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## Transcriptome analysis of *Schistosoma mansoni* larval development using serial analysis of gene expression (SAGE)

A. S. TAFT<sup>1,†</sup>, J. J. VERMEIRE<sup>1,§,†</sup>, J. BERNIER<sup>2</sup>, S.R. BIRKELAND<sup>2</sup>, M. J. CIPRIANO<sup>2</sup>, A. R. PAPA<sup>2</sup>, A.G. MCARTHUR<sup>2</sup>, and T. P. YOSHINO<sup>1,\*</sup>

<sup>1</sup>Department of Pathobiological Sciences, School of Veterinary Medicine, University of Wisconsin-Madison, 2115 Observatory Drive, Madison, WI, USA

<sup>2</sup>Josephine Bay Paul Center for Comparative Molecular Biology and Evolution, Marine Biological Laboratory, Woods Hole, MA, USA

### SUMMARY

Infection of the snail, *Biomphalaria glabrata*, by the free-swimming miracidial stage of the human blood fluke, *Schistosoma mansoni*, and its subsequent development to the parasitic sporocyst stage is critical to establishment of viable infections and continued human transmission. We performed a genome-wide expression analysis of the *S. mansoni* miracidia and developing sporocyst using Long Serial Analysis of Gene Expression (LongSAGE). Five cDNA libraries were constructed from miracidia and *in vitro* cultured 6- and 20-day-old sporocysts maintained in sporocyst medium (SM) or in SM conditioned by previous cultivation with cells of the *B. glabrata* embryonic (Bge) cell line. We generated 21 440 SAGE tags and mapped 13 381 to the *S. mansoni* gene predictions (v4.0e) either by estimating theoretical 3' UTR lengths or using existing 3' EST sequence data. Overall, 432 transcripts were found to be differentially expressed amongst all 5 libraries. In total, 172 tags were differentially expressed between miracidia and 6-day conditioned sporocysts and 152 were differentially expressed between miracidia and 6-day unconditioned sporocysts. In addition, 53 and 45 tags, respectively, were differentially expressed in 6-day and 20-day cultured sporocysts, due to the effects of exposure to Bge cell-conditioned medium.

### Keywords

*Schistosoma mansoni*; *Biomphalaria glabrata*; SAGE; miracidia; sporocyst; development

### INTRODUCTION

Schistosomiasis is a major public health problem, mainly in the tropics, with an estimated 200 million individuals infected and 650 million living in endemic areas (<http://www.who.int/schistosomiasis/en/>). The causative agents are digenetic parasitic flatworms of the genus *Schistosoma*, which have a complex life cycle involving 2 obligate hosts ; a

\*Corresponding author: Department of Pathobiological Sciences, School of Veterinary Medicine, University of Wisconsin-Madison, 2115 Observatory Drive, Madison, WI, 53726, USA. Tel: +608 263 6002. Fax: +608 265 8122. yoshinot@svm.vetmed.wisc.edu.

§Current address: Department of Pediatrics, Yale University School of Medicine, New Haven, CT, USA.

†These authors contributed equally to this work.

mammalian definitive host (human) and snail intermediate host. Adult male and female worms, depending on the species, inhabit the vasculature of the urinary plexus or mesenteric venules surrounding the large intestine, where they produce eggs that escape to the environment in urine or feces. In freshwater, free-swimming miracidia contained within eggs hatch and, upon finding a suitable snail intermediate host, directly penetrate the snail mantle to initiate an infection (Basch, 1991). Post-penetration, the miracidium sheds its ciliary epidermal plates, during which time a tegumental syncytium covers the entire larval surface as it transforms to the parasitic primary or mother sporocyst stage (Basch and DiConza, 1974). This developmental transition from free-living to parasitic state within the snail host is crucial to successful establishment of infections, and is presumed to involve dramatic physiological changes both at the biochemical and molecular levels. However, to date few studies have focused on the molecular basis of larval transformation or the identification of genes regulating subsequent asexual development of and embryogenesis within subsequent sporocyst generations.

Previously, high-throughput cDNA or oligonucleotide DNA microarrays have been used to compare the expression of thousands of genes in a variety of *S. mansoni* stages, including adult male and female worms (Hoffman *et al.* 2002; Fitzpatrick *et al.* 2005; Verjovski-Almeida *et al.* 2007; Waisberg *et al.* 2007) as well as larval stages, including daughter sporocysts and cercariae (Jolly *et al.* 2007). In a recent study Vermeire *et al.* (2006) identified a large number of gene expression changes in *S. mansoni* miracidia compared to 6-day *in vitro* cultured sporocysts using a DNA microarray (MA) spotted with oligonucleotide probes corresponding to ESTs or full-length mRNAs derived mainly from adult worms, eggs, and cercariae, with <5% originating from miracidia or sporocyst sequences. Despite this over-representation by cercarial and adult genes, approximately 60% of the array probes, representing individual mRNA transcripts, were expressed in miracidia and/or sporocysts, with a significant number being differentially expressed between these stages (Vermeire *et al.* 2006). Considerable overlap in transcribed genes thus exists between stages. However, microarrays are limited to analysing only previously identified transcripts. In this way they constitute a 'closed' gene expression profiling platform, limited to predetermined or known sets of genes.

Serial Analysis of Gene Expression (SAGE) is a sequence based gene-expression profiling tool that can be utilized to generate quantitative transcriptional profiles of genes in an organism. In SAGE, a short sequence tag from a unique position of each mRNA molecule is used to uniquely identify the source gene from within the genome (Velculescu *et al.* 1995). Sequence tags are isolated from an mRNA sample and are linked together to form long concatenated molecules that are cloned and sequenced. The population of tags defines patterns of expression of individual genes. Quantification of all tags provides a relative measure of gene expression (i.e., mRNA abundance). SAGE thus provides both the identity of expressed genes and levels of their expression. SAGE constitutes an 'open' platform, providing a rapid and comprehensive approach for elucidation of quantitative gene expression patterns not dependent upon prior availability of transcript information. In addition, the sequences generated can be used to identify previously unknown genes through the application of tag-based reverse transcription- polymerase chain reaction (RT-PCR), i.e.,

use of tag sequences to design primers for amplifying unknown cDNA sequences, leading to gene identification and elucidation of function.

In preliminary experiments involving gene expression patterns across the entire *S. mansoni* life cycle, SAGE has demonstrated excellent potential for stage-associated gene profiling (Williams *et al.* 2007). Recently, SAGE has been used to identify transcriptional changes in adult worms in response to nitric oxide exposure (Messerli *et al.* 2006). Ojopi *et al.* (2007) used SAGE to identify expressed transcripts in pooled adult male and female worms, but did not identify any stage-related changes in transcript levels. The SAGE tags used by Ojopi *et al.* (2007) were only 14 bp long, potentially leading to errors in data analysis ; specifically the problem of a single tag matching multiple transcripts.

In the present study, LongSAGE was used to compare gene expression profiles for *S. mansoni* miracidia, 6-day *in vitro* cultured primary sporocysts and 20-day *in vitro* cultured sporocysts to quantitatively assess commonly- and differentially-expressed genes during early larval development of this parasite. LongSAGE is a highly specific quantitative method of gene expression profiling that generates 21 bp tags, of which theoretical modelling predicts that >99.8% are expected to match only once to a human-sized genome (Saha *et al.* 2002). We combined SAGE data with the gene predictions/ annotations of the *S. mansoni* genome (v4.0e) and modelled theoretical 3' UTR lengths for genes without 3' EST sequence data to generate the most up-to-date analysis of the *S. mansoni* larval transcriptome during establishment of intramolluscan infections. The 6- and 20-day time-points were chosen to determine transcriptional changes associated with 2 very important developmental time-points in early larval development. After 6-days of *in vitro* culture the miracidia have all transformed and are transitioning from a free-living to parasitic stage and at 20-days the *in vitro*-cultured sporocysts are beginning to form brood chambers to produce the second generation of daughter/secondary sporocysts. In addition, we also investigated the effects of sporocyst exposure to snail Bge cell products, predicted to enhance larval growth/embryogenesis on gene expression (Yoshino and Laursen, 1995).

## MATERIALS AND METHODS

### Parasite culture

*Schistosoma mansoni* (NMRI strain) eggs were recovered from the livers of mice at 7–8 weeks post-infection as described by Yoshino and Laursen (1995). Miracidia were hatched from the eggs in sterile artificial pond water and concentrated on ice in conical polypropylene centrifuge tubes. Miracidia were isolated at 15-min intervals over a 2 h period. Cold-immobilized miracidia were either centrifuged for 1 min at 500 g and immediately harvested for total RNA or were pooled and transferred to a 24- well culture plate to permit transformation into primary sporocysts followed by culturing for 6 or 20 days under normoxic conditions at 26 °C in either *S. mansoni* sporocyst medium (SM; Ivanchenko *et al.* 1999) or SM previously conditioned with *Biomphalaria glabrata* embryonic (Bge) cells. Conditioned SM was used in these larval SAGE studies to determine the effect of snail-derived components on sporocyst gene expression. Control (unconditioned) and conditioned media inwells containing sporocysts were changed at 2-day intervals. At 6-day (6d) and 20-day (20d) developmental time-points total RNA from all

cultured sporocysts (6d control, 6d conditioned, 20d control and 20d conditioned) was isolated using TRIzol<sup>®</sup> reagent (Invitrogen, Carlsbad, CA) according to the manufacturer's instructions.

