

Schistosoma mansoni, Strain NMRI Exposed Biomphalaria glabrata, Strain NMRI

Catalog No. NR-21962

For research use only. Not for human use.

Contributor and Manufacturer:

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Product Description:

Flatworm Classification: Schistosomatidae, Schistosoma

Species: Schistosoma mansoni

Strain: NMRI

Original Source: Schistosoma mansoni (S. mansoni), strain NMRI was isolated in the 1940s from S. mansoni eggs obtained from infected Puerto Rican school children.¹

Host: The Biomphalaria glabrata (B. glabrata), strain NMRI snail host is an albino snail that is highly susceptible to the NMRI strain of S. mansoni. It was derived from a cross between pigmented Puerto Rican (susceptible) snails and Newton's M-line snails.¹

Comment: S. mansoni, strain NMRI was brought to the Naval Medical Research Institute (NMRI) in 1945, where it was maintained until 1971, at which time it was transferred to the Biomedical Research Institute.²

S. mansoni is a species of trematode worm which causes the chronic parasitic disease Schistosomiasis. Worldwide, more than 200 million people are infected and nearly 700 million are at risk, primarily in areas with poor sanitation that lack access to safe drinking water.⁴

Infection occurs through contact with larval-stage schistosomes (cercariae) that are released by freshwater snails. Upon exposure to infested water, these larvae penetrate human skin and travel through blood vessels to the liver where they mature and deposit eggs. Some of these eggs are then passed through human feces into water to re-infect the snail host and continue the parasite's life cycle. Schistosome eggs that remain in the human body cause an immune response and damage to internal organs.⁴

Material Provided:

NR-21962 consists of up to 100 B. glabrata, strain NMRI, infected with S. mansoni, strain NMRI, in a petri dish.¹

Packaging/Storage:

NR-21962 is packed in moist paper towels in a Petri dish encased in bubble-wrap and shipped overnight in cardboard boxes (425 mm x 225 mm x 225 mm). To insure viability, snails should be placed in suitable aquaria at 26°C to 28°C.

Note: Infected snails are exposed to miracidia only a few days before shipment. Laboratories should be aware and take caution of possible cercarial exposure following a 4 week prepatent period after exposure to miracidia (depending on the temperature and species). BEI Resources recommends consulting personnel in your institution's Occupational Safety and Health office for clearance to receive and maintain the infected snails.

Growth Conditions:^{5,6}

Food Source:

Romaine lettuce or Nostoc blue-green algae grown on a layer of autoclaved mud may be used as food sources (see Appendix I for details).

Maintenance of B. glabrata:

Depending on the number of snails to be maintained, they may be kept in plastic trays (approximately 1.5 L volume) or larger aquaria (40 to 110 L) in filtered tap water that has been aerated for 2 to 3 days (pH ~ 7.1). The ambient temperature should be 26°C to 28°C, and the aquaria should be under continuous illumination from a 40-watt fluorescent bulb.

Propagation of B. glabrata:

1. Place ten snails in plastic trays containing aerated tap water and Styrofoam (75 mm x 120 mm). Maintain trays at 26°C to 28°C under continuous illumination.
2. At the end of one week, remove egg masses from the trays and Styrofoam and transfer to Petri dishes that contain aerated tap water.
3. After one week, 50 newborn snails 0.6 mm to 0.8 mm in shell diameter are transferred with a pipette to each Petri dish that contains water, mud and blue-green algae.
4. Each week, transfer and maintain snails at a density of 25 snails per dish. After 14 days, 3 mm to 5 mm shell diameter snails may be infected with S. mansoni or used to initiate new cultures.

Collecting Cercariae:

1. Screen infected snails individually under a dissecting microscope in a Petri dish containing a small volume of aerated tap water for the presence of Schistosoma sporocysts, which appear as light colored flecks that are visible through the shell.
2. Place infected (patent) snails in a beaker where moisture is maintained but does not contain water. Let beaker stand for 15 to 30 minutes at 26°C to 28°C under ceiling illumination.
3. Rinse snails with a small volume of water. Add a second volume of water (100 mL per 100 snails). Place uncovered beaker under a 15-watt white fluorescent light, 18 inches above the beaker for 20 minutes. Decant water containing cercariae into another beaker. On average, 2000 to 4000 cercariae may be obtained from each infected snail.
4. To estimate the cercarial density, stir the beaker that contains cercariae continuously with a magnetic stir bar, and remove 0.1 mL aliquots from the center of the suspension.
5. Place each aliquot in a 10-cell depression slide. Add

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one drop of Lugol's Iodine solution to each cell prior to counting under a dissecting microscope. Determine the cercariae density per volume. This suspension may subsequently be diluted, or concentrated by pipetting out the excess liquid.

