

# Identification of the *Boudicca* and *Sinbad* retrotransposons in the genome of the human blood fluke *Schistosoma haematobium*

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*Schistosomes have a comparatively large genome, estimated for Schistosoma mansoni to be about 270 megabase pairs (haploid genome). Recent findings have shown that mobile genetic elements constitute significant proportions of the genomes of S. mansoni and S. japonicum. Much less information is available on the genome of the third major human schistosome, S. haematobium. In order to investigate the possible evolutionary origins of the S. mansoni long terminal repeat retrotransposons Boudicca and Sinbad, several genomes were searched by Southern blot for the presence of these retrotransposons. These included three species of schistosomes, S. mansoni, S. japonicum, and S. haematobium, and three related platyhelminth genomes, the liver flukes Fasciola hepatica and Fascioloides magna and the planarian, Dugesia dorotocephala. In addition, Homo sapiens and three snail host genomes, Biomphalaria glabrata, Oncomelania hupensis, and Bulinus truncatus, were examined for possible indications of a horizontal origin for these retrotransposons. Southern hybridization analysis indicated that both Boudicca and Sinbad were present in the genome of S. haematobium. Furthermore, low stringency Southern hybridization analyses suggested that a Boudicca-like retrotransposon was present in the genome of B. truncatus, the snail host of S. haematobium.*

Key words: schistosome - long terminal repeat retrotransposon - mobile genetic element - reverse transcriptase - *Bulinus*

The life cycle of schistosomes, dieocious trematodes of the order Digenea, involves parasitism of both humans and genera of aquatic or amphibious snails specific to each species: *Biomphalaria* for *Schistosoma mansoni*, *Oncomelania* for *S. japonicum*, and *Bulinus* for *S. haematobium*. In the case of *S. mansoni* and *S. japonicum*, adult worms inhabit the portal system blood vessels and mesenteric veins of the intestines of their human hosts. Much progress has been made toward sequencing and sequence annotation of the genomes of *S. mansoni* and *S. japonicum* (El-Sayed et al. 2004), including characterization of much of their transcriptomes (Hu et al. 2003, Verjovski-Almeida et al. 2003), and the construction of a bacterial artificial chromosome (BAC) library representing an eight-fold coverage of the *S. mansoni* genome (Le Paslier et al. 2000). In contrast to the burgeoning wealth of sequence information for *S. mansoni* and *S. japonicum*, relatively little is known about the genome sequence of *S. haematobium*; less than seventy *S. haematobium* nucleotide sequences were present in GenBank as of February 2006, compared with over 100,000 each for *S. mansoni* and *S. japonicum*. The genomic DNA of *Schistosoma* is indistinguishable among *S. mansoni*, *S. japonicum*, and

*S. haematobium* by physical measurement (Hilyer 1974, Rollinson et al. 1997) and their chromosomal karyotypes are similar in appearance (Hirai et al. 2000).

Mobile genetic elements (MGEs) represent a major force driving the evolution of eukaryotic genomes (Charlesworth et al. 1994, Kidwell & Lisch 1997, Kazazian 2004) and play an important role in the establishment of genome size (Petrov et al. 2000). In schistosomes, more than half of the genome appears to be composed of, or derived from, repetitive sequences, to a large extent from retrotransposable elements (Brindley 2005, Copeland et al. 2005b). Schistosome MGEs include SINE-like retrotransposons (Spotila et al. 1989, Drew & Brindley 1995), long terminal repeat (LTR) retrotransposons (Laha et al. 2001, 2004, Foulk et al. 2002, Copeland et al. 2003, 2005a, DeMarco et al. 2004), non-LTR retrotransposons (Laha et al. 2002a, b, Brindley et al. 2003), and DNA transposons related to bacterial IS1016 insertion sequences (Feschotte 2004). The long terminal repeat retrotransposable elements, i.e. the LTR retrotransposons, are similar in structure and life cycle to retroviruses (Garfinkel et al. 1991, Havecker et al. 2004), differentiated from true retroviruses mainly by their lack of an envelope-encoding gene. They are of interest for their potential for horizontal transmission, as well as their ability to shed light on phylogenies of their host organisms when solely vertically transmitted.

