Use of Cell-Free Circulating Schistosome DNA in Serum, Urine, Semen, and Saliva To Monitor a Case of Refractory Imported Schistosomiasis Hematobia

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This case of imported refractory schistosomiasis has highlighted the usefulness of cell-free parasite DNA as a diagnostic marker to assess active schistosome infection. In contrast to the rapid disappearance of ova in urine, parasite DNA remained persistent in several other specimen types even after the fourth treatment with praziquantel. This result was consistent with the presence of morphologically intact ova in bladder biopsy samples and with the corresponding symptoms.

CASE REPORT

In January 2009, a previously healthy Japanese man, 21 years of age, sought medical attention after an approximately 2-month history of hematuria, discomfort during urination, and hematospermia. He had returned from a 5-month trip (May to October 2008) around various African countries. During the trip, he bathed in Jinja (Uganda), Lake Malawi (Malawi), and the Niger River and specifically developed itchy skin after swimming in the Dogon region of Mali. Urinalysis revealed the presence of Schistosoma haematobium ova. The patient was treated with praziquantel (PZQ; 40 mg/kg of body weight twice a day [b.i.d.] orally [p.o.] for two consecutive days). After the first PZQ treatment, the patient had a fever of 38°C, a relatively high white blood cell (WBC) count (10,000/mm³), and eosinophilia (30%).

For detection of ova, sediment from 10 ml of urine and 250 μl of semen was examined by microscope. Parasite ova were detected in urine on day 1 of the first PZQ treatment course, and detection results were negative after that. On the other hand, ova in semen were detected until 101 days after the first PZQ treatment.

Cell-free schistosome DNA in bodily fluids was detected by conventional PCR and/or sequence capture-PCR. The genetic examination for parasite DNA in the patient’s specimen was approved by the Bioethics Committee of Dokkyo Medical University (approval no. 1969), and the patient’s consent was obtained. Prior to DNA extraction, 3.5 ml of urine was concentrated to 140 μl using an Amicon Ultra-15 centrifugal filter system with an UltraCel-100K membrane (Millipore Ireland Ltd., Cork, Ireland). DNA was extracted from concentrated urine, serum, and supernatant of semen (140 μl) each using a QIAamp viral RNA minikit (Qiagen Sciences), and DNA from the sediment of 500 μl of semen was extracted using a NucleoSpin tissue system (Macherey-Nagel GmbH & Co. KG, Düren, Germany) according to the manufacturers’ instructions. Approximately 2 ml of the saliva sample was collected using an Oragene-DNA self-collection kit (DNA Genotek Inc., Canada); DNA was extracted according to the manufacturer’s instructions. The primer pair CF (5'-GATCGTA-AATTTGGA/TACTGC) and CR (5'-CCAACCATACACATATG ATG) was designed to amplify a part of the schistosome mitochondrial cytochrome c oxidase subunit 1 (CO1) gene, which is common to at least 4 human schistosome species (S. mansoni [253 bp] and S. haematobium, S. japonicum, and S. mekongi [254 bp]). The primer pair ShF (5'-AGTCGTGCAGATTCTAAAAC) and CR was designed to amplify S. haematobium CO1 (365 bp), and the primer pair SmF (5'-TCCCTATCAATTTGAG AGG) and CR was designed to amplify S. mansoni CO1 (479 bp) (1). Sequence capture-PCR is the combination of purifying and concentrating methods for target nucleic acid in clinical samples and PCR. As described in the literature (2–5), crude DNA samples were purified and concentrated using both 5'-biotinylated CF and CR and magnetic beads with streptavidin, and then PCR was performed. We were able to amplify S. haematobium DNA in the patient’s bodily fluids; however, S. mansoni DNA was not amplified. Parasite DNA in bodily fluids was detected even 1 month after the fourth PZQ treatment (303 days after the first treatment).

Using cystoscopy, we identified massive polypoid lesions in the urinary bladder wall, which gradually decreased in number with repeated PZQ treatments (Fig. 1A). However, histopathological findings from examinations of bladder biopsy sites of polypoid lesions with petechiae revealed parasite ova with intact cells even 2 months after the third treatment (255 days after the first treatment) (Fig. 1B). Finally, 184 days after the fourth PZQ treatment (457 days after the first treatment), most of the lesions diminished to trace levels and the subjective symptoms were resolved; thereafter, the patient was routinely treated with praziquantel (PZQ; 40 mg/kg of body weight twice a day [b.i.d.] orally [p.o.] for two consecutive days).
examined in follow-up checks. The therapeutic process and examination results are summarized in Table 1.