### **Bge cell culture and production of conditioned media**

The *Biomphalaria glabrata* embryonic (Bge) cell line (ATCC CRL 1494) was used to produce snail cell-conditioned sporocyst medium (SM) for use in sporocyst *in vitro* culture experiments. Bge cells were maintained in 250-ml tissue culture flasks (Falcon<sup>™</sup>, BD Biosciences, San Jose, CA) containing Bge medium (Hansen, 1976) supplemented with heat-inactivated 10% fetal bovine serum (cBge), penicillin G (0.06 mg/ml) and streptomycin sulfate (0.05 mg/ml) at 26 °C under normal atmospheric conditions. Bge cells were grown to confluence, washed once with snail phosphate-buffered saline (sPBS; Yoshino, 1981) pH 7.2, suspended in sPBS by gentle spraying of buffer to detach them from the flask wall, removed from the flasks, and washed an additional 2 times with sPBS. The cells were then resuspended in SM supplemented with 5% heat-inactivated fetal bovine serum and cultured in 250-ml flasks for 48 h prior to use in sporocyst culture experiments. After the 48 h incubation period these media were considered conditioned (designated 'conditionedSM') and were removed from the flasks containing Bge cells, centrifuged for 10 min at 500 g at 4 °C to remove cellular debris, and immediately introduced into parasite culture experiments as described above.

### **Long Serial Analysis of Gene Expression (LongSAGE)**

SAGE libraries were constructed using 30 µg of total RNA isolated from miracidia and 6d- and 20d-old *in vitro*-cultured primary sporocysts following the I-SAGE Long Kit protocol (Invitrogen, Carlsbad, CA) following the manufacturer's instructions, with the exception of the use of pGEM3Z as the cloning vector. One Shot TOP10 Electrocomp *E. coli* cells (Invitrogen) were transformed with recombinant pGEM3Z clones containing SAGE concatemers by electroporation using an Eppendorf (Westbury, NY) *E. coli* electroporation apparatus. Plasmid sequencing templates were prepared from 1.2 ml cultures using alkaline lysis as performed by a RevPrep Orbit robotic workstation (GeneMachines, San Carlos, CA). Sequencing reactions were run on an ABI 3730xl capillary DNA sequencer. Recombinant pGEM3Z clones containing SAGE concatemers were sequenced using only the M13F primer to avoid duplicate sampling of SAGE tags due to overlapping bidirectional sequences from individual clones.

Sequences collected were analysed with software created specifically for *Schistosoma* SAGE analysis. The SAGE software extracts ditag sequences from the ABI 3730xl results according to the SAGE sequence grammar, passes out individual SAGE tags, and reduces all SAGE tags to a table of uniqueSAGE tag sequences and their observed frequencies among all of the *Schistosoma* SAGE libraries. In cases where a ditag sequence was sampled more than once, only 1 representative was used in generating tag frequencies (Emmersen, 2008a). Tags that contained base call ambiguities or bases with PHRED (Ewing *et al.* 1998; Ewing and Green, 1998) values of less than 10 (10% or greater chance of incorrect base call) were excluded from analyses (Emmersen, 2008b). Additional putative sequencing error was removed by identifying SAGE tag sequences that did not have a perfect sequence match in

the set of genome project gene predictions and that did not appear more than once in any of the SAGE libraries. As such, tags appearing at least twice in at least one SAGE library but that did not have a sequence match within the predicted genes were assumed to be from legitimate rare transcripts, from allelic variants, or unsequenced regions of the genome and were retained for analysis of differential gene expression.

SAGE tag sequences were mapped to genes predicted by the ongoing genome project (version 4.0e) using custom software created specifically for *Schistosoma* SAGE. As SAGE tags are generated from the 3'-most NlaIII restriction site of the transcript, we included observed (EST) or theoretical 3' UTR sequences when assigning tags to genes. Of all the genes predicted by the ongoing genome project (version 4.0e), 5334 had empirical EST data for 3' UTR lengths. In these cases, the 3' UTR sequence used for assignment of SAGE tags to genes was that predicted by the ESTs. The 99% confidence of the observed 3' UTR lengths was determined (1388 bp) and for genes without EST data this length was used to predict 3' UTR sequences from the genome.

In order to identify potential differentially-expressed gene between stages, tags were assigned an R-value, the log-likelihood ratio statistic of Stekel *et al.* (2000), which scores tags by their deviation from the null hypothesis of equal frequencies. Higher scores represent a greater deviation from the null hypothesis, while scores close to zero represent near constitutive expression. For this study, an R-value of 4 was used as a conservative measure to denote significant differences in gene expression between compared libraries (Stekel *et al.* 2000). In addition, Fisher's exact test was used to compare the effects of Bge cell-conditioning on the 6-d and 20-d sporocysts.

### Quantitative PCR

Total RNA was extracted using TRIzol as described above. Single-stranded cDNA was synthesized using the SuperScript III First-Strand Synthesis System for RT-PCR (Invitrogen, Carlsbad CA). Quantitative real-time PCR (qPCR) primers were designed using Primer Express software (Applied Biosystems, Foster City, CA). qPCR was performed in a GeneAmp 7300 PCR apparatus in a 96-well format using SYBR green chemistry. Theoretical tag numbers were approximated by setting the highest Ct value to 1 (Smp\_134670.2) and each increasing cycle number corresponds to a 2-fold increase in transcript abundance. This calculation assumes that amplification efficiency is similar for each primer set.

## RESULTS

### SAGE libraries

Five SAGE libraries were constructed from miracidia, 6-day unconditioned sporocysts, 6-day Bge cell-conditioned sporocysts, 20-day unconditioned sporocysts, and 20-day Bge cell-conditioned sporocysts, resulting in 68 450, 68 044, 60 171, 30 684 and 52 666 sequenced tags, respectively, after removal of sequencing error (Williams *et al.* 2007). The total number of unique SAGE tag sequences obtained from all 5 libraries was 21 440, including 8180 detecting sense transcription of a single gene, 4544 detecting anti-sense transcription

of a single gene, 625 unresolved among possible sense transcription of several genes, 32 unresolved among possible antisense transcription of several genes, and 8059 not assigned to a gene. Of the 8059 unassigned tags, 2836 matched the genome but were not associated with any predicted gene or known transcript, 1900 matched the genome in multiple locations, and 3233 did not match the genome at all. Of the 13 185 transcripts predicted by the ongoing genome project, 12 879 (96.7%) contained at least one NlaIII site and are detectable by SAGE. All SAGE data were deposited to GenBank's GEO database under Accession number GSE9722.

### Gene abundance

Of the 21 440 unique tag sequences observed in the 5 SAGE libraries, 30.4% had a frequency of 1, 37.1% a frequency of 2–5, 10.8% a frequency of 6–9, 19.8% a frequency of 10–99, and 1.9% a frequency of  $\geq 100$  at their highest observed abundance. Of the most abundant tags (frequency  $\geq 100$ ), 253 detected sense transcription of a single gene, 7 detected anti-sense transcription of a single gene, 39 were unresolved among multiple sense and/or anti-sense transcripts, and 113 were not assigned to a gene. For genes assigned both sense and anti-sense SAGE tags, a strong inverse correlation existed between the antisense : sense frequency ratio and the sense tag frequency (Fig. 1).

Using Blast2GO (Conesa *et al.* 2005) we identified level 3 molecular function gene ontology (GO) categories for 99 of the most abundant 150 tags (Fig. 2). The most highly represented functional categories included genes involved in transcription, translation, ion binding and oxidoreductase activity. There were 2147 SAGE tags uniquely expressed in miracidia and 9739 SAGE tags uniquely expressed in sporocysts. A comparison of the level 2 molecular function gene ontology categories between uniquely expressed miracidia and sporocyst transcripts revealed 2 categories, binding and catalytic activities, dominated gene expression in both miracidia and sporocysts, comprising >80% of the total transcriptome (Table 1). Although the overall pattern of unique miracidia and sporocyst transcript GO categories is similar, this may represent different transcripts performing similar molecular functions in the various stages. GO categories enriched in unique sporocyst transcripts include structural molecules, plus antioxidant and enzyme regulator activities, while chaperone activity was enriched in unique miracidial transcripts. In addition, we identified 911 SAGE tags uniquely expressed in miracidia and 3608 SAGE tags uniquely expressed in sporocysts corresponding to non-predicted and unknown transcripts or SAGE tag sequences not matching the genome.

