Korean J Parasitol. Vol. 48, No. 4: 343-346, December 2010 DOI: 10.3347/kjp.2010.48.4.343

## Prevalence of *Haplorchis taichui* in Field-Collected Snails: A Molecular Approach

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**Abstract:** The prevalence of the cercarial stage of an intestinal trematode, *Haplorchis taichui*, in thiarid snails (Gastropoda: Thiaridae) was investigated using light microscope and species-specific PCR procedures. A total of 988 snails were collected from Mae Taeng district, Chiang Mai province, northern Thailand, which comprised of 3 species; *Melanoides tuber-culata, Tarebia granifera*, and *Thiara scabra*. The overall prevalence of pleurolophocercous cercariae was 21.7% as determined by the morphology. For genetic detection of *H. taichui* infection in snails, 2 primers Hapt\_F (5<sup>´</sup>-GGCCAACGCAA-TCGTCATCC-3<sup>´</sup>) and Hapt\_R (5<sup>´</sup>-GCGTCGGGTTTCAGACATGG-3<sup>´</sup>), were used. The genomic DNA of *H. taichui*, which was used as a positive control, gave an amplification of the 256 bp fragment. The overall prevalence of *H. taichui* from specific PCR was 9.7%. The proportion of *H. taichui* among the pleurolophocercous cercariae in this study was 44.9%.

Key words: Haplorchis taichui, pleurolophocercous cercaria, thiarid snail, PCR, Thailand

The intestinal trematode, *Haplorchis taichui*, is a medically important parasite infecting humans and livestock. Haplorchiasis has become one of the major public health threats in Asia and in parts of Africa and the Americas [1-3]. *H. taichui* is the most frequently reported species among the minute intestinal flukes from eastern Asia, including Thailand [4-7]. This parasite has an aquatic life cycle, using freshwater snails as the first [8] and cyprinid fish as the second intermediate hosts, with definitive hosts being fish-eating mammals [9,10]. Previous reports showed that metacercariae of 2 heterophyids, *H. taichui* and *Haplorchoides* sp., had high prevalence of infection in cyprinoid fish collected from Chiang Mai and Lamphun provinces of northern Thailand [11]. This was also found in a later study where a high prevalence (83.9%) of these 2 heterophyid trematodes was also recorded from the same area [10].

Many species of heterophyid trematodes have been reported from pleurolophocercous cercariae, such as *H. taichui, Haplorchis pumilio*, and *Stellantchasmus falcatus* [12-14]. About 100 species of snails have been reported to act as intermediate hosts for the trematodes especially in thiarid snails which harbor the larvae

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of intestinal trematodes, including H. taichui [12,15].

Traditional methods of detecting cercarial infections are usually performed by exposing the snails to light or dissection. It is difficult to determine the parasite stages to the species level since they appear morphologically very similar to other trematodes [16]. In addition, only a very small percentage of snails can be found to be infected when testing shed cercariae [17]. Specific identification of cercariae from field collected snails is difficult, but necessary for epidemiological studies.

The development of a molecular approach for cercarial detection in infected snails is necessary and will be useful, such as PCR. Various conventional PCR assays have been developed to detect *H. taichui* DNA in feces, definitive hosts, and intermediate hosts. Van et al. [18] separated larvae and adult stages of *H. taichui* and *H. pumilio* using ITS-2 gene sequencing. Wongsawad et al. [6] established the specific primer of *H. taichui* by generating the 256 bp amplicon which had shown a positive result. The objectives of our study were to determine the prevalence of *H. taichui* infections in snails by using PCR to gain information on the epidemiology of parasitic infections, and developing effective control measures.

Field collections were made during March 2009 to February 2010. Thirty individuals of each thiarid snails were collected by handpicking and scoop methods. Seven sites were selected based on the relevance of public health concerns in and around the Mae Ngad Somboonchon dam in Mae Taeng district, Chiang Mai province, such as rice paddy, irrigation canal, dam, and

Received 5 October 2010, revised 16 November 2010, accepted 18 November 2010.

