

# Sm14 gene expression in different stages of the *Schistosoma mansoni* life cycle and immunolocalization of the Sm14 protein within the adult worm

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## Abstract

Sm14 is a 14-kDa vaccine candidate antigen from *Schistosoma mansoni* that seems to be involved in cytoplasmic trafficking of fatty acids. Although schistosomes have a high requirement for lipids, they are not able to synthesize fatty acids and sterols *de novo*. Thus, they must acquire host lipids. In order to determine whether Sm14 is present in different stages of the life cycle of the parasite, we performed RT-PCR. Sm14 mRNA was identified in all stages of the life cycle studied, mainly schistosomulum, adult worm and egg. Additionally, we used a rabbit anti-Sm14 polyclonal antibody in an indirect immunofluorescence assay to localize Sm14 in adult worm sections. The basal lamella of the tegument and the gut epithelium were strongly labeled. These tissues have a high flow of and demand for lipids, a finding that supports the putative role of Sm14 as an intracellular transporter of fatty acids from host cells.

## Key words

- Sm14
- Fatty acid-binding protein
- FABP
- Immunolocalization
- RT-PCR

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Schistosomiasis is a chronic parasitic disease which affects more than 200 million people worldwide causing extensive liver damage (1). Although drugs are available to treat schistosomiasis, the large extension of endemic areas and the constant reinfections make chemotherapy an ineffective approach to control the disease (2). Therefore, much effort has been devoted to research for vaccine development. Recently, the World Health Organization selected six molecule candidates to compose a subunit vaccine against schistosomiasis (3). One of them is Sm14, a 14-kDa *Schistosoma mansoni* antigen that induces partial protection in mice following

vaccination and cercarial challenge (4). Sm14 is a cytoplasmic fatty acid-binding protein (FABP) and its ability to bind to palmitic and linolenic acids *in vitro* has been demonstrated (5).

Schistosomes have a strong requirement for lipids in order to synthesize and maintain their complex membrane systems (6). Fatty acids act as precursors for lipid and phospholipid synthesis, and therefore play an important role in the life cycle of the parasite, including membrane formation, functioning as lipid anchors for proteins, sexual maturation, and regulation of egg production (6). However, schistosomes are unable to syn-

thesize fatty acids and sterols *de novo*, and thus must acquire lipids from host cells (7). The parasite is capable of synthesizing only phospholipids and triacylglycerols from precursors obtained from the host such as the low-density lipoproteins (LDL) of the blood in which the parasites reside (8). LDL receptors have been identified on the surface of schistosome membranes (9). The fraction of fatty acids can subsequently be modified, mainly by chain elongation (7).

FABPs appear to be involved in the acquisition and utilization of fatty acids in different organisms (10). There are at least two main classes of FABPs, a cytoplasmic and a plasmalemmal FABP (11). Cytoplasmic FABP has been identified in a variety of parasites, including *Fasciola hepatica* (12), *Schistosoma mansoni* (5) and *S. japonicum* (13). Plasmalemmal FABP seems not to be present in trematodes.

The aim of the present study was to first determine the gene expression profile of Sm14 in different stages of the life cycle of the parasite and then to localize the Sm14 protein within adult *S. mansoni* by indirect immunolabeling using rabbit anti-Sm14 polyclonal serum followed by fluorescence microscopy.

For detection of Sm14 in different stages of the parasite life cycle, we used RT-PCR. Eggs were recovered from macerated liver of infected mice and adult worms were perfused from the portal vein of 42-day-infected mice. The cercariae were obtained from *Biomphalaria glabrata* infected with a single miracidium *in vitro* and recovered using artificial light. The schistosomula were mechanically transformed according to Ramalho-Pinto et al. (14) and cultured *in vitro* in RPMI 1640 medium (Gibco BRL, Rockville, MD, USA) until day 7 that corresponds to the lung stage worm.

Total RNA was extracted from newly transformed schistosomula, *in vitro* cultured schistosomula, adult worms, cercariae and eggs using the Rneasy total RNA system (Qiagen Inc., Valencia, CA, USA). The RNA

samples were quantified by spectrophotometry and 2.5 µg was used to synthesize cDNAs using oligo d(T) as a primer. cDNAs were then amplified using specific primers for Sm14 (Sm14f: 5'-CGGTGGTCGTTCAA GCGTATCAGAAGC-3' and Sm14r: 5'-CCCTCGCAAATATCAGTGTTCATTTG-3'). To standardize the quantities of cDNA used, we also amplified the cytochrome oxidase C1 gene (housekeeping) from *S. mansoni* using specific primers (CoxF: 5'-AAACTGCGAGGTTGAGCAGG-3' and CoxR: 5'-CAACACTAAACATAAAAA CAGATAG-3'). Five pmoles of each primer was used in PCR, which consisted of a denaturation step at 90°C for 1 min, a primer annealing step at 50°C for 30 s, and an extension step at 72°C for 1 min. This cycle was repeated 30 times for Sm14 and 22 times for Cox amplification. The samples were resolved on 4% acrylamide gels and stained with ethidium bromide and densitometry analysis of bands was carried out using the ImageMaster VDS® videodocumentation system and software (1996, version 2.0; Pharmacia Biotech Inc., San Francisco, CA, USA).