### Differential expression of genes during development in miracidia, 6-day sporocysts and 20-day sporocysts

We identified 432 differentially expressed tags during larval development by comparison of all 5 SAGE libraries (R = 4). Differentially expressed sense SAGE tags are listed in Table 2, while all tags are shown in supplemental data file 1. Due to the limits of space, only tags with R = 7 are shown. The major classes of genes upregulated in miracidia compared to sporocysts were calcium-binding proteins (SME16, Calcineurin, 22.6 kDa tegument-associated antigen and synaptogagmin), heat shock proteins (HSP70, HSP90 and HSP27) and genes involved in cellular energy production (mitochondrial carrier protein, lactate

dehydrogenase and phosphoglycerate kinase). The major classes of proteins upregulated in sporocysts compared to miracidia were associated with transcription and translation (elongation factor 1-alpha, polyadenylate binding protein, and many ribosomal proteins). Some heterogeneity may exist in the transcript levels of individual miracidia or sporocysts; however, the use of large populations of parasites more accurately reflects the transcriptomic profiles of the developmental stages sampled.

### Differential expression of genes between conditioned and unconditioned sporocysts

Using Fisher's Exact test (Fisher, 1922) with a cutoff of  $P < 0.01$ , we identified 53 tags that were differentially expressed between 6-day unconditioned and 6-day conditioned sporocysts, 22 of which were sense tags (Table 3). Forty-five tags were differentially expressed ( $P < 0.01$ ) between 20-day unconditioned and 20-day conditioned sporocysts, 19 of which were sense tags (Table 3). Fifteen of these differentially expressed genes were in higher abundance in unconditioned media than in conditioned media. Transcripts differentially expressed due to the effects of conditioning include HSP90, thioredoxin reductase, elongation factor 1-alpha, multiple ribosomal proteins, and proteins of unknown function. Seven tags were found to be differentially expressed in both 6-day and 20-day datasets.

### Quantitative PCR

Since SAGE is quantitative in nature, we selected 3 genes exhibiting of high, medium and low expression levels in the miracidial stage and used real-time qPCR to independently examine their transcript levels within a miracidial cDNA pool. Those genes with the highest tag number (Smp\_071390, Smp\_009760 and Smp\_067800) correlated significantly in abundance within the larval cDNA pool by qPCR (Fig. 3). Similar correlations of tag number and transcript abundance by qPCR can be observed for genes with medium and low tag numbers (Fig. 3). These results indicate that tag numbers are predictive of relative levels of the transcript abundance.

## DISCUSSION

SAGE allows for the dissection of complex processes involving the interaction of multiple genes or gene families, such as stage-specific differentiation or response to external stimuli, on a transcriptome-wide level. The quantitative nature of SAGE also enables one to analyse thousands of transcripts from a given sample simultaneously, allowing for greater coverage of expressed mRNAs and the detection of low abundance transcripts that may be missed using methodologies like DNA microarrays (Velculescu *et al.* 1995). SAGE has been employed in numerous areas of biological, medical, and pharmaceutical research and has proved to be an excellent tool for comparing gene expression profiles between normal and abnormal cells occurring in a diseased state (e.g., tumors; St-Croix *et al.* 2000), differentially-treated, experimental cell populations (de Waard *et al.* 1999) or whole organisms sampled over a period of development (e.g., *Drosophila* : Lee *et al.* 2005 or parasites: Palm *et al.* 2005; Skuce *et al.* 2005; Kronstad, 2006).

Previously, only 2 studies have undertaken large-scale gene expression studies focusing on intramolluscan larval schistosomes. Vermeire *et al.* (2006) investigated miracidia and 4-day *in vitro* cultured sporocysts using a 7335 feature oligonucleotide microarray and Jolly *et al.* (2007) identified gene transcript changes between cercariae, daughter sporocysts recovered from infected snails, and adult worms utilizing an 11 998 feature array. We identified a large number of stage-associated transcripts that correlated with the Vermeire *et al.* study (2006), despite differences in culture conditions and sporocyst age. For example, both studies found SME16, p40 egg antigen, myosin light chain, phosphoenol-pyruvate carboxykinase (PEPCK), and secretory glycoprotein kappa-5 to be stage-associated with miracidia. Similarly, calreticulin, polo-like kinase (smPLK), 14-kDa fatty acid binding protein (Sm14; M60895) and a protein similar to insulin induced gene-1 were significantly associated with the sporocyst stage in both studies, demonstrating that similar results may be obtained using both SAGE and microarray methodologies in these stages. Differential gene expression for these transcripts was confirmed by real-time quantitative PCR (Vermeire *et al.* 2006) and further serves to validate SAGE as a quantitative transcriptomic method when applied to schistosomes. Williams *et al.* (2007) also demonstrated, using semi-quantitative reverse transcription PCR (RT-PCR) that overall expression levels and changes in expression levels correlate well between SAGE and RT-PCR in stage-specific comparisons.

Based on GO assignments, various functional groups were identified by SAGE as being over-represented in differentially expressed genes, uniquely expressed genes and/or genes having very high expression levels (SAGE tag frequencies). These functional groups are discussed below.

### Heat-shock proteins (HSP)/chaperonins

HSPs play important roles in folding, secretion, regulation, assembly, translocation and degradation of other proteins and as such are critical to numerous biochemical and molecular cell processes (Brown *et al.* 2007). The expression levels of HSP86/90, HSP40/DnaJ, HSP70/BiP, USP (universal stress protein-like) and p40 egg antigen (HSP27) were significantly higher in miracidia than 6- or 20-day sporocysts. HSP90 is an ATP-dependent chaperone involved in the activation and trafficking of proteins (Young *et al.* 2004). The p40 egg antigen contains 2 alpha-crystallin domains and exhibits high homology to small heat shock proteins from *Drosophila* (HSP27; Nene *et al.* 1986). The biological significance of *S. mansoni* miracidia secreting such high levels of a small heat shock protein homologue remains unclear. However, the secretion of HSPs by parasitic helminths is emerging as a common theme (Nene *et al.* 1986; Cai *et al.* 1996; Knudsen *et al.* 2005; Craig *et al.* 2006; Cass *et al.* 2007). Knudsen *et al.* (2005) identified HSP86/90, HSP70 and HSP60 in cercarial secretions, while p40 and other HSP/chaperone family members are secreted by schistosome eggs (Cai *et al.* 1996; Cass *et al.* 2007) or found in the empty eggshells of newly hatched *S. japonicum* miracidia (Liu *et al.* 2006). In a recently completed proteomic study, excretory-secretory proteins (ESP) released by *in vitro*-cultured *S. mansoni* miracidia, HSP/chaperonins were identified as a major constituent (Guillou *et al.* 2007; Wu *et al.* 2008). In the mammalian host, schistosome-secreted HSPs, like p40, serve as powerful immunogens eliciting production of pro-inflammatory cytokines resulting in extensive tissue fibrosis (Cai *et al.* 1996). Recently, it has been proposed that HSPs represent important



'danger signals' that, upon binding to macrophages/monocytes receptors (e.g., toll-like or scavenger receptors), stimulate release of pro-inflammatory cytokines or chemokines (Binder *et al.* 2004). Because induction of inflammatory fibrosis and granuloma formation around eggs is required for their efficient excretion from the host, the schistosome parasite appears to manipulate the host immune response to their advantage, by increased expression of HSPs and other immunogens (Binder *et al.* 2004).

The transition from free-living miracidium to parasitic sporocyst is accompanied by morphogenetic and physiological changes (Vogel and Seidel, 1972). Upregulation of HSPs may very well represent a stress response to these changes. During larval transformation the shedding, and subsequent degeneration, of ciliary epidermal plates during formation of the sporocyst tegument appears to represent a major source of excreted larval proteins and thus likely represents the source of the abundant HSPs found in larval ESP (Guillou *et al.* 2007; Wu *et al.* 2008). Yet to be answered, however, is the question of whether or not HSPs released by the parasite serve to alert or suppress the snail host's immune system.

