REFRACTIVENESS OF PHYSA CUBENSIS (PFEIFFER) AND APPLEXA MARMORATA (GUILDING) TO FASCIOLA HEPATICA (L.)

Vigueras and Moreno\(^2\) reported *Physa cubensis* (fig. 1), the most common fresh-water snail in Cuba, to be an intermediate host for the liver-fluke, *Fasciola hepatica*.

*P. cubensis*\(^3\) is very common in ponds, creeks, drainage canals, and rivers in Puerto Rico. Another physid snail, *Aplexa marmorata* (fig. 1), is found occasionally by us in association with *Lymnaea columella* and *Lymnaea cubensis*, the intermediate snail hosts of *F. hepatica* in Puerto Rico.

It is highly important to know if these two physid snails of Puerto Rico serve as intermediate hosts for *F. hepatica*, if control measures are to be instituted against fascioliasis.

The specimens of *P. cubensis* used in this study came from an egg mass laid by an individual snail; those of *A. marmorata* from egg masses laid by four snails. Both species were collected from a small fresh water pond at the Agricultural Experiment Station, Río Piedras. Identification of both species was based on their description by Richards.\(^4\)

*P. cubensis* was exposed to *F. hepatica* infection as follows: 1, 100 1- to 2-day-old snails were introduced in a small (10.5 cm.) finger-bowl which contained about 1,000 miracidia in 100 ml. dechlorinated tap water; 2, 100 21- to 26-day-old snails were put in a small finger-bowl which contained 1,000 miracidia in 200 ml. dechlorinated tap water; 3, 50 one- to 2-month-old snails were put in a small finger-bowl containing 1,000 miracidia in 400 ml. dechlorinated tap water; and 4, 10 1-day-old, 10 26-day-old, 10 1½-month old *P. cubensis* and 10 21-day-old *L. cubensis* as control were placed in individual wells of a complement-fixation plastic tray, each containing 10 miracidia in dechlorinated tap water. The snails were observed under a

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\(^{2}\) Vigueras, I. and Moreno, A., "Physa cubensis" (Mollusca), Un Nuevo Hospedero Intermediario de "Fasciola hepatica" (Trematoda), Memorias de la Sociedad Cubana de Historia Natural "Felipe Poey" 12 (1): 74, 1938.


dissecting microscope for penetration by the miracidia. The exposure lasted for 4 to 4½ hours at a room temperature of 29° to 31° C.

The *P. cubensis* cultures were kept in 12-gallon plastic aquaria, half-filled with rain water and the *L. cubensis* (control) in individual culture baskets as described by de León. The snails used to observe penetration of the miracidia were eliminated after the observation. A piece of decaying *malanga* leaf (*Caladium* sp.) was fed once a week. A second trial was conducted following a similar procedure.

Half the snails of each age-group and all the control snails (*L. cubensis*) in the first and second trials were gently crushed between two glass slides 3 weeks after exposure and their “livers” separated from other tissues and shell fragments. The “livers” then were pressed between a glass slide and a cover slip and examined under the microscope for *F. hepatica* rediae. The remaining snails in both trials were examined similarly 6 weeks after exposure.

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Fig. 1.—*Physa cubensis* Pfeiffer (above) and *Aplexa marmorata* Guilding (below).
No miracidium penetrated *P. cubensis* after more than 4 hours of exposure; almost all the miracidia penetrated into *L. cubensis* in 15 to 30 minutes. No redia was observed in any of the three age-groups of *P. cubensis* exposed to *F. hepatica*. Many rediae were found in almost all the control snails 3 weeks after exposure. These observations indicate that *P. cubensis* is refractive, apparently, to *F. hepatica*.

*A. marmorata* was exposed to *F. hepatica* infection as follows: 1, 40 one- to 2-day-old snails were placed in a small finger-bowl which contained about 250 miracidia in 50 ml. dechlorinated tap water; 2, 30 21- to 26-day-old snails were put in a small finger-bowl which contained 500 miracidia in 100 ml. dechlorinated tap water; 3, 20 2-month-old snails were held in a finger bowl containing 500 miracidia in 100 ml. dechlorinated tap water; and 4, 10 one-day-old and 10 7-day-old *L. cubensis*, as control, were placed in individual wells of a complement-fixation plastic tray, each containing five miracidia in dechlorinated tap water. Penetration of the miracidia into the snails was observed under a dissecting microscope. The exposure lasted 4 to 4½ hours at a room temperature of 29° to 31°C.

*A. marmorata* and *L. cubensis* (control) were kept in individual culture baskets. The snails used to observe penetration of the miracidia were eliminated after the observation.

The snails were dissected for rediae using the same method as in *P. cubensis*.

None of the miracidia penetrated *A. marmorata* after more than 4 hours of exposure; almost all the miracidia penetrated into *L. cubensis* in 15 to 30 minutes. No redia was observed in any of the three age-groups of *A. marmorata* exposed to *F. hepatica*. However, many rediae were found in almost all the control snails 3 weeks after exposure. Probably, *A. marmorata* is also refractive to *F. hepatica*. Furthermore, microscopic examination of the “livers” of 67 *A. marmorata,* collected from a swamp in Canóvanas, failed to reveal *F. hepatica* rediae, whereas 9 percent and 25 percent of *L. columella* and *L. cubensis*, respectively, were infected.

Our results are not in accord with Vigueras and Moreno, who reported that *P. cubensis* is an intermediate host of *F. hepatica* in Cuba. Our negative results are supported by Wright (cited by Taylor) in his statement that all snails serving as intermediate hosts for *F. hepatica* throughout the world belong to a closely-related group belonging to the genus *Lymnaea*. Moreover, as early as 1883, Thomas (cited by Pantelouris) found *P.*
*fontinalis* to be refractory to *F. hepatica*. We exposed large numbers of *P. cubensis* and *A. marmorata* of an inclusive age spectrum to 10 miracidia each, noting that penetration was not apparent. The rediae observed by Vigueras and Moreno\textsuperscript{10} in natural infections might not have been those of *F. hepatica*. However, it also is possible that the Cuban strain of *P. cubensis* is distinctive in being an intermediate host of at least marginal importance.

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\textsuperscript{10} Vigueras and Moreno, *loc. cit.*