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Fig. 1. Map showing the 7 collecting sites in Mae Taeng district (★), Chiang Mai province, Thailand. 1. Rice paddy (N21<sup>°</sup>16.486<sup>′</sup> E0.4<sup>°</sup>09.359<sup>′</sup>), 2. River (N21<sup>°</sup>17.380<sup>′</sup> E0.50<sup>°</sup>02.067<sup>′</sup>), 3. Mae Ngad Somboonchon dam (N21<sup>°</sup>19.725<sup>′</sup> E0.50<sup>°</sup>03.645<sup>′</sup>), 4. Rice paddy (N21<sup>°</sup>13.134<sup>′</sup> E0.50<sup>°</sup>07.982<sup>′</sup>), 5. Irrigation canal (N21<sup>″</sup>12.864<sup>′</sup> E0.50<sup>°</sup> 08.488<sup>′</sup>), 6. Stream (N21<sup>″</sup>12.992<sup>′</sup> E0.50<sup>°</sup>07.862<sup>′</sup>), and 7. Khek river (N19<sup>°</sup>08.775<sup>′</sup> E 0.99<sup>°</sup>00.586<sup>′</sup>).

river (Fig. 1) during each of 3 seasons (hot-dry, rainy, and cooldry). The snail specimens were identified using Brandt's taxonomic key [19] and were examined for cercarial infections by crushing. The types of cercariae were morphologically identified under a compound microscope, and only pleurolophocercous cercariae were preserved in 70% alcohol for DNA extraction.

The genomic DNA was extracted and purified from individual pleurolophocercous cercariae samples found in each snail using commercial GF-1 tissue DNA extraction kit (Vivantis, Malaysia) according to the manufacturer's instructions. The genomic DNA of each sample was eluded in 50  $\mu$ l of the elution buffer provided with the kit and stored at -20°C.

The *H. taichui* specific primers were tested for specificity prior to perform the detection of cercarial stage by attempting to amplify them with all 6 adult trematode DNAs, including *H. taichui*, *H. pumilio*, *S. falcatus*, *Centrocestus caninus*, *Haplorchoides* sp. and *Opisthorchis viverrini*. These mentioned worms are widely distributed with a high prevalence of infection in Chiang Mai province, Thailand as same as they have been reported frequently in a past decade [4,5].

PCR amplification for the test of specificity and detection of cercarial stages of *H. taichui* were also conducted using *H. taichui* specific primers. There were 5'-GGCCAACGCAATCGTCATCC-3' (Hapt\_F) as a forward primer and 5'-GCGTCGGGTTTCA-



Fig. 2. Seasonal prevalence of pleurophocercous cercariae using morphology, and *H. taichui* cercariae using PCR in the infected thiarid snails, collected in Mae Taeng district, Chiang Mai province of northern Thailand during March 2009 to February 2010.

GACATGG-3<sup>'</sup> as a reverse primer [6]. The PCR amplifications were carried out in a final volume of 20  $\mu$ l, including 1  $\mu$ l of DNA template of pleurolophocercous cercariae from each snail sample, 0.6  $\mu$ l of each primer (Hpt\_F and Hpt\_R), 0.3  $\mu$ l of MgCl<sub>2</sub>, dNTPs, and *Taq* DNA polymerase. The amplification procedure involved an initial denaturation step at 95°C for 5 min, then 35 cycle including denaturation at 95°C for 45 sec, primer annealing at 68°C for 45 sec, extension at 72°C for 1 min, and final extension at 72°C for 7 min. PCR products were analyzed after electrophoresis separation at 50 volt for 45 min on 2.5% agarose gels stained with ethidium bromide in TBE buffer. Gels were visualized by a Kodak Digital Camera (Gel Logic 100).

A total of 988 collected snails were identified to be 3 species, including *Melanoides tuberculata, Tarebia granifera,* and *Thiara scabra.* Pleurolophocercous cercariae were found in 214 snails, with an overall prevalence of 21.7%, as detected by the morphology. The highest prevalence of infection was found in the cool-dry season (November - February) (30.87%), followed by hot-dry season (March - May) (23.45%), and rainy season (May-October) (15.74%) (Fig. 2).

When *H. taichui* specific primers were applied in combination to DNA from all 6 parasites, only 256 bp specific fragment of *H. taichui* was generated and there was no cross reaction with any other tested trematodes (Fig. 3). This result confirmed the specificity of *H. taichui* specific primers which were appropriate for detection of cercariae in snails.