*Boudicca*, the first fully characterized LTR retrotransposon from the genome of *S. mansoni* (Copeland et al. 2003, 2004), belongs to the gypsy-like retrotransposons, one of three distinct major groups of LTR retrotransposons: the *Gypsy/Ty3* group, the *Copia/Ty1* group and the *Pao/BEL* group (Xiong et al. 1993). *Sinbad*, another LTR retrotransposon from *S. mansoni*, is a member

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of the *Pao/BEL* group (Copeland et al. 2005a). Here, we investigated the possibility of the presence of homologues of *Boudicca* and *Sinbad* in genomes phylogenetically related to *S. mansoni* and genomes of host species parasitized by these schistosomes, using Southern hybridization analyses. Six platyhelminths were examined including the three major schistosome species, *S. mansoni*, *S. japonicum*, and *S. haematobium*, two non-schistosome trematodes, *Fasciola hepatica* and *Fascioloides magna*, and a (free-living) planarian, *Dugesia dorotocephala*. *D. dorotocephala*, a member of the class Turbellaria, served as a representative of platyhelminths phylogenetically distant from the schistosomes. The liver flukes *F. hepatica* and *F. magna*, like schistosomes, are members of class Trematoda, order Digenea. Unlike the schistosomes, flukes of this family are monocious, live as adults within the ducts of the liver rather than the blood vessels, and enter the host through the ingestion of vegetation contaminated with encysted larval forms known as metacercariae. *F. hepatica*, the sheep liver fluke, is primarily a parasite of sheep and other domesticated ruminants, but is also known to infect humans. *F. magna*, the deer liver fluke (renowned for its remarkable size and also known as the “giant fluke” and the “large American liver fluke”) is a wildlife parasite of deer, but can also infect other large ruminants, including cattle and sheep (Swales 1936). Further, to investigate host-parasite mediated horizontal transfer as a source of these elements, the genomes of three snail hosts corresponding to each of the major schistosomes, *Biomphalaria glabrata*, *Oncomelania hupensis*, and *Bulinus truncatus*, were examined in the same Southern blot as the six flatworms. *Homo sapiens* genomic DNA was also included in the analysis, since humans are the mammalian definitive host to all three major schistosome species.

#### MATERIALS AND METHODS

**Source of DNA** - Sources of genomic DNA used in the study were as follows. *S. mansoni*: cercariae (Puerto Rican strain), *S. japonicum*: adult worms (Chinese strain, Anhui Province), *S. haematobium*: adult worms (Egyptian strain), *O. hupensis*, *B. truncatus*, and *B. glabrata*: adult snails, *H. sapiens*: buffy coat isolated from non-schistosome infected human whole blood (provided by Fran Krogstad, Tulane University), *F. hepatica*: adult worms from Okeechobee, FL, US (provided by Dr Ray Kaplan, University of Georgia, Athens, GA, US) and Sydney, Australia (provided by Dr John Dalton, University of Technology, Sydney, Australia), *F. magna*: adult worms isolated from wild deer, central Florida, US (provided by Dr Ray Kaplan), and *D. dorotocephala*: adult worms (Carolina Biological Supply Co., Burlington, NC, US).

**Extraction of genomic DNA** - Samples preserved in ethanol were vacuum dried to remove the ethanol before extraction of genomic DNA. Genomic DNAs were extracted from the samples using the AquaPure Genomic DNA Purification system (Bio-Rad). In brief, tissue was lysed overnight at 55°C in a solution containing Proteinase K and, then treated with RNase for 30 min at 37°C followed by 45 min at 65°C. For whole blood, serum and most red blood

cells were removed by centrifugation and isolation of the buffy coat. The buffy coat was then also subjected to treatment with a red blood cell lysis solution (Bio-Rad). Following RNase treatment, samples were cooled, after which protein was removed by precipitation with salt. The supernatant was removed and DNA was precipitated with isopropanol, washed with ethanol, hydrated in sterile distilled water, and its concentration and purity determined by spectrophotometric analysis at OD<sub>260</sub> and OD<sub>280</sub>.

