributed in the environment than the gastropods. This enhanced life span and dispersal of emerged L3 would increase their availability on temporal and

spatial scales and may eliminate the necessity for direct gastropod–muskox contact.

Spatial and temporal overlap of muskoxen with gastropods at high densities may be limited by seasonal migrations (Gunn and Fournier, 2000) or seasonal dietary shifts (thus changes in habitat use) by muskoxen (Robus, 1981; O'Brian, 1988; Gunn and Sutherland, 1997). For example, gastropod densities may be greatest in muskox wintering areas and yet their availability to muskoxen would be restricted in the winter. If, however, L3 emerge and survive ambient conditions on the vegetation, they may be available during the winter when gastropods are hibernating but muskoxen are present.

In the present study L3 emerged over an extended time period. A similar pattern in nature could result in L3 emerging on to many different areas of the vegetation during normal summer gastropod activity. If arctic gastropods exhibit similar circadian activity patterns, influenced by temperature, humidity, and wind, as has been reported for temperate species (Dainton, 1989; Kaufmann, 1990; Rollo and Shibata, 1991), they are probably not available to the muskoxen at all times of the day. Larval emergence, however, would provide multiple foci of infection from a single gastropod, on vegetation or in the water, available 24 hr a day, increasing the probability for transmission compared to that dependent on ingestion of the gastropod. This strategy may be particularly important in parasite transmission from heavily infected gastropods.

Larval emergence may be linked to the life history of the gastropods and may increase L3 availability after death of the IH. For example, if in some years, as has been proposed for *E. rangiferi* in Norway (Halvorsen and Skorping, 1982), larvae of *U. pallikuukensis* require 2 summer seasons to develop to L3, it would be the large slugs, infected the previous year, that would contain L3 by the following midsummer. In southern Ontario, Canada, large *D. laeve* (those that have overwintered) disappear from the population in August and September (Lanckester and Anderson, 1968). If a similar life cycle occurs in the Arctic, without emergence, all L3 in these large slugs would be lost in the second summer when the slugs died. Alternatively, L3 emerging on to vegetation or into the water before the IH dies may remain available for transmission. Observations in the present study and by other authors (Rose, 1957; Cheng and Alicata, 1964) suggest that L3 emergence may increase in unhealthy gastropods.

In conclusion, based on laboratory and field data, emergence of L3 from the IH appears to be an inherent trait of *U. pallikuukensis*. This alternate pathway for transmission may be of importance in the natural host–parasite system by increasing larval availability spatially and temporally, perhaps extending or eliminating the narrow and seasonally defined windows of transmission. Recognition that emergence of L3 of *U. pallikuukensis*, and perhaps other protostrongylids, may be a natural and epidemiologically significant phenomenon broadens our current understanding of transmission dynamics and life history patterns for lungworms (e.g., Anderson, 1992). To understand this phenomenon further, and to test the hypotheses outlined in this paper, it is important to determine the extent of L3 emergence in the field, the survival and subsequent infectivity of emerged larvae under natural conditions, the factors affecting emergence, and how the life history and behavioral patterns of

the gastropod intermediate hosts relate to the life history strategies of muskoxen definitive hosts.

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