For the detection of *H. taichui* in snails, 256 bp fragments were generated in different pleurolophocercous cercariae (Fig. 4). Genomic DNA of *H. taichui* adult worm was used as the



Fig. 3. The 256 bp specific fragment (arrow) demonstrated the specificity of Hap\_F and Hap\_R specific primers, as confirmed by the positive result only in *H. taichui* DNA sample. Lane M, 100 bp molecular size markers; Lane Ht, *H. taichui*; Lane Hp, *H. pumilio*; Lane St, *S. falcatus*; Lane Cc, *Centrocestus caninus*; Lane Ha, *Haplorchoides* sp.; Lane Ov, *Opisthorchis viverrini*; Lane N, negative control.

positive control, and also gave an amplification product of 256 bp. This result provided the effective evidence by revealing that not only *H. taichui* that was developed from pleurolophocercous cercariae, but also there were other trematode species which were generated from the same cercarial type, like *H. taichui*. The overall prevalence of *H. taichui* infected snails was 9.7% (Fig. 2). The cool-dry season had the highest prevalence of 12.2%, followed by the hot-dry (9.5%) and rainy seasons (8.6%), respectively. The proportion of *H. taichui* among the pleurolophocercous cercarial population was 44.9% (96/214) (Fig. 5).

The prevalence of pleurolophocercous cercariae in naturally infected *M. tuberculata, T. granifera,* and *T. scabra* in this study was relatively high (21.7%), and higher than in the previous study (5.3%) because the crushing method which was employed in this study show more sensitivity and accuracy than those of shedding method that acquired previously [13]. Kumchoo et al. [4] reported high prevalence (91.4%) of *H. taichui* in cyprinoid fish from Mae Taeng district of Chiang Mai province. Nithikathkul and Wongsawad found a high infection rate of *H. taichui* and *Haplorchoides* sp. (83.90%) in this area [10].

Our results showed that a high level (44.9%) of pleurolophocercous cercarial populations infected in thiarid snails was *H. taichui*, whereas it can be proposed that the remainders can develop into other trematode species, such as *H. pumilio*, *S. falcatus*, and *Procerovum varium* [12-14,19,20]. From this, it seems that the prevalence of trematode infection in definitive and sec-



Fig. 4. Representative agarose gel PCR profiles of *Haplorchis taichui* isolated from thairid snails. Lane M, molecular size maker; Lane Ht, *H. taichui*; Lanes 1-16, DNA samples of pleurolophocercous cercariae infected in snails (Lanes 1-3, 7, 8, 11, 15,16; positive results of *H. taichui* PCR specific fragment), (Lanes 4-6, 9,10, 12-14: negative results of cercariae infected in snails); Lane N, negative control.



Fig. 5. Proportion of *Haplorchis taichui* cercariae infected in thiarid snails (n = 214), as determined by PCR.

ond intermediate hosts collected from these areas may be very high. This conclusion is also supported by the peculiar ecological conditions of water resources in Mae Taeng district, which represents a complex and diverse freshwater ecosystem. There are good conditions for the cercarial stage of *H. taichui* to develop in the molluscs and then fish which lead to perpetuation in their life cycle.

Freshwater snails are important for medical and veterinary public health points of view. About 100 species of snails have been reported to act as intermediate hosts, such as Thiaridae, and have been recorded to harbor larval trematodes [22]. Our study found that *M. tuberculata* and *T. granifera* serve as intermediate hosts of *H. taichui*. The PCR approach conducted in this study confirmed the high prevalence of *H. taichui*, comprising almost a half of the pleurolophocercous cercariae. This test is suitable for epidemiological studies, and control programs for the snail intermediate hosts, and also for the detection of human infections. An essential and immediate applica-

tion of this PCR method should be a metacercarial survey in fish of natural water resources in order to monitor infected areas. This may be a way for the long-term control of haplorchiasis in northern of Thailand.

## ACKNOWLEDGEMENTS

We greatly appreciate use of the Applied Parasitology Research Laboratory and Economic Plants Genome Research and Service Center, Department of Biology, Faculty of Science, Chiang Mai University, Thailand. Special thanks are given to Mr. Sukson Chuboon and Mr. Tanu Marayong for their assistance in collecting specimens. We also thank Prof. Duangduen Krailas and Ms. Wivitcuta Dechruksa for confirming the identification of snail specimens. We thank Dr. J.F. Maxwell for editing our manuscript.

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