

**Oligo(dT) Generated Complementary DNA from *Schistosoma haematobium*, Egyptian Strain, Miracidia**

**Catalog No. NR-49833**

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**Product Description:** Complementary DNA (cDNA) was synthesized from total RNA extracted from *Schistosoma haematobium*, Egyptian strain, miracidia, using the ProtoScript<sup>®</sup> II First Strand cDNA Synthesis Kit (New England BioLabs<sup>®</sup>). The kit provides an anchored oligo-[d(T)<sub>23</sub>VN] primer which forces the primer to anneal to the beginning of the polyA tail increasing the yield of 3' end poly(A)-primed cDNAs.

**Lot<sup>1,2</sup>: 63725308**

**Manufacturing Date: 10JUN2015**

TEST	SPECIFICATIONS	RESULTS
Concentration	Report results	1.0 µg in 20 µL per vial (0.05 µg/µL)
OD <sub>260</sub> /OD <sub>280</sub> Ratio	1.70 to 2.00	1.80
Qualification by RT-PCR Amplification of 28S ribosomal RNA gene <sup>3</sup>	~ 290 base pair amplicon	~ 290 base pair amplicon (Figure 1)

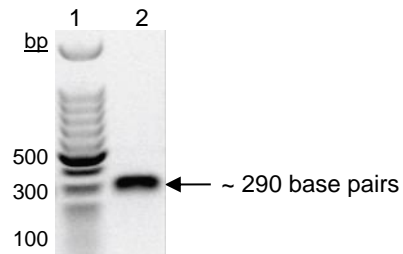
<sup>1</sup>QC testing was performed by the Biomedical Research Institute, Rockville, Maryland (NIH-NIAID Contract HHSN272201000005I).

<sup>2</sup>cDNA was synthesized from total RNA by ProtoScript<sup>®</sup> II First Strand cDNA Synthesis Kit (New England BioLabs<sup>®</sup>, Inc.) according to the manufacturer's instructions, using Oligo d(T)<sub>23</sub>VN.

<sup>3</sup>Primers were designed to amplify the nucleotide region 39 to 326 of the *Schistosoma mansoni* (*S. mansoni*) 28S ribosomal RNA gene (GenBank: Z46503.1). Cross-amplification of the 28S gene from other *Schistosoma* species has been observed.

**Figure 1: Amplification of 28S Ribosomal RNA Gene for Qualification by RT-PCR**

Lane 1: 100 base pair ladder  
Lane 2: ~ 290 base pair amplicon from *S. mansoni* 28S ribosomal RNA gene



**Date:** 08 JAN 2016

**Signature:** 

BEI Resources Authentication

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