**Gene probes, labeling of gene probes with <sup>32</sup>P-dCTP, Southern hybridization** - The *Boudicca*-specific probe was produced by PCR amplification using the Le Paslier et al. (2000) bacterial artificial chromosome BAC 53-J-5 as a template and 5'-AACTGCAGATGCACGGAATCA CCGACT (forward) and 5'-GCTCTAGACTAAGATT CAGTCGGCAGATGC (reverse) primers, with restriction sites for *Pst* I and *Xba* I added to facilitate cloning into plasmid vectors. The probe, targeting part of the *gag* gene of *Boudicca*, was 385 bp in length, spanning residues 694-1078 of the 5858 nucleotides of the 53-J-5 copy of *Boudicca* (AY662653). Identity was confirmed by sequencing. The *Sinbad* probe was obtained by PCR amplification of a fragment of the LePaslier et al. (2000) bacterial artificial chromosome BAC 30-H-16 using the primers 5'-CGCGGATCCAAGAGAAAAACCTTGATAGAC and 5'-CCGGAATTCCTGTCTGAAGATAAAAGAGC. This fragment was cloned into pBluescript and its identity was confirmed by sequencing (Accession AY871176). The probe spans residues 2457 to 2823 of the BAC 33-N-3 copy of the *Sinbad* retrotransposon, the region of *pol* between the conserved protease and reverse transcriptase domains (Copeland et al. 2005a). The pSM389 probe is a 3.1 kb fragment encompassing the 3' end of the intergenic spacer region and the 5' end of the 18S exon of the ribosomal RNA gene of *S. mansoni* (Vieira et al. 1991). This probe has been shown previously to hybridize not only with *S. mansoni* genomic DNA, but also with several species of *Biomphalaria* (Knight et al. 1991). This schistosome ribosomal gene probe was employed here alongside the retrotransposon based probes in order to verify the integrity and quantity of the genomic DNAs in the different lanes of the Southern blots (Fig. 1).

*S. mansoni*, *S. japonicum*, *S. haematobium*, *O. hupensis*, *B. truncatus*, *B. glabrata*, *H. sapiens*, *F. hepatica*, *F. magna*, and *D. dorotocephala* gDNA (~10 µg/lane) were digested with *Hind* III. Digested gDNA, along with 100 ng each of the probes described above, was size fractionated by electrophoresis through a 0.8% agarose gel, transferred to a nylon membrane (Zeta-Probe GT, Bio-Rad) by capillary action (Southern 1975), and UV-light cross-linked to the membrane.

The probes described above were labeled with <sup>32</sup>P radiolabeled dCTP using the RadPrime DNA Labeling System (Invitrogen). Five hundred nanograms of probe DNA were labeled according to the manufacturer's instructions. After pre-hybridization at 65°C in 0.25 M Na<sub>2</sub>HPO<sub>4</sub>, 7% SDS, 1 mM EDTA for 15 min, the radiolabeled probe was added to 20 ml of fresh 0.25 M Na<sub>2</sub>HPO<sub>4</sub>, 7% SDS, 1 mM EDTA and membranes were allowed to hybridize at 65°C in this solution overnight for high strin-

gency hybridizations and 42°C overnight for low stringency hybridizations. The membranes were then washed twice in 40 mM Na<sub>2</sub>HPO<sub>4</sub>, 5% SDS, 1mM EDTA for 1 h at 65°C (high stringency hybridizations) or 42°C (low stringency hybridizations) followed by two washes in 40 mM Na<sub>2</sub>HPO<sub>4</sub>, 1% SDS, 1mM EDTA, for 1 h each (65°C for high stringency hybridizations and 42°C for low stringency hybridizations). The membranes were then rinsed briefly in 2× SSC and placed in clear plastic sheet protectors attached to BioMax amplification screens (Kodak). The signals from the <sup>32</sup>P-labeled gene probes were detected using x-ray film (Fuji).

Separate blots were used for the *Sinbad* and *Boudicca* hybridizations, and both these blots were subsequently stripped and probed with pSM389 as a positive control. The blots were stripped as follows. After briefly rinsing the membranes in 2 × SSC, the membranes were washed four times by immersion in 0.1% SDS, 0.1 × SSC at 100°C followed by washing (37°C, rocking) for 15 min as the solution cooled. No radioactivity was detected from the membranes by Geiger counter after these four washes.

**EST searches** - EST searches for *Boudicca* and *Sinbad* elements in the *S. japonicum* transcriptome were carried out using the complete sequence of each retrotransposon (accession number AY662653 for *Boudicca* and AY506538 for *Sinbad*). The search parameters were set to search the EST database of *S. japonicum* only, with “Schistosoma japonicum[organism]” entered into the “Limit by entrez entry” box of the “options for advanced blasting” section.