### Calcium-interactive proteins

The divalent cation  $\text{Ca}^{++}$  is used as a cellular signal or ionic cofactor involved in diverse metabolic processes, including secretion, metabolism, muscle movement and neuronal function (Bhattacharya *et al.* 2006). Likewise, molecular interactions with calcium appear to play important roles in several physiological processes that govern miracidial infection of the snail host, especially its initial development to the parasitic sporocyst stage. Host entry, miracidium-sporocyst transformation, muscle movement and larval motility, and enzyme regulation all appear to be calcium-dependent processes (Sponholtz and Short, 1976; Knabe *et al.* 1982; Noel *et al.* 2001). For example, calcium chelators and pharmacological calmodulin antagonists have been shown to inhibit schistosome egg hatching and/or miracidial transformation (Katsumata *et al.* 1988, 1989; Kawamoto *et al.* 1989). Other studies have shown that calcium mobilization plays a role in cercarial penetration processes, possibly by Ca-regulation of protease activities during infection (Lewert *et al.* 1966; Fusco *et al.* 1991) or within penetration glands (Dresden and Edlin, 1975). The finding that excystment of *Paragonimus ohirai* metacercariae is a  $\text{Ca}^{++}$ -dependent process (Ikeda, 2004) indicates that the role of calcium as a regulator of larval development may be functionally conserved across trematode species. In the present study, 5 calcium-signalling or binding molecules were found to have stage-associated expression. Calcineurin B and SME16 were found to be highly expressed in miracidia, while calreticulin and calmodulin were mainly associated with 6-day and/or 20-day-old sporocysts. These results are consistent with the findings of recent gene expression analyses of calponin, SME16, calreticulin and calpain in miracidia of *S. japonicum* (Liu *et al.* 2006) and SME16 and calreticulin in *S. mansoni* miracidia (Vermeire *et al.* 2006; Guillou *et al.* 2007). Calreticulin, a versatile protein typically associated with chaperone activity, also functions as a storage form for calcium and a signalling molecule involved in regulating calcium homeostasis (Gelebart *et al.* 2005). Recently, it was shown that a calreticulin-like protein from endoparasitoid wasp venom fluid inhibits haemocyte spreading behaviour and thus prevents encapsulation within its lepidopteran host (Zhang *et al.* 2006). In this light, the finding of a calreticulin in ESP of *S.*

*mansoni* primary sporocysts (Guillou *et al.* 2007) suggests that it may have a role as a parasite defensive mechanism against haemocyte encapsulation.

Calmodulin is a multifunctional protein involved in the regulation of a variety of cellular processes. In mammalian cells, calmodulin functions in the activation of protein kinases, smooth muscle contraction and calcium channel regulation (Bers and Guo, 2005). Most pertinent to our studies was an earlier finding that the hatching of *S. mansoni* eggs appears to be a Ca<sup>++</sup>/calmodulin-dependent process (Katsumata *et al.* 1989). The fact that some HSPs (e.g., HSP90 and HSP70) contain calmodulin-binding domains (Song *et al.* 2007) implies potential molecular and functional interactions between these distinct molecular groups.

Since larval hatching from eggs is greatly facilitated by active larval motility, the finding of calponin, an actin- and tropomyosin-binding protein that acts as a regulator of smooth muscle contraction and motility (Winder *et al.* 1998), is consistent with its predicted involvement in hatching or other functions requiring miracidial motility. Synaptotagmin, Ca<sup>++</sup> sensors in a family of membrane-trafficking proteins involved in exocytosis and neurotransmitter release (Yoshihara and Montana, 2004), also is upregulated in miracidia compared to sporocysts. This protein may function as a signalling protein during the miracidial stage.

Two other Ca<sup>++</sup>-binding protein transcripts, SME16 and a small 8 kDa protein containing predicted EF-hand domains, also show differential expression in miracidia. Both SME16 and a protein with significant homology to the Ca-binding 8 kDa dynein light chain have been identified in ESP from *S. mansoni* miracidia (Wu *et al.* 2008). It now appears that SME16, previously shown to be a highly-expressed egg protein, originates within the fullyformed miracidium, although its function in larval development or metabolism is still unknown (Moser *et al.* 1992).

### Egg antigens

Transcripts of several abundant proteins found in soluble egg antigen (SEA) (Cass *et al.* 2007) were found to be highly expressed in miracidia (Vermeire *et al.* 2006). These transcripts included p40 egg antigen, PEPCK, secretory glycoprotein kappa-5, SME16 and several forms of glutathione-S-transferases (GSTs). This finding suggests that these antigens arise from the miracidia within the egg, or possibly other egg tissues (e.g., von Lichtenberg's membrane) (Neill *et al.* 1988), capable of synthesizing and secreting proteins from the egg. Neill *et al.* (1988) showed by electron microscopy that the von Lichtenburg's membrane, a structure that forms a thin cellular envelope surrounding the developing miracidium, contains ribosomes, and it was suggested that this tissue may be the source of SEA detected by immune sera and be responsible for promoting granuloma formation around the egg. Recent proteomic analysis of *S. mansoni* egg excretions and their immunolocalization supports this notion (Cass *et al.* 2007). However, the observation that most 'egg antigen' mRNAs are being transcribed in high abundance within miracidia suggests that these proteins are synthesized within the miracidium itself, are then released, and finally exported by some yet unknown mechanism through the tissues surrounding the larvae and out through the cribriform pores of the parasite's egg shell (Neill *et al.* 1988). El Ridi *et al.* (1996) demonstrated the *in situ* binding of mouse L-selectin to the surface of egg-

encased miracidia, implying external intact protein transport across the von Lichtenberg membrane, thus providing a basis for hypothesizing that reverse transport (miracidium-to-egg-shell surface) is highly possible. Further characterization of larval *S. mansoni* excretory-secretory proteins will help to further our understanding of the molecules released from the ciliated larvae during egg development, hatching and penetration of the molluscan intermediate host.

## Antioxidants

RNA transcripts for *S. mansoni* glutathione peroxidase (GPx) were found to be present at high levels in both miracidia and sporocysts, although a greater abundance was observed in miracidia mRNA populations. This observation is consistent with earlier microarray analyses in which GPx mRNA was found to be preferentially expressed in miracidia when compared to sporocysts (Vermeire *et al.* 2006) and in preliminary analyses of larval transcripts by SAGE (Williams *et al.* 2007). However, even though lower than in miracidia, GPx transcript abundance in sporocysts was still much higher (by 10-fold) than those of peroxiredoxins (Prxs), another prominent class of reactive oxygen-scavenging enzymes expressed mainly in sporocysts (Vermeire and Yoshino, 2007). *S. mansoni* GPx has been biochemically characterized as a primary lipid hydroperoxide reductant, but the molecule also possesses a hydrogen peroxide metabolizing activity (Mei *et al.* 1996). Another protein potentially involved in the *S. mansoni* redox pathway is translationally-controlled tumor protein (TCTP), which was the sixteenth highest expressed gene transcript from the 5 larval libraries. Studies of recombinant *Brugia malayi* TCTP showed that this protein possesses antioxidant activity, can be reduced by thioredoxin, and is upregulated upon host infection (Gnanasekar and Ramaswamy, 2007). In yeast, it is upregulated by treatment with H<sub>2</sub>O<sub>2</sub> (Bonnet *et al.* 2000). The presence of GPx and TCTP in such high abundance in miracidia and sporocysts suggests that it may provide protection for both the developing miracidia within eggs in the definitive host as well as some measure of protection against oxidative damage in the sporocyst stage. It seems plausible that GPx and TCTP may also aid sporocysts in their defence against reactive oxygen species (ROS) naturally occurring in the haemoglobin-rich plasma environment of the snail host (Hahn *et al.* 2001a) or produced by snail haemocytes during an encapsulation reaction (Hahn *et al.* 2001b; Bender *et al.* 2005). Recent studies have shown that Prx expression can be induced in *S. mansoni* mother sporocysts upon exposure to *B. glabrata* embryonic cells or exogenous hydrogen peroxide *in vitro* (Coppin *et al.* 2003; Vermeire and Yoshino, 2007). Thioredoxin, another redox pathway enzyme found to be moderately expressed in all 5 of our *S. mansoni* libraries, had highest expression levels in 6- and 20-day sporocysts cultured in snail cell-conditioned medium. GST-26 and GST-28, also possessing anti-oxidant activities, were more highly expressed in sporocysts than in miracidia.

## Effects of cultivation in *Biomphalaria glabrata* embryonic (Bge) cell-conditioned medium

Co-cultivation of *S. mansoni* primary sporocysts with the Bge cell line results in the production of embryos of secondary (=daughter) sporocysts by 15 days in culture and fully formed daughter sporocysts from 20 days and after (Yoshino and Laursen, 1995). However, due to the complexity of performing SAGE analysis on a multi-organism dataset, we utilized Bge cell-conditioned medium to investigate the influence of host molecules on sporocyst

gene expression and larval development. Earlier experiments (Coppin *et al.* 2003; Vermeire *et al.* 2004) have demonstrated that factors secreted from Bge cells influence gene expression during *in vitro* development of *S. mansoni* sporocysts. For example, in sporocysts, glutaminyl t-RNA synthetase (GlnRS) transcripts were shown to be 3-fold higher and T-complex protein 1 subunit gamma (SmTCP-1) transcripts 1.3-fold higher in response to Bge cell-conditioned medium (Coppin *et al.* 2003). Our data indicate GlnRS transcripts are 2-fold higher in 6-day sporocysts cultured in conditioned medium and 1.3-fold higher in 20-day larvae. Vermeire *et al.* (2004) demonstrated that G-alpha subunit 1 (G-alpha1) and SmPLK gene expression was higher in 4-day sporocysts cultured in Bge cell-conditioned Bge medium than in unconditioned medium. In contrast, our data showed that SmTCP-1 and SmPLK were lower in conditioned 6-day sporocysts than in 6-day unconditioned sporocysts. The discrepancies between our data and those of Coppin *et al.* (2003) and Vermeire *et al.* (2004) are likely due to the differences in culture media used and/or the length of time cultured.