For immunofluorescence analysis, the specific antiserum was produced by immunization of two rabbits with recombinant Sm14 (4). The rabbits were injected three times at two-week intervals with 50 µg of recombinant Sm14 emulsified in Freund's complete adjuvant subcutaneously and the last immunization was performed without adjuvant. Sera were collected 10 days after the final boost. The specificity of the polyclonal antibody was confirmed by Western blot analysis (data not shown).

*S. mansoni* Puerto Rican strain maintained in BALB/c mice was used in the immunolocalization assay. Adult worms were obtained by perfusion of mice 42 days after infection (15). The recovered parasites were maintained in RPMI 1640 culture medium (Gibco BRL). About eight worms were included in resin (Tissue-Tek® 4583 O.C.T.

compound; Sakura Finetek, Torrance, CA, USA) and quickly frozen by immersion in liquid nitrogen. These blocks were cut with a microtome at  $-20^{\circ}\text{C}$  into sections of approximately 6-7  $\mu\text{m}$ . The cryosections were fixed with acetone on glass slides for 3-4 min at room temperature. Nonspecific binding sites were blocked by incubation of slides with naive goat serum diluted 1:10 in 0.15 M PBS, pH 7.2, for 30 min at room temperature. A rabbit anti-Sm14 antibody diluted 1:100 in PBS plus 10% (v/v) naive goat serum was added to each slide and kept at room temperature for 1 h. As a negative control, we used normal rabbit serum as primary antibody (data not shown). Unbound antibodies were removed with three washes of 10 min each with PBS. Slides were then incubated with goat anti-rabbit IgG conjugated with FITC (fluorescein isothiocyanate, Vector Laboratories, Burlingame, CA, USA) diluted 1:50 in 10% (v/v) naive goat serum. Slides were then washed in PBS again, dried and covered with Vectashield<sup>®</sup> fluorescent medium (Vector Laboratories) before a coverslip was applied. Slides were then kept at  $4^{\circ}\text{C}$  in the dark and immunofluorescence was observed using a fluorescence microscope (Nikon, Melville, NY, USA).

Using RT-PCR, we were able to identify transcripts for Sm14 in all parasite stages studied (Figure 1A). Using the values of Cox amplification to normalize the Sm14 mRNA levels we plotted a graph with the mean value of two experiments performed independently to determine the relative quantities of Sm14 mRNA (Figure 1B). Higher levels of Sm14 transcripts were detected in 6- and 7-day-cultured schistosomula, adult worms or eggs compared to the other parasite life cycle stages studied. Since the lung schistosomulum of *S. mansoni* is the main target of protective immunity in vaccinated mice and there is evidence of cross-reactivity among schistosomulum, adult worm, or egg antigens, we speculate that the higher expression of Sm14 in these parasite stages may be related to the ability of this

antigen to induce protection against infection in mice. However, further studies should be performed using a more powerful technique such as “real-time” PCR to confirm the Sm14 expression levels found within these parasite stages.