## RESULTS

***Sinbad*-like elements present in *S. haematobium*** - Standard (high stringency) Southern hybridization was used to search for *Sinbad*-like retrotransposons in the genomes of *S. japonicum* and *S. haematobium*, in the genomes of helminths phylogenetically related to these schistosomes, and in the genomes of host organisms parasitized by schistosomes (human and snails). The pattern of restriction fragments of *S. mansoni* genomic DNA was consistent with that of previous experiments. The *Sinbad* element was found only in *S. mansoni* and *S. haematobium*. Its presence was not evident in the genomes of any of the other flatworms, or in the snail hosts, or in *H. sapiens* (Fig. 2, the signal seen in the *Dugesia* lane is extended radiation from the positive control lane’s signal.) The *S. haematobium* signal showed a clear clustering in the ~11-14 kb range, whereas the signal in *S. mansoni* was intense to large fragments of > 15 kb in size (Fig. 2), a reflection of the general *Hind* III digestion pattern for these genomes (Fig. 1B).

In the 1-4 kb range, the patterns shown by the two species were similar (Fig. 2). *Hind* III is not expected to cut within *Sinbad*, as indicated by the fully sequenced *Sinbad* copy found in Le Paslier et al. (2000) BAC library clone 33-N-3 (Copeland et al. 2005a) but has been found to cut once in a *Sinbad* fragment found at the end of the BAC 30-H-16 clone of Le Paslier et al. (2000).

***Boudicca*-like elements present in *S. haematobium*** - These same ten genomes were examined for the presence

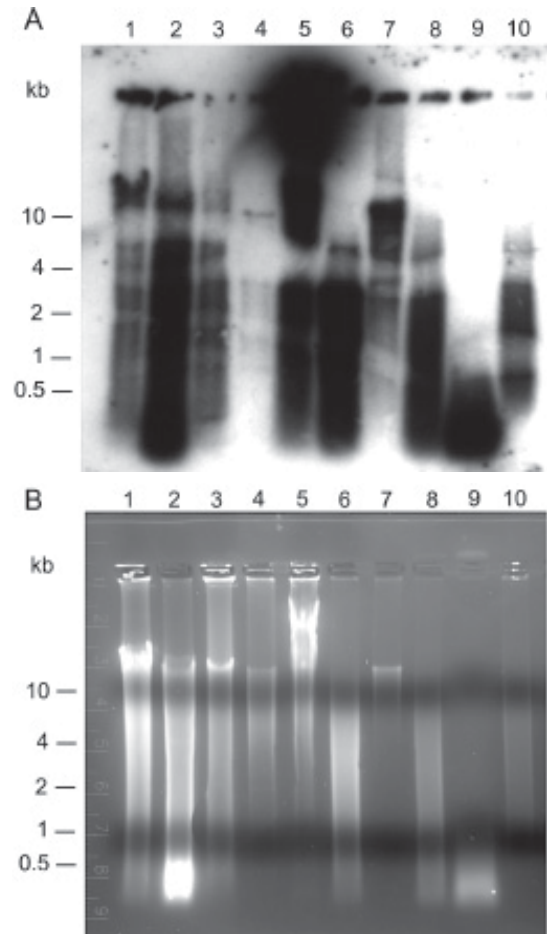


Fig. 1A: Southern hybridization analysis of *Hind* III digested genomic DNA from 10 genomes, probed with a radiolabeled *Schistosoma mansoni* ribosomal gene specific probe and exposed to X-ray film for 14 days. Size scale is shown to the left, in kilobases. Lanes - 1: *Oncomelania hupensis*; 2: *Bulinus truncatus*; 3: *Biomphalaria glabrata*; 4: *Homo sapiens*; 5: *S. mansoni*; 6: *S. japonicum*; 7: *S. haematobium*; 8: *Fasciola hepatica*; 9: *Fascioloides magna*; 10: *Dugesia dorotocephala*; B: ethidium stained gel of *Hind* III digested genomic DNA from 10 genomes used for the blot depicted in (A). Size scale is shown to the left, in kilobases. Lanes - 1: *O. hupensis*; 2: *B. truncatus*; 3: *B. glabrata*; 4: *H. sapiens*; 5: *S. mansoni*; 6: *S. japonicum*; 7: *S. haematobium*; 8: *F. hepatica*; 9: *F. magna*; 10: *D. dorotocephala*.