Overall, 22 sense tags were found to be differentially expressed between 6-day conditioned and unconditioned sporocysts, of which 14 were upregulated in 6-day conditioned sporocysts compared to 8 transcripts upregulated in unconditioned sporocysts. This compares to 19 sense tags being differentially expressed between 20-day conditioned and unconditioned sporocysts, of which 16 were more highly expressed in unconditioned versus conditioned media. The majority of upregulated genes in both 6- and 20-day sporocysts involved transcriptional and translational processes, possibly reflecting a high degree of mitotic and protein synthetic activity associated with somatic and germinal tissue growth at this stage of larval development. Thirty-two % of the sense tags differentially expressed between 6-day conditioned and unconditioned sporocysts were higher in unconditioned media whereas 84% of the sense tags differentially expressed between 20-day conditioned and unconditioned were higher in unconditioned media. Differences in the gene expression in early versus late developing larvae under different culture conditions suggest that snail cell components may be playing different roles in regulating gene expression throughout sporocyst development.

### Anti-sense transcripts

Unlike traditional oligonucleotide microarrays, SAGE analysis can identify anti-sense transcripts. These transcripts have been previously identified in *S. mansoni* (Waisberg *et al.* 2007) and other parasitic organisms, including *Leishmania* (Dumas *et al.* 2006), *Onchocerca volvulus* (Erttmann *et al.* 1995), *Plasmodium falciparum* (Gunasekera *et al.* 2004) and *Trypanosoma brucei* (Liniger, 2001). Anti-sense SAGE tags are highly represented in these parasites, constituting 17% and 21.5% of all tags and in *Plasmodium* and *Toxoplasma gondii*, respectively (Patankar *et al.* 2001; Radke *et al.* 2005). Moreover, in *P. falciparum* (Gunasekera *et al.* 2004) and other organisms (Farrell and Lukens, 1995; Luther *et al.* 1998; Hastings *et al.* 2000) there exists a significant inverse relationship between anti-sense and sense tag frequencies that have been speculated as being a novel form of post-transcriptional gene regulation. In our study, 35% of the total mapped tags appear to be anti-sense tags and a strong inverse correlation between sense to anti-sense transcription exists. Gene loci containing higher levels of antisense tags contain lower levels of sense transcripts and vice-

versa, supporting the notion that antisense transcription may be a novel form of post-transcriptional regulation. Three mechanisms of anti-sense-mediated post-transcriptional regulation have been proposed, (a) anti-sense transcripts bind to the complementary sense transcripts, targeting it for RNAi mediated decay; (b) interfering with mRNA elongation or; (c) binding to the sense transcript and interfering with translation (Gunasekera *et al.* 2004).

## Conclusions

We have utilized LongSAGE to profile gene expression changes during early larval schistosome development, targeting the transition from free-living miracidium to the snail-parasitic mother sporocyst stage. This study represents the largest and most comprehensive transcriptomic analysis of gene expression changes during the earliest stages of intramolluscan larval *S. mansoni* development. We identified genes potentially involved in parasite growth and development, including many genes that are expressed in a stage-associated manner, thereby increasing our knowledge of putative regulatory networks in establishment of molluscan schistosome infections. Although the *in vitro* culture system employed in this study may not exactly mimic *in vivo* development, parasites in this *in vitro* system appear to develop normally (Basch and DiConza, 1974) and daughter sporocyst production is attainable (Yoshino and Laursen, 1995). Moreover, susceptible strains of *B. glabrata*, when injected with *in vitro*-cultured sporocysts, develop fully patent infections (Granath and Yoshino, 1984). Also, the miracidia recovered using our isolation procedure likely represent a heterogeneous population with regards to age or maturation synchrony, and may account for variation in individual gene expression between stages, especially those with low transcript numbers. However, because our analyses encompass large population sizes, data on differential gene expression captures a transcriptomic profile of the majority of parasites represented in those populations. In addition, we have identified 4519 SAGE tags uniquely expressed in either the miracidial or sporocyst stage corresponding to non-predicted transcripts, unknown transcripts or unsequenced regions of the genome. These transcripts may represent important developmental processes crucial to the survival of these individual stages. As the *S. mansoni* genome is further annotated, these results can be updated with additional SAGE tag mappings thereby identifying and further elucidating the function of these stage-specific transcripts.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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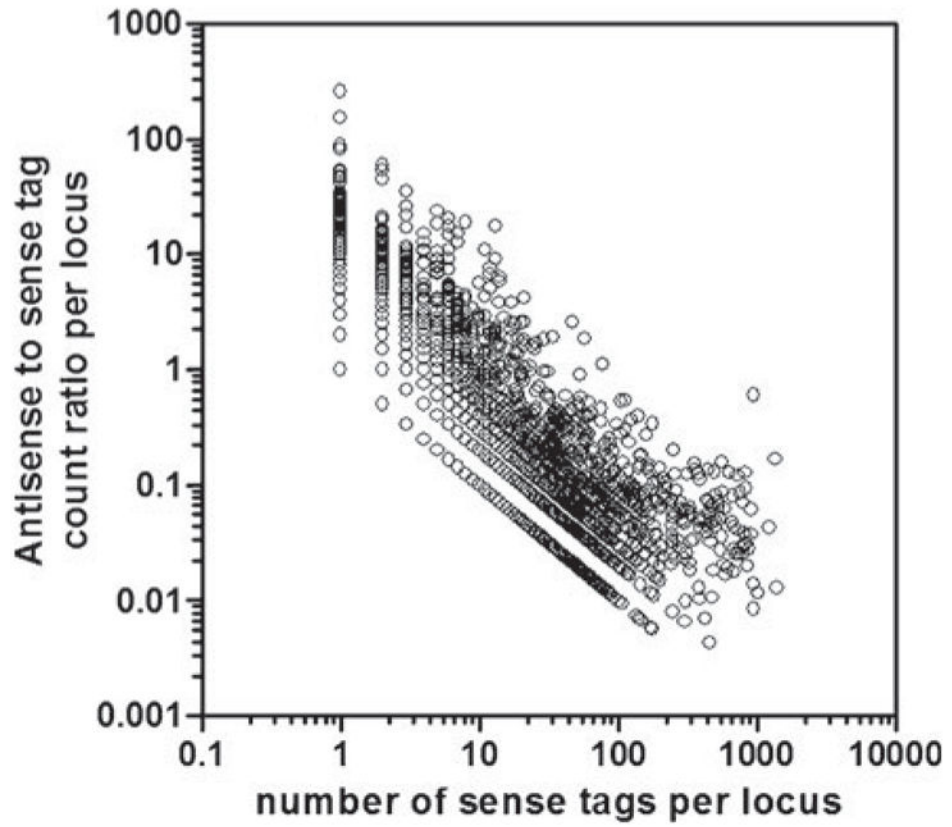


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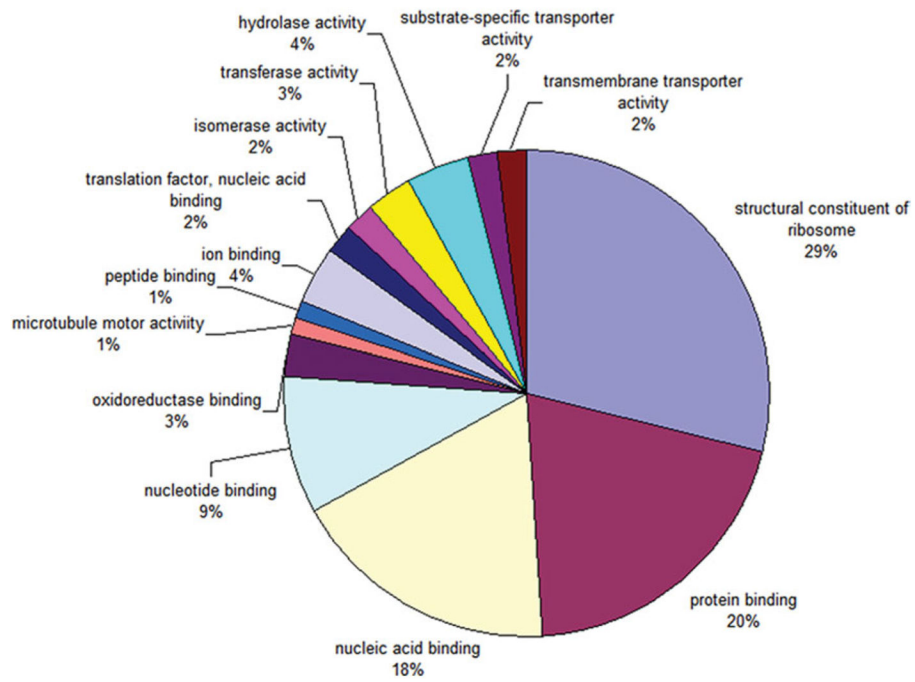
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## APPENDIX A. SUPPLEMENTARY DATA