The current diagnostic methods of schistosomiases rely on an ovum-related phenomenon: detection of ovum- and parasite-specific antibodies and ovum-induced pathology. Although detection of ova is still the gold standard for diagnosing schistosomiasis, it is difficult to detect long-standing/chronic infection and low-level infection in patients after repeated treatments; moreover, detection of ova is ineffective in the early stage of infection (prepatent period). The cell-free circulating schistosome DNA consists of the fragments of parasite-derived DNA that exist in the host’s bodily fluids. It was detected in plasma/serum and urine of schistosomiasis patients (6, 7, 8, 9). In our study, the primer pairs (CF and CR, ShF and CR, and SmF and CR) had been originally designed to amplify the mitochondrial CO1 gene, which enabled us to differentiate

<table>
<thead>
<tr>
<th>Days post-PZQ treatment</th>
<th>Detection of ova</th>
<th>Detection of occult blood (urine)</th>
<th>Detection of DNA (PCR/sc-PCR)</th>
<th>Corresponding cystoscopy panel in Fig. 1A</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>Urine</td>
<td>Semen</td>
<td>Bladder biopsy specimen</td>
<td>Serum</td>
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<tr>
<td>Treatment 1</td>
<td>1 day post</td>
<td>100 days post</td>
<td>52 days post</td>
<td>39 days post</td>
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<td>102</td>
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<td>457</td>
<td>355</td>
<td>254</td>
<td>184</td>
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a (−), not detected; (+), detected; PZQ, praziquantel; sc-PCR, sequence capture-PCR; ND, not done.

b PZQ treatment 1 consisted of 40 mg/kg of body weight/day for 2 days; PZQ treatment 2 consisted of 40 mg/kg/day for 2 days; PZQ treatment 3 consisted of 40 mg/kg/day for 2 days; PZQ treatment 4 consisted of 60 mg/kg/day for 2 days.

c Results for ova in semen at day 101 were first negative, however, later, when reexamined retrospectively, the results were positive.
four species of human schistosomes (1). It was confirmed that the patient was infected with *S. haematobium* and was not infected with *S. mansoni*.

Because schistosomiasis (japonica) has been eliminated from Japan, it is unlikely that the patient will get reinfected now that he has returned home. Parasite ova in urine, a common testing material for schistosomiasis hematoxia, were detected for only 1 week after the first PZQ treatment. On the other hand, ovum was detected in semen even 101 days after the first PZQ treatment and indicates the significance of semen as a testing material for schistosomiasis hematoxia. We had expected the cell-free parasite DNA to disappear soon after PZQ treatment along with the ova. However, it was detected in the bodily fluids even after 1 month after the fourth PZQ treatment (303 days after the first treatment). The idea of the reliability of persistent parasite DNA was supported by the presence of morphologically intact parasite ova in the bladder biopsy sample, pathology, symptoms, and other signs; the patient reported subjective symptoms of discomfort with urination even 2 months after the third PZQ treatment (255 days after the first treatment) and sometimes noticed hematuria during therapy.

It has been reported that schistosome DNA was detected in serum for as long as 10 to 19.3 weeks after a single PZQ treatment (8, 10). One possible explanation for the lingering worm DNA detection may have been unsatisfactory treatment. Immature ova/worms are refractory to PZQ treatment (11–14). Multiple treatments with PZQ are needed with intervals between the treatments to allow maturation of immature worms/ova (2 to 4 weeks) (13–15). In the present case, the persistent parasite DNA was detected even after repeated PZQ treatments and it indicates that this was a refractory case.

To date, there have been no reports on PZQ resistance in areas of schistosomiasis hematoxia endemicity (15–17). On the other hand, several cases refractory to repeated PZQ administration have been reported in countries where the disease is nonendemic, i.e., imported cases (18–20). It is speculated that schistosomiasis patients, who are rare in countries where the disease is nonendemic, tend to be followed up by detailed examinations, such as cystoscopy and bladder biopsy sampling, in addition to detection of ova, thus enabling identification of refractory cases. By the conventional gold standard, the examination of ova in urine, the present case would have been considered cured soon after the first PZQ treatment. Chronic infection with *S. haematobium* is carcinogenic to humans (21, 22). In the present study, we have demonstrated the limitations of detection of ova as a method of therapeutic evaluation. Cell-free schistosome DNA may be a potential tool to monitor active worms/ova to reduce the risk of cancer triggered by prolonged *S. haematobium* infection because of inaccurate diagnosis. Therefore, further studies of novel diagnostic techniques such as cell-free schistosome DNA detection to determine/assess active infection are required.

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We declare no competing financial interests.

**REFERENCES**