of a *Boudicca*-like retrotransposon, also using standard Southern hybridization. As with *Sinbad*, only *S. mansoni* and *S. haematobium* were found to be positive for *Boudicca* (Fig. 3). Both *S. mansoni* and *S. haematobium* exhibited areas of strong signal at ~8-11 kb and ~3 kb (Fig. 3). On the basis of the fully sequenced copy of *Boudicca* (from BAC 53-J-5), *Hind* III is expected to cut twice within the retrotransposon, at positions 4470 and 4793 of the 5858 bp element, yielding a ~300 bp product. The *S. haematobium* signal showed a distinct band at ~500 bp (Fig. 3), and *S. mansoni* gave strong signals for smaller fragments. As this copy of *Boudicca* exhibits several mutations (Copeland et al. 2003), the *Hind* III restriction map may differ in other *Boudicca* copies.

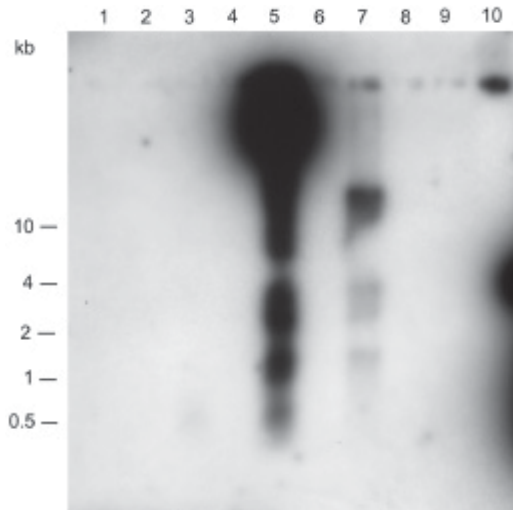


Fig. 2: Southern hybridization analysis of *Hind* III digested genomic DNA from 10 genomes, probed with a radiolabeled *Sinbad* specific probe and exposed to X-ray film for 14 days. Size scale is shown to the left, in kilobases. Lanes - 1: *Oncomelania hupensis*; 2: *Bulinus truncatus*; 3: *Biomphalaria glabrata*; 4: *Homo sapiens*; 5: *Schistosoma mansoni*; 6: *S. japonicum*; 7: *S. haematobium*; 8: *Fasciola hepatica*; 9: *Fascioloides magna*; 10: *Dugesia dorotocephala*.

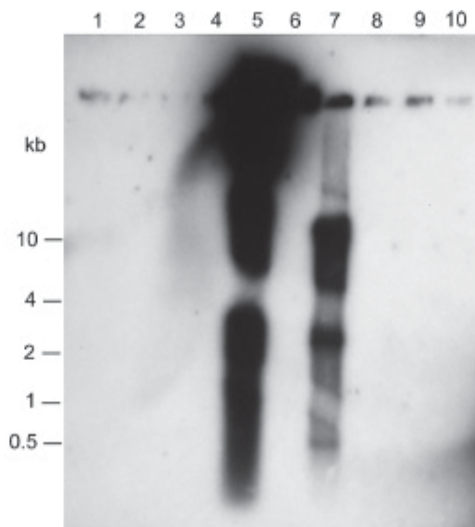


Fig. 3: Southern hybridization analysis of *Hind* III digested genomic DNA from 10 genomes, probed with a radiolabeled *Boudicca* specific probe and exposed to X-ray film for 10 days. Size scale is shown to the left, in kilobases. Lanes - 1: *Oncomelania hupensis*; 2: *Bulinus truncatus*; 3: *Biomphalaria glabrata*; 4: *Homo sapiens*; 5: *S. mansoni*; 6: *S. japonicum*; 7: *Schistosoma haematobium*; 8: *Fasciola hepatica*; 9: *Fascioloides magna*; 10: *Dugesia dorotocephala*.

*S. mansoni* ribosomal gene probe - As a positive control, a probe based on the 18S ribosomal subunit and preceding intergenic spacer of *S. mansoni* (Ali et al. 1991) was used in a final standard Southern hybridization. This probe, known as pSM389, has been shown previously to hybridize not only with *S. mansoni* genomic DNA, but also with several species of *Biomphalaria* (Knight et al. 1991). We hypothesized that the conserved nature of ri-

bosomal genes would allow this probe to be used as a positive control for all of the genomes examined in this study. The probe did indeed hybridize to all species, with strong signals for the trematodes and snails and a weak but discernible signal for the human DNA (Fig. 1a). From these results, it can be concluded that the gDNA of these ten species transferred successfully to the blot, and that negative results for species such as *S. japonicum* do not reflect a lack of cross-linked DNA.

