**Strongyloides ratti**: In Vitro and In Vivo Activity of Tribendimidine

Jennifer Keiser1*, Kai Thiemann1, Yvette Endriss1, Jürg Utzinger2

1 Department of Medical Parasitology and Infection Biology, Swiss Tropical Institute, Basel, Switzerland, 2 Department of Public Health and Epidemiology, Swiss Tropical Institute, Basel, Switzerland

Abstract

**Background:** Strongyloidiasis is a truly neglected tropical disease, but its public health significance is far from being negligible. At present, only a few drugs are available for the treatment and control of strongyloidiasis.

**Methodology/Principal Findings:** We investigated the activity of tribendimidine against third-stage larvae (L₃) of *Strongyloides ratti* in vitro and against juvenile and adult stages of the parasite in vivo. *S. ratti* larvae incubated in PBS buffer containing 10–100 μg/ml tribendimidine died within 24 hours. A single 50 mg/kg oral dose of tribendimidine administered to rats infected with 1-day-old *S. ratti* showed no effect. The same dose administered to rats harboring a 2-day-old infection showed a moderate reduction of the intestinal parasite load. Three days post-exposure a significant reduction of the immature worm burden was found. Administration of tribendimidine at doses of 50 mg/kg and above to rats harboring mature *S. ratti* resulted in a complete elimination of the larval and adult worm burden. For comparison, we also administered ivermectin at a single 0.5 mg/kg oral dose to rats infected with adult *S. ratti* and found a 90% reduction of larval and 100% reduction of adult worms.

**Conclusion/Significance:** Tribendimidine exhibits activity against *S. ratti* in vitro and in vivo. The effect of tribendimidine in humans infected with *S. stercoralis* should be assessed.

Introduction

An estimated 30–100 million people are infected with *Strongyloides stercoralis*, the causative agent of strongyloidiasis, and yet this is a truly neglected tropical disease [1]. One explanation is that current diagnostic tools have limitations [2]. Whilst *S. stercoralis* mainly occurs in tropical and subtropical areas, endemic foci also occur in temperate regions such as Spain or the United States [3]. Serious clinical problems have been observed in *S. stercoralis*-infected patients who are immunocompromised due to a co-infection with human T-cell leukaemia virus type 1 (HTLV-1) or HIV, or corticosteroid-treated patients [4]. However, the global burden of strongyloidiasis is currently not known. The growing evidence that an infection with *S. stercoralis* is a risk factor for biliary tract cancer needs to be considered when estimating the burden of strongyloidiasis [5].

As with other nematodes (*Ascaris lumbricoides, hookworms* and *Trichuris trichiura*), and trematodes (e.g. *Schistosoma spp.*), there are only a few drugs available for the treatment and control of strongyloidiasis [6]. Albendazole and mebendazole were found to be safe, but multiple treatment courses repeated over several weeks were required to achieve acceptable cure rates [7]. Another benzimidazole, thiabendazole, is highly efficacious (two treatment courses of 25–50 mg/kg are commonly given for 3–4 days 2 weeks apart), but severe adverse events, including liver dysfunction and neuropsychiatric symptoms, have been observed [8]. Ivermectin, a semi-synthetic macrocyclic lactone, which was developed as a veterinary anthelmintic, is safe and efficacious, and is now the drug of choice for strongyloidiasis [9]. Ivermectin resistance in humans infected with *S. stercoralis* has not been reported thus far, but host and parasite-specific resistance to the drug has been reported in veterinary medicine [9].

Tribendimidine is an aminophenyldimidine derivative of amidan. Tribendimidine is safe and has a broad spectrum of activity against numerous nematode species, including *A. lumbricoides, Enterobius vermicularis* and the hookworms (*Ancylostoma duodenale* and *Necator americanus*) [10]. Tribendimidine has been approved by Chinese regulatory authorities in 2004 [10] and phase IV trials have been completed recently [11]. Efforts are ongoing to secure a western registration for tribendimidine, in order for the treatment to be considered for global soil-transmitted helminthiasis control usage. In our recent work, we have documented the in vivo activity of tribendimidine against a number of trematodes, namely the intestinal fluke *Echinostoma caproni* [12] and the two liver flukes *Clonorchis sinensis* and *Opisthorchis viverrini* [13].

Here, we investigated the in vitro activity of tribendimidine against third-stage larvae (L₃) of *Strongyloides ratti*. Moreover, we evaluated the dose-response relationships of single oral doses of tribendimidine against adult *S. ratti* harbored in rats, and assessed the in vivo activity of tribendimidine against different immature stages of the parasite in the rat model.

---

*Citation:* Keiser J, Thiemann K, Endriss Y, Utzinger J (2008) *Strongyloides ratti*: In Vitro and In Vivo Activity of Tribendimidine. PLoS Negl Trop Dis 2(1): e136. doi:10.1371/journal.pntd.0000136

*Editor:* Charles King, Case Western Reserve University School of Medicine, United States of America

*Received:* August 29, 2007; *Accepted:* October 29, 2007; *Published:* January 23, 2008

*Copyright:* © 2008 Keiser et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

*Funding:* This study was funded by the Swiss National Science Foundation (project no. PPOOA-114941; J. Keiser) and (project no. PPOOB-102883; J. Utzinger). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

*Competing Interests:* The authors have declared that no competing interests exist.

*E-mail: jennifer.keiser@unibas.ch*
In vivo studies

Freshly harvested *S. ratti* L$_3$ were washed 3 times with PBS buffer and incubated in 6-well microtiter plates (Costar) containing 4 ml PBS buffer (pH 7.3). 800 *S. ratti* L$_3$ were used for each control and experimental group. The worms were incubated in three serial drug dilutions of tribendimidine, i.e. 100, 10 and 1 mg/ml for up to 96 hours (h). Each experiment was carried out in triplicate and then repeated once. The control wells contained tribendimidine: 66.6% of *S. ratti* showed no movement when exposed to this concentration. The worms died within 24 h. No effect was observed with the lowest concentration (1 mg/ml) of tribendimidine: 66.6% of *S. ratti* were still active after an incubation period of 72 h. Ninety-six h post-incubation 50.4% of *S. ratti* showed no movement when exposed to this concentration, similar to the control group, where 60.0% of the worms were found to