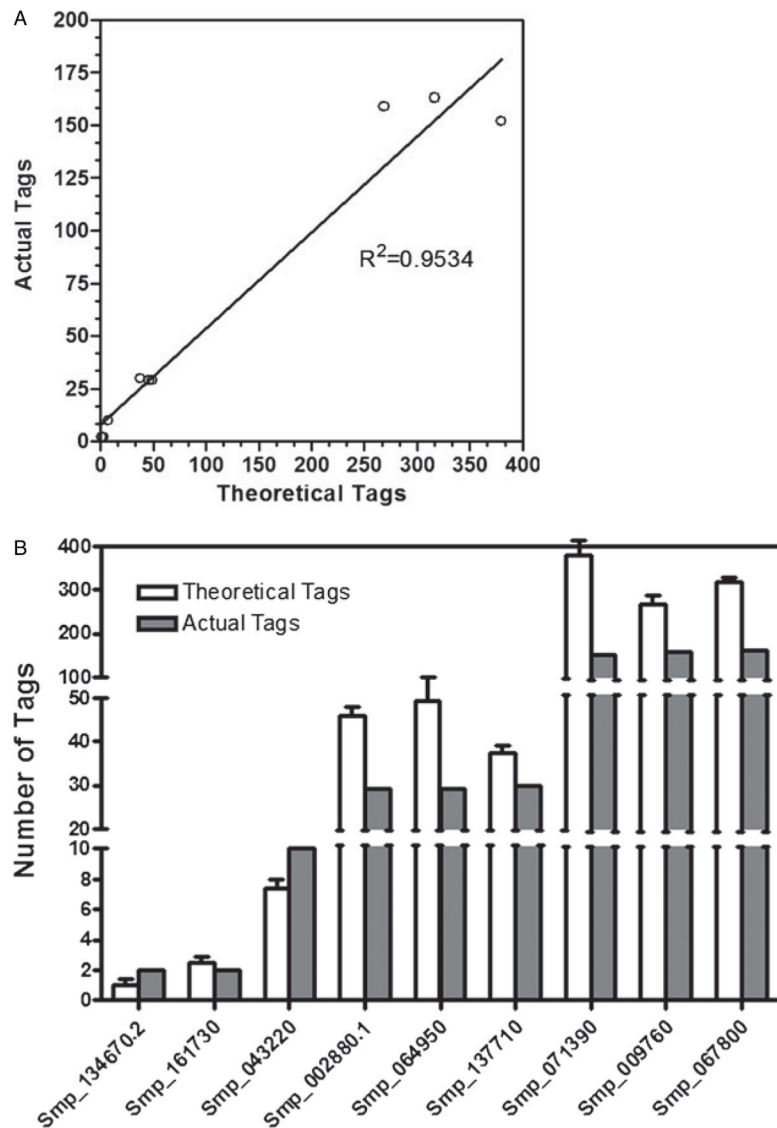
The following additional data files are available with the online version of the paper. Supplemental data file 1 contains all tags differentially-expressed ( $R = 4$ ) during larval development. Supplemental data file 2 contains all tags differentially expressed ( $P < 0.01$ ) between 6-day conditioned and unconditioned sporocysts and supplemental data file 3 contains all tags differentially expressed ( $P < 0.01$ ) between 20-day conditioned and unconditioned sporocysts.



**Fig. 1.** Correlation between anti-sense : sense frequency ratio and the sense tag frequency for genes assigned both sense and anti-sense SAGE tags.



**Fig. 2.** Level 3 molecular function GO categories for the most highly expressed transcripts in the 5 *Schistosoma mansoni* SAGE libraries. Percentages represent the frequency of each term.

**Fig. 3.**

(A) Quantitative PCR examination of high, medium and low expressed transcripts. (A) Theoretical transcript numbers are compared to actual transcript numbers for nine transcripts: Smp\_134670.2 (expressed protein of unknown function), Smp\_161730 (RNA binding protein), Smp\_043220 (expressed protein of unknown function), Smp\_002880.1 (ATP synthase alpha), Smp\_064950 (calcineurin B), Smp\_137710 (putative drug transporter), Smp\_071380 (adenylate kinase) and Smp\_067800 (fibrillin 2). (B) The correlation between actual transcript levels and theoretical transcript levels.

**Table 1**

Transcripts uniquely expressed in *Schistosoma mansoni* miracidia (2,147 SAGE tags corresponding to 448 transcripts) or sporocysts (9739 SAGE tags corresponding to 2104 transcripts) within various GO categories, also included are unique unknown SAGE tags

Gene ontology category	Transcripts uniquely present in miracidia (%)	Transcripts uniquely present in sporocysts (%)
binding	222 (49.6)	1030 (48.9)
catalytic activity	144 (32.1)	662 (31.5)
transporter activity	22 (4.9)	119 (5.7)
transcription regulator activity	18 (4)	93 (4.4)
molecular transducer activity	14 (3.1)	49 (2.3)
enzyme regulator activity	13 (2.9)	37 (1.8)
translation regulator activity	5 (1.1)	18 (0.9)
structural molecule activity	4 (0.9)	66 (3.1)
motor activity	4 (0.9)	21 (1)
auxiliary transport protein activity	1 (0.2)	3 (0.1)
chaperone activity	1 (0.2)	1 (0.05)
antioxidant activity	0 (0)	3 (0.1)
Unknown SAGE tags	911	4519

**Table 2**

Sense tags differentially expressed (R 7) between the 5 libraries (Values represent the individual tag percentage abundance in a given library.)

SAGE tag sequence	R-Value	Miracidia	6-day unconditioned sporocysts	6-day conditioned sporocysts	20-day unconditioned sporocysts	20-day conditioned sporocysts	Description
<i>(a) Highest expression in miracidia</i>							
CATGTTTGTTGTTAAATTCGGA	162-18	0-00764	0-00113	0-00138	0-00062	0-00053	Smp_049240 Heat-shock protein beta-1 (HspB1) (HSP 27), putative
CATGTTTTTCAAAATAAATAATT	123-62	0-00313	0-00003	0-00003	0	0	Smp_096390.2 calcium-binding protein (SME16), putative
CATGACAAAAGCACAAACATTGT	100-45	0-00257	0-00001	0-00003	0-00003	0	Smp_126100 hypothetical protein
CATGTATTTTATCTCCTTTT	73-03	0-00222	0-00006	0-00007	0-00007	0-00009	Smp_071390 adenylate kinase, putative
CATGAATTTAAACAATTGTGA	55-67	0-00133	0	0	0	0	Smp_154690 ribosomal protein related
CATGTGGACAGACAAAAA	51-01	0-00149	0-00003	0-00005	0-00003	0-00004	Smp_176700 expressed protein
CATGATTAATTTGAATGTTAA	36-19	0-00238	0-00029	0-00047	0-00114	0-00085	Smp_067800 fibrillin 2, putative
CATGATCACTAAATAAATGTC	33-09	0-00095	0-00006	0	0	0-00004	Smp_097490 P25 alpha-related
CATGGGTCAAAATCTGGTTAA	32-74	0-00108	0-00007	0-00012	0	0-00002	Smp_078590 expressed protein
CATGTACATCCTGTATTTCT	31-98	0-00224	0-00038	0-00005	0-00081	0-00047	Smp_003770 histone h1/h5, putative
CATGGAAGAAAGTGGATTAGCT	25-14	0-0047	0-00197	0-00273	0-00261	0-00177	Smp_072330.2 heat shock protein, putative
CATGCGACAAATCAATAAAAA	24-47	0-00058	0	0	0	0	Smp_122820 expressed protein
CATGTTCAATATTAGTATTCA	18-07	0-00133	0-00028	0-00048	0-00033	0-00021	Smp_145300 peptidylglycine alpha hydroxylating mono-oxygenase, putative
CATGGCTATTGATATGAAATA	15-32	0-00045	0	0-00005	0	0-00002	Smp_096790 expressed protein
CATGCTTGCGGTTGCTTCTC	14-62	0-00041	0-00001	0-00002	0	0	Smp_179260 alpha-galactosidase/alpha-n-acetylgalactosaminidase, putative
CATGTCAACTAGTTTTTCAA	14-01	0-00076	0-00013	0-00012	0-00007	0-00013	Smp_074930 zinc finger protein, putative
CATGAAGGAAAGATAAGAAAT	13-9	0-00047	0-00003	0-00002	0-00003	0-00002	Smp_005710.1 egg protein CP391S, putative
CATGTATACGTGCACATTGTT	12-68	0-00066	0-00012	0-00008	0-0001	0-00008	Smp_079010 camp-dependent protein kinase type II-alpha regulatory subunit, putative
CATGCAAGTTAAATTAATGAAA	12-63	0-00105	0-00019	0-0003	0-00033	0-00036	Smp_020920 DNAj homolog subfamily B member 4, putative
CATGGAATTAATAAGAAATGAG	12-22	0-00194	0-00091	0-00086	0-00078	0-00065	Smp_163710 expressed protein
CATGAATAACTAAGGGTTACT	11-84	0-00034	0	0-00002	0	0-00002	Smp_173150 cd63 antigen-like
CATGGACTTAGATTAATGGGA	11-52	0-00184	0-00066	0-00078	0-00113	0-00084	Smp_069130.2 heat shock protein 70 (hsp70)-4, putative
CATGTGTTATTGAGAGTAAAA	10-48	0-00127	0-00034	0-0008	0-00055	0-00046	Smp_051210.1 importin alpha 3.4, putative
CATGGAGGAATAGTTAAGGTA	10-38	0-00114	0-00037	0-00033	0-00036	0-00042	Smp_193420 hypothetical protein
CATGTCATATGTTTATACCTT	9-78	0-00044	0-0001	0-00007	0	0-00004	Smp_047900 erk1/2, putative
CATGTTTAAATCCAAATGGATG	9-75	0-00039	0-00003	0-00002	0-00007	0-00006	Smp_140000 tetraspanin-CD63 receptor, putative