*Low stringency hybridization suggests Boudicca-like elements in B. truncatus* - In addition to the high stringency Southern hybridizations described above (overnight hybridization at 65°C), low stringency hybridizations (overnight hybridization at 42°C) were performed on the same blots with both the *Boudicca* and *Sinbad* probes. The *Sinbad* low stringency results resembled the high stringency results shown in Fig. 2, with only the *S. mansoni* and *S. haematobium* lanes exhibiting a positive signal (not shown). The *Boudicca* blot, however, exhibited a pattern significantly different from that of the high stringency hybridization (Fig. 4). Though the *S. mansoni* and *S. haematobium* lanes exhibited the strongest signals, the next strongest signal, unexpectedly, was found in the *B. truncatus* lane. The signal for *Bulinus* (the snail host of *S. haematobium*) was stronger than that of *S. japonicum*, almost as strong as that of *S. haematobium*, and significantly stronger than any of the signals from the other genomes examined.

In all three blots, the strong signal toward the top of the blot seen in *S. mansoni* but not in *S. haematobium* reflects the general pattern of *Hind* III digested *S. mansoni* genomic DNA, with a large proportion of relatively undigested DNA (large fragments > 15 kb in size) compared to other genomes examined (Fig. 1B). Likewise, the strong signal around 11-14 kb in *S. haematobium* reflected the general restriction pattern for that schistosome.

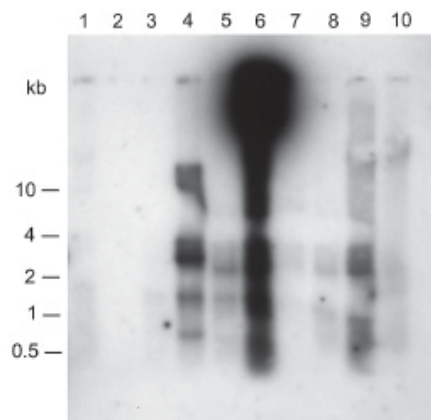


Fig. 4: low stringency Southern hybridization analysis of *Hind* III digested genomic DNA from 10 genomes, probed with a radiolabeled *Boudicca* specific probe and exposed to X-ray film for 30 days. Size scale is shown to the left, in kilobases. Lanes - 1: *Dugesia dorotocephala*; 2: *Fascioloides magna*; 3: *Fasciola hepatica*; 4: *Schistosoma haematobium*; 5: *S. japonicum*; 6: *S. mansoni*; 7: *Homo sapiens*; 8: *Biomphalaria glabrata*; 9: *Bulinus truncatus*; 10: *Oncomelania hupensis*.

## DISCUSSION

Of the three major species of schistosomes, *S. haematobium* is the least understood in terms of genome sequence. Until now, no LTR retrotransposons have been found in *S. haematobium*, though the presence of the (SINE-like) Sma elements (Ferbeyre et al. 1998) indicates that non-LTR retrotransposons are probably present. This study represents the first evidence of the presence of LTR retrotransposons in the genome of *S. haematobium*. Given the general paucity of genomic data for *S. haematobium*, and the fact that *Boudicca* and *Sinbad* elements make up over 4% of the *S. mansoni* genome (Copeland et al. 2005b), these findings significantly advance the exploration of the genome of this important human parasite. In addition, transposable elements are of interest for their possible utility in applications related to both sequencing and manipulation of genomes, including transgenesis and gene knockouts, generation of cell culture lines, generation of mutant libraries, and increased accuracy in whole genome shotgun sequencing (Izsvak et al. 1997, Handler 2001, Hirochika 2001, Tammi et al. 2003). Knowledge of these elements from *S. haematobium* will aid in the process of molecular genetic characterization of the *S. haematobium* genome.

