**Product Information Sheet for NR-34793**

**Schistosoma mansoni**, Strain PR-1, Exposed Golden Syrian LVG Hamsters

**Catalog No. NR-34793**

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: PR-1
Host: Mesocricetus auratus (Golden Syrian LVG Hamster)
Original Source: Schistosoma mansoni (S. mansoni), strain PR-1 was collected from infected snails in Arecibo, Puerto Rico in 1950.1
Comment: Strain PR-1 was maintained by NIH/NIAID until 1978, when it was brought 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.2

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-infest 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.2

**Material Provided:**
NR-34793 consists of male Golden Syrian LVG hamsters from Charles River Laboratory that have been exposed to S. mansoni, strain PR-1.

**Packaging/Storage:**
S. mansoni, strain PR-1, exposed Golden Syrian LVG hamsters are placed in transfer cages with adequate food and water source and shipped overnight. Upon arrival they should be immediately placed in cages at the recipient institute’s animal facility.

**Collection of Schistosoma miracidia:**
1. Euthanize hamster by intraperitoneal injection of 0.3 mL sodium pentobarbital (65 mg/mL) with heparin (10000 units/mL).
2. Remove liver and small and large intestines. Rinse tissues in 1.2% NaCl. If using intestines, remove and wash with 1.2% NaCl repeatedly.
3. Blend liver/intestines in filtered tap water that has been aerated for 2 to 3 days (conditioned water) for 20 seconds in a Waring blender. Centrifuge homogenate for 5 minutes (300 x g) at room temperature.
4. Pour off supernatant. Add 5 mL conditioned water and shake tube vigorously for several seconds. Dilute suspension at least 100-fold in conditioned water. For optimal hatching, use conditioned water between 26°C and 28°C.
5. Place suspension in darkened side-arm flask. Make sure that water fills the unpainted sidearm.
6. Direct a light source at exposed unpainted part of side arm. Miracidia will swim to this area after hatching and collect within the unpainted side arm within 10 to 20 minutes.
7. Remove miracidia from the side arm using a fine-tipped Pasteur pipette and place in a Petri dish that contains conditioned water.

**Citation:**
Acknowledgment for publications should read “The following reagent was obtained through BEI Resources, NIAID, NIH: Schistosoma mansoni, Strain PR-1, Exposed Golden Syrian LVG Hamsters, NR-34793”.

**Biosafety Level:**
1


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References:

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