Figure 2 shows a longitudinal section of a

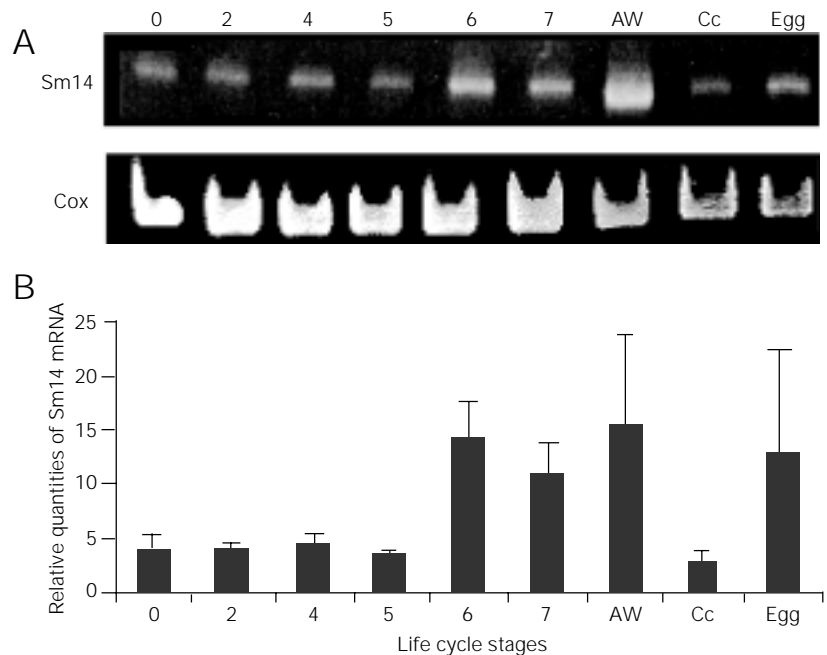


Figure 1. A, Amplification of cDNA from a *Schistosoma mansoni* newly transformed schistosomulum (0), and from 2, 4, 5, 6 or 7-day-cultured schistosomula; adult worm (AW); cercariae (Cc) or egg using specific primers for Sm14 or Cox. B, The data represent relative quantities of specific Sm14 mRNA in relation to the Cox transcripts (intensities of Sm14 in pixels  $\times 100$ /intensities of Cox in pixels).

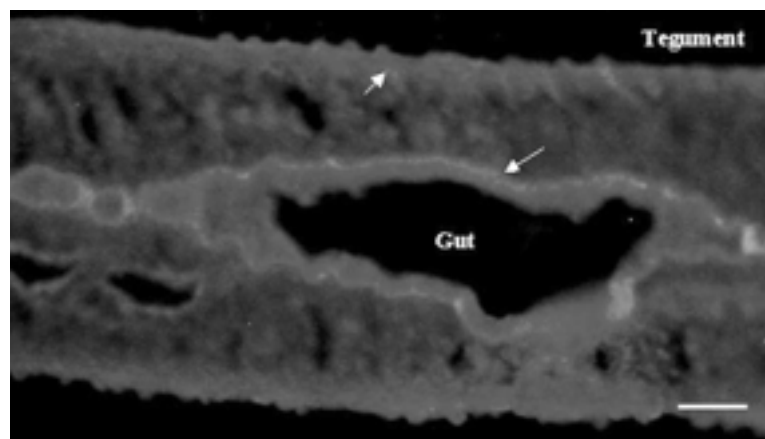


Figure 2. Localization of Sm14 by indirect immunofluorescence staining in adult worm cryosections. The sections reacted with rabbit anti-Sm14 polyclonal antibody diluted 1:100. The arrows indicate the basal lamella of the gut and of the tegument. Bar = 100  $\mu\text{m}$ .

male adult worm where the gut is clearly visible in the center. The major structure sharply delineated is the basal lamella of the gut which was revealed as a thin bright line in the immunofluorescence reaction. The basal lamella underlying the tegument was also strongly positive. Therefore, Sm14 was localized by anti-Sm14-specific antibodies in the basal lamella of the gut and underlying the tegument. Both tissues are related to flow of and demand for fatty acids and these results corroborate the putative role for Sm14. As a positive control, we used a rabbit anti-SWAP IgG as the primary antibody and the labeling observed was not localized to specific structures, but was broadly distributed throughout all tissues (data not shown). A previous study by Moser et al. (5) demonstrated that Sm14 was present on tubercles, which are structures located on the dorsal surface of the parasite rich in lipid globules, but not in the muscle layers or the tegument. Thus, our findings bring new insights into the localization and putative function of Sm14 in *S. mansoni*.

The localization of Sm14 shown in this study is related to other FABPs, which are preferably localized in tissues with a high flow of or demand for fatty acids (10). In mammals, there are three major cytoplasmic

FABPs: intestinal, heart and liver FABPs. Intestinal and liver FABPs are abundant in the gut epithelium, associated with delivery and metabolism of fatty acids from the diet (16). Heart FABP is expressed at high levels in the myocardium, associated with  $\beta$ -oxidation to generate power to the muscle (17). Although Sm14 has a higher phylogenetic relationship to heart FABP, it is known that *S. mansoni* is unable to use lipids as a power source (18,19).

Furthermore, Sm14 is localized in tissues near the interfaces of parasite/host contact, such as the basal lamella of the tegument and the epithelium of the gut. This suggests a putative role for Sm14 as a second element in the uptake and transport of fatty acids, but it is not clear whether this uptake occurs through the tegument or the gut because both tissues are related to the flow of and demand for fatty acids. Hockley and McLaren (20) demonstrated that the captured fatty acids go to the basal lamella of the tegument and are stored in the gut or in the esophageal gland. Furthermore, our results are similar to Gobert's findings (21), who recently localized *S. japonicum* FABP in the subtegumental region of male adult worms and vitellinic glands of female worms using electron microscopy and colloidal gold staining.

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