*S. haematobium* is considered to be much more closely related to the other African species, *S. mansoni*, than to *S. japonicum*, an Asian species. In addition, evidence based on ribosomal RNA gene sequence, gene order of the mitochondrial genomes, and C-banding patterns of nuclear genome karyotypes suggests that *S. japonicum* is ancestral to both *S. mansoni* and *S. haematobium* (Snyder & Loker 2000, Le et al. 2000, Hirai et al. 2000). The presence of *Sinbad* and *Boudicca* in *S. haematobium* and *S. mansoni*, but not *S. japonicum*, is consistent with this phylogeny, and could have been brought about by the acquisition of the elements after the split from the *S. japonicum* lineage but before the divergence of *S. mansoni* and *S. haematobium*. Alternatively, ancestral elements of *Boudicca* and *Sinbad* could have been present in a basal ancestral schistosome, and subsequently diverged into different elements or were lost from the *S. japonicum* genome after the split between the *S. japonicum* and *S. mansoni/S. haematobium* lineage.

If these retrotransposons had ancient origins in the ancestors of *S. mansoni*, it would be expected that the genomes of related species would also include at least vestigial forms of these elements. However, neither the moderately related turbellarian and trematode genomes, nor the closely related *S. japonicum* genome appeared to contain these elements. Thus far, no *Boudicca* elements have been identified in *S. japonicum*, and the only *Pao*-like element from *S. japonicum*, *Tiao* (Fan & Brindley 1998) has not been fully characterized. A BLAST search of the *S. japonicum* EST database with the complete sequence of the *Boudicca* retrotransposon (AY662653) did reveal a match to a reverse transcriptase (BU723196) with moderate similarity (bit score = 97) to that of *Boudicca*, indicating that *S. japonicum* may encode, and transcribe, *Gypsy*-group LTR retrotransposons from the *CsRn1/Kabuki* clade. This clade, of which *Boudicca* is a member, con-

sists of elements so far found only in the genomes of *Bombyx mori*, *Chlonorchis sinensis*, *Paragonimus westermani*, and *S. mansoni*, and is defined by a unique cysteine box structure (Copeland et al. 2005b). A corresponding BLAST search using the complete sequence of the *Sinbad* retrotransposon (AY506538) failed to yield any significant matches. More detailed characterization of *Tiao* and any *S. japonicum* elements from the *CsRn1/Kabuki* clade, would help to elucidate this phylogeny.

The results of this study indicate either a relatively recent origin (after the split between the *S. japonicum* and *S. mansoni/S. haematobium* lines) for the *Boudicca* and *Sinbad* retrotransposons via horizontal transmission, the loss of these elements in the *S. japonicum* lineage, or a high degree of divergence of ancestral retrotransposons of *Boudicca* and *Sinbad* after the split between the *S. japonicum* and *S. mansoni/S. haematobium* lineage. Previously, elements closely related to *Boudicca* have been found only in platyhelminths and insects (Bae et al. in press, Copeland et al. 2005b), whereas elements known to be closely related to *Sinbad* have been limited to schistosomes and fish (Copeland et al. 2005a). The presence of a robust positive signal for *Sinbad* in *S. haematobium* but absence in *S. japonicum*, the other platyhelminths, and the host genomes, even under low stringency conditions, supports the possibility of a horizontal transfer event from some other genome after the divergence of *S. japonicum* from the *S. mansoni/S. haematobium* line. The finding of a *Sinbad*-like element in the zebrafish *Danio rerio*, combined with the presence of other *Pao*-like elements in the genomes of aquatic, but not terrestrial, vertebrates (Copeland et al. 2005a), suggests a possible origin for *Sinbad* in fish.

The robust positive signal for *Boudicca* in *Bulinus* under low stringency conditions raises the possibility of a horizontal transfer event of an ancestral *Boudicca*-like element from an ancestral snail to an ancestral schistosome. Especially intriguing is the strength of the signal; the *Bulinus* signal was not only stronger than that of the platyhelminth genomes but was also stronger than that of the host genome of *S. mansoni*, *Biomphalaria*. The evolutionary path suggested by this pattern is a jump from an ancestral *Bulinus* snail genome to an ancestral *S. haematobium/S. mansoni* genome, which in turn points to *Bulinus* or an ancestor of *Bulinus* as the original host of this ancestral schistosome. One caveat, considering the low stringency conditions of the experiment, is that it is plausible for an element only moderately related to *Boudicca* to generate a strong signal if it were very high in copy number. The fact that the positive signal in *Bulinus* only emerged under low stringency conditions therefore necessitates further inquiry as to the identity of any *Boudicca*-like elements in this genome, and invites further studies of *Boudicca*-like elements in other snail genomes.

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