SAGE tag sequence	R-Value	Miractdia	6-day unconditioned sporocysts	6-day conditioned sporocysts	20-day unconditioned sporocysts	20-day conditioned sporocysts	Description
CATGTTTCTTGTGTATCCA	9.55	0.00039	0.00004	0.00007	0	0.00004	Smp_076010 expressed protein
CATGTTGACACCAAACTGGT	9.23	0.00082	0.00024	0.00023	0.00029	0.00017	Smp_043120 universal stress protein, putative
CATGATTTTATCAAAATTAGTT	8.82	0.00042	0.00004	0.00005	0.00003	0.00009	Smp_138760 expressed protein
CATGCAAAACACCAAGTGATGT	8.52	0.00102	0.00047	0.00025	0.00088	0.00049	Smp_146510 expressed protein
CATGTACTTTGATCAACACGA	8.3	0.00051	0.00013	0.00007	0.00007	0.00011	Smp_188780 hypothetical protein
CATGTACTGATCTGGATAAA	8.27	0.00072	0.00121	0.00103	0.00205	0.00158	Smp_054160 Glutathione S-transferase 28 kDa (GST 28), putative
CATGGTTATGTAAITGCTTTT	8.24	0.00108	0.00031	0.00058	0.00062	0.00038	Smp_079270 crp-related
CATGATGTTTTTGGCTTTGTTT	8.23	0.00085	0.00019	0.00032	0.00026	0.00036	Smp_151250 er lumen protein retaining receptor, putative
CATGGTTGGCACAAGCTGAAG	8.08	0.00129	0.0006	0.0004	0.00059	0.00066	Smp_122840 Hypothetical protein, putative
CATGAAITTAACAAAAA	7.93	0.0002	0	0	0.00003	0	Smp_154690 ribosomal protein related
CATGCCACCACCTTGTTAACT	7.77	0.00105	0.00046	0.0003	0.00081	0.00049	Smp_006390 cystatin B, putative
CATGCTGATTTGTTTGCCAT	7.67	0.00039	0.00007	0.00003	0.00003	0.00009	Smp_136930 hypothetical protein
CATGGCTTACGATTAAGCG	7.66	0.00023	0	0.00003	0.00003	0	Smp_171960 short-chain dehydrogenase, putative
CATGGAAGTCATTATCTCAIT	7.64	0.00031	0.00001	0.00012	0	0.00004	Smp_068500 hypothetical protein
CATGTACATTTGTTTCTCC	7.64	0.00117	0.00041	0.00048	0.00055	0.00049	Smp_089000.1 translocon-associated protein, delta subunit, putative
CATGACGTTTATAITGATCAA	7.57	0.00037	0.00006	0.00003	0.00003	0.00006	Smp_063330 hypothetical protein
CATGCAITTCATTGTGTTTT	7.55	0.00066	0.00013	0.00018	0.00026	0.00021	Smp_061250 mitochondrial carrier protein, putative
CATGGGGTGGTCAAGAGCACT	7.45	0.00085	0.00028	0.00022	0.00059	0.00053	Smp_167270 expressed protein
CATGTTTTGGAGCACACGTT	7.11	0.00019	0	0	0	0.00002	Smp_168390 tnf receptor-associated factor, putative
CATGAAGTTGAATCTTTTGT	7.01	0.00034	0.00003	0.0001	0.00029	0.00004	Smp_030250 NF- $\kappa$ B subunit

(b) highest expression in 6-day sporocysts

CATGAACTAATCCCAATATAT	34.51	0.00012	0.00147	0.00163	0.00023	0.00046	Smp_153370 expressed protein
CATGCTGATATGTATGCGTT	22.26	0.00006	0.00068	0.0007	0.0001	0	Smp_017430 Sh23, putative
CATGATTGCATAAATAAACTA	12.87	0.00107	0.00226	0.00286	0.00212	0.00178	Smp_031310 40S ribosomal protein S26, putative
CATGTGAATAAGTAGAAATAA	12.77	0.00161	0.00339	0.00347	0.00303	0.00289	Smp_017280 ubiquitin, putative
CATGCTAAATACAACCTGCAAT	12.54	0.0015	0.00351	0.00269	0.00235	0.00241	Smp_105320 ribosomal protein smL37a, putative
CATGTATCATTTATGATATGTT	11.55	0.00001	0.00062	0.00045	0.00029	0.00028	Smp_169090 conserved hypothetical protein
CATGCTGCAGTAATCGCACAT	11.11	0.00146	0.00312	0.00263	0.00212	0.00178	Smp_142180 poly(ADP-ribose) glycohydrolase, putative
CATGATTTGATTTCTCTACAA	10.83	0.00131	0.00285	0.00269	0.00235	0.00186	Smp_047200.3 60S ribosomal protein L3, putative
CATGTAAGTGACTGGAAAT	7.92	0.00123	0.00256	0.00224	0.00186	0.00177	Smp_013470 ribosomal protein S2, putative