Citation:

Acknowledgment for publications should read "The following reagent was obtained through BEI Resources, NIAID, NIH: *Schistosoma mansoni*, Strain NMRI Exposed *Biomphalaria glabrata*, Strain NMRI, NR-21962."

Biosafety Level: 2

Appropriate safety procedures should always be used with this material. Laboratory safety is discussed in the following publication: U.S. Department of Health and Human Services, Public Health Service, Centers for Disease Control and Prevention, and National Institutes of Health. Biosafety in Microbiological and Biomedical Laboratories. 5th ed. Washington, DC: U.S. Government Printing Office, 2009; see <http://www.cdc.gov/biosafety/publications/bmbl5/index.htm>.

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References:

1. F. A. Lewis, Personal Communication.
2. Lewis, F. A., et. al. "Large-Scale Laboratory Maintenance of *Schistosoma mansoni*, with Observations on Three Schistosome/Snail Host Combinations." J. Parasitol. 72 (1986): 813-829. PubMed: 3546654.
3. Carvalho E. M. and A. A. M. Lima. "Schistosomiasis (Bilharziasis)." In: Goldman L, Ausiello D, eds. Cecil Medicine. 23rd ed. Philadelphia, Pa: Saunders Elsevier; 2007: Chap 376.
5. Lewis, F. "Schistosomiasis." Curr. Protoc. Immunol. Chapter 19 (2001): Unit 19.1. PubMed: 18432750.
6. Bruce, J. I. and Y.-S. Liang. "Cultivation of Schistosomes and Snails for Researchers in the United States of America and Other Countries." J. Med. Appl. Malacol. 4 (1992): 13-30.

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APPENDIX I: PREPARATION OF MUD-BASED MEDIA FOR GROWTH OF *NOSTOC* ALGAE⁵Equipment

Autoclave

Spatula(s)

Materials

Mud, or soil source

Chicken manure

Calcium carbonate (pulverized limestone)

Clay

0.06% sodium nitrate solution prepared with aged tap water

Nostoc (stock cultures can be obtained from Ward's Biological Supply, Rochester, NY)

Plastic Petri dishes (25 mm x 100 mm)

Stainless steel baking pan (250 mm x 375 mm x 75 mm)

Procedure

1. The proportions of dried mud, lime and chicken manure needed for good growth of *Nostoc* will likely vary, depending on the richness of the soil obtained. A small amount of clay may be necessary for cohesion of the mud mound that will be placed in the Petri dishes. The following describes the current proportions of each component for the soil. Trial and error will be the rule, rather than the exception, to accommodate the apparent richness [or lack thereof] of soils in different regions.
2. The soil and site chosen ideally should be one where there is considerable sedimentation (e.g. a stream bed bottom) or topsoil. Soil should be obtained where no known herbicides or pesticides have been used.
3. The mud or soil brought back to the laboratory from the field site should be strained through a series of crude screens to remove rocks and other large debris. Once it is a fine consistency, it should be completely dried before use.
4. Mix 3 kilograms of dried mud with 90 grams lime (pulverized limestone), and 30 grams dried chicken manure. To this mixture add enough tap water to make a paste. Place the mud mixture in a large stainless steel baking pan and cover with aluminum foil. The depth of the mud should be no more than about 100 mm. Autoclave for a continuous 2 hours.
5. Once the mud is autoclaved and cooled to room temperature, use a sterile spatula (spatulas should be wiped down periodically with gauze drenched in pure alcohol) and place about 40 grams of the still wet mud in the center of a Petri dish and form a smooth and solid mud mound about 15 mm high and 60 mm in diameter. If the mud has dried too much during autoclaving and needs some additional liquid to make it easier to spread, add a few mL of sterile 0.06% nitrate solution and mix thoroughly. To expedite the spreading process, one can use two curved sterile spatulas to stir a third to half of the mud in the steel pan (adding the sterile 0.06% nitrate solution as needed) before spreading it into the Petri dishes. This ensures consistency of the ingredients in the mud that is placed in each Petri dish.
6. Once the mud mounds have been formed in the Petri dishes, cover the mud mound with 0.06% nitrate solution and add about 2 mL of a suspension of *Nostoc* (in 0.06% nitrate solution) to seed the plate for new growth. Be sure not to flood the Petri dish with liquid, so that the lid does not become wet with the growth medium.
7. Cover and place under fluorescent lighting (40 watt, cool-white fluorescent) at 25°C to 27°C for 1 to 3 weeks. For best results the lights should be about 1 foot above the Petri dishes.
8. The preparation is suitable for feeding to the snails once a solid mat of the *Nostoc* has grown over the surface.