SAGE tag sequence	R-Value	Miracidia	6-day unconditioned sporocysts	6-day conditioned sporocysts	20-day unconditioned sporocysts	20-day conditioned sporocysts	Description
CATGTATAAACACAGATGGAAC	7.57	0.00047	0.0011	0.00101	0.00059	0.0004	Smp_002180 transcription factor btf3, putative
CATGGACAAGTACCATCCAGG	7.48	0.00095	0.00198	0.00199	0.00179	0.00141	Smp_132300 kif-3, putative
CATGTAATGAGAAAAAGTGATT	7.35	0	0.00028	0.00033	0.00016	0.00019	Smp_124750 expressed protein
<i>(c) highest expression in 20-day sporocysts</i>							
CATGAAAAATGTTAGATGTTGC	21.93	0.00025	0.00125	0.00138	0.00189	0.00177	Smp_113760 anti-inflammatory protein 16, putative
CATGATGCATCAAAATTAGAGA	19.45	0.00298	0.00228	0.00269	0.00355	0.00534	Smp_032560.2 expressed protein
CATGGACCCAAAAGTGTGGGA	18.44	0.00117	0.00241	0.00268	0.00326	0.00346	Smp_102070 Glutathione S-transferase 26 kDa (GST 26), putative
CATGTTACCACCAAACTTTTG	10.61	0.00202	0.00306	0.00264	0.00433	0.00357	Smp_146190 40S ribosomal protein S21e, putative
CATGCCATCCGTCAGCATAGA	10.41	0.00299	0.00219	0.00176	0.00391	0.00304	Smp_022640 60S ribosomal protein L13 (BBC1 protein homolog), putative
CATGCCATCAGCCTGTGCTGT	9.78	0.00361	0.00391	0.00407	0.00603	0.00547	Smp_029820 60S acidic ribosomal protein P2, putative
CATGTATCGTTCTATATTAAT	9.53	0.00015	0.00069	0.00068	0.00088	0.00085	Smp_095360 fatty acid binding protein, putative
CATGCTTATATGCCTATAGTG	9.09	0.00101	0.00188	0.00151	0.00257	0.00211	Smp_095540 cytochrome C oxidase, subunit II, putative
CATGAAGCCCTGTCGCAAAATC	8.32	0.00111	0.00169	0.00188	0.00257	0.00234	Smp_084460 60S ribosomal protein L28, putative
CATGTTATTTTAGGTAGGTTT	7.55	0.00086	0.00054	0.00066	0.00121	0.00148	Smp_181810 cytochrome c oxidase subunit 3, putative
CATGAGTAAATATGATACGCTA	7.35	0.00101	0.00118	0.00133	0.00212	0.00203	Smp_032950 Calmodulin (CaM), putative
CATGCACAGACAGCTGTAGTT	7.29	0.00262	0.00253	0.00271	0.00443	0.00353	Smp_041650 40S ribosomal protein S27, putative
<i>(d) lowest expression in miracidia</i>							
CATGGGATTCGGTTTATTTGA	12.15	0.00349	0.00526	0.00548	0.00678	0.00497	Smp_009690.1 60S acidic ribosomal protein P0, putative
CATGAACCAATCCAGGTGAGAT	9.97	0.00226	0.00403	0.00283	0.00394	0.00264	Smp_099870.1 elongation factor 1-alpha (eF-1-alpha), putative
CATGCTTATCTGTGTCTTC	8.89	0.00164	0.00306	0.00278	0.00244	0.00319	Smp_179420 expressed protein
CATGTTTTTGTGTGAAAAAAA	8.66	0.0001	0.00028	0.0006	0.00062	0.00013	Smp_102020 expressed protein
CATGATCCCGAAATGTATGTC	8.02	0.00015	0.00076	0.00058	0.00068	0.0007	Smp_030370 calreticulin autoantigen homolog precursor, putative
CATGAACAACCTGATTCAAAT	7.96	0.00003	0.00035	0.00032	0.00016	0.00051	Smp_064300 expressed protein
CATGTTCCGGAAGATTCGTTCT	7.56	0.0012	0.00253	0.00204	0.00218	0.00199	Smp_038510 60S ribosomal protein L6, putative

Table 3

Differential expression of transcripts due to the effects of conditioning with Bge excretory-secretory products, examined using Fisher's Exact test ( $P < 0.01$ )

SAGE Tag sequence	6-day unconditioned sporocysts	6-day conditioned sporocysts	20-day unconditioned sporocysts	20-day conditioned sporocysts	Description
<i>(a) higher abundance in 6-day unconditioned compared to 6-day conditioned sporocysts</i>					
CATGAACCATCCAGGTGAGAT	0.403	0.283	0.394	0.264	Smp_099870.1 elongation factor 1-alpha (ef-1-alpha), putative
CATGTTGAACGTGAAAACTACT	0.035	0.012	0.023	0.034	Smp_154530 mitochondrial ATP synthase B chain, putative
CATGTCAATCGTTCCCTTGCAC	0.201	0.116	0.15	0.156	Smp_174950 40S ribosomal protein S19, putative
CATGCTTAATAACAGCAGCAA	0.051	0.02	0.036	0.03	Smp_064830 survival motor neuron protein, putative
CATGAACCATCCCGGTGAGAT	0.209	0.136	0.156	0.116	Smp_189530 elongation factor 1-alpha (ef-1-alpha), putative
CATGTCAATTTGCGAAAAATG	0.015	0	0.007	0.002	Smp_154230.2 expressed protein
CATGAGTTTTATCGAAATGAT	0.012	0	0	0.002	Smp_084540.1 expressed protein
<i>(b) higher abundance in 6-day conditioned compared to 6-day unconditioned sporocysts</i>					
CATGGAAGAAAGTGGATTAGCT	0.197	0.273	0.261	0.177	Smp_072330.2 heat shock protein, putative (HSP90)
CATGTATAAAAAGCATCCTTTT	0.037	0.071	0.059	0.072	Smp_007260 ATPase, putative
CATGGTGATTATTTTTTGT	0.076	0.13	0.055	0.082	Smp_048660 nucleolar protein nop56, putative
CATGTGTTATTTGAGTAAAA	0.034	0.08	0.055	0.046	Smp_051210.1 importin alpha 3.4, putative
CATGTAATTTATTTTATATTA	0.01	0.038	0.026	0.017	Smp_073590 GTP-binding protein-like protein
CATGGAAAAATATATGATTTG	0.097	0.15	0.108	0.101	Smp_164650 lupus la ribonucleoprotein, putative
CATGAAATATACATATATATAT	0.001	0.018	0.013	0.011	Smp_002080 s-adenosylmethionine synthetase, putative
CATGTAATAAAGAGTAAATTT	0.003	0.03	0.013	0.011	Smp_052710 expressed protein
CATGTGAGAAAAGATACACTTT	0.022	0.053	0.026	0.053	Smp_027080 golgi membrane protein sb140 (yip1b), putative
CATGAATAAAAATATTTGCTTT	0.044	0.083	0.072	0.089	Smp_086480 Antigen Sm21.7, putative
CATGTTATGTAITGTTATTTT	0	0.013	0.01	0.006	Smp_153500 tyrosine protein kinase, putative
CATGCTCACAAATTTTTTTTGT	0	0.013	0.003	0.006	Smp_016600 solute carrier family 1 (glial high affinity glutamate transporter, putative
CATGAACCCCTTAITGTTATTT	0.009	0.03	0.007	0.011	Smp_022090 ribose-phosphate pyrophosphokinase 1, putative
CATGGCTGGGGAATATAAATA	0.019	0.071	0.049	0.023	Smp_135690 hypothetical protein
CATGTTTTTGTGTGAAAAAAA	0.028	0.06	0.062	0.013	Smp_102020 expressed protein
<i>(c) higher abundance in 20-day unconditioned compared to 20-day conditioned sporocysts</i>					
CATGGGATTCGGTTTATTTGA	0.526	0.548	0.678	0.497	Smp_009690.1 60S acidic ribosomal protein P0, putative
CATGAACCATCCAGGTGAGAT	0.403	0.283	0.394	0.264	Smp_099870.1 elongation factor 1-alpha (ef-1-alpha), putative

SAGE Tag sequence	6-day unconditioned sporocysts	6-day conditioned sporocysts	20-day unconditioned sporocysts	20-day conditioned sporocysts	Description
<i>(a) higher abundance in 6-day unconditioned compared to 6-day conditioned sporocysts</i>					
CATGTAATAAACACTTCAGCG	0.09	0.081	0.104	0.049	Smp_044280 fibrillarlin, putative
CATGAAAGTTGAAATCTTTTGGT	0.003	0.01	0.029	0.004	Smp_030250 NF-YA subunit
CATGTGGTTGAAATAAAATTTGC	0.22	0.203	0.257	0.163	Smp_096750 expressed protein
CATGGCTCTCCTTTACTCGAT	0.053	0.057	0.059	0.015	Smp_017280 ubiquitin, putative
CATGAAAGTTACTTTCACAAAAT	0.032	0.02	0.075	0.028	Smp_048430.1 thioredoxin reductase, putative
CATGACATCCAAAGCGGTGTTT	0.003	0.01	0.023	0.002	Smp_105210 zinc finger protein, putative
CATGAGCGAGAGGAAACGCGAA	0.059	0.042	0.062	0.021	Smp_103320 nuclear movement protein nude, putative
CATGGGGAAAAAGTATAAAATAA	0.007	0.007	0.016	0	Smp_130540 alanine aminotransferase, putative
CATGAAAGAGAGCATTAAATATT	0.004	0.007	0.016	0	Smp_074370 expressed protein
CATGTAACATAAATATACTTTG	0.003	0.01	0.026	0.004	Smp_040450 expressed protein
CATGCAGCTTGTGTGAATATG	0.016	0.012	0.026	0.002	Smp_132640 phosphatidylinositol synthase, putative
CATGAAACAAGTATGATCCCAT	0.012	0.012	0.023	0.002	Smp_024390 microsomal signal peptidase 25 kD subunit, putative
CATGTTTTTGTGTGAAAAAAA	0.028	0.06	0.062	0.013	Smp_102020 expressed protein
CATGTTGATACTAGAGTTTCT	0.003	0	0.02	0	Smp_140760 expressed protein
<i>(b) higher abundance in 20-day conditioned compared to 20-day unconditioned sporocysts</i>					
CATGCCAAAGTTTACAGTTAAA	0.069	0.078	0.075	0.156	Smp_105970 expressed protein
CATGACTATTCGGGTGGGTAC	0.001	0.002	0	0.025	Smp_131110 expressed protein
CATGATGCATCAAAATTAGAGA	0.228	0.269	0.355	0.534	Smp_032560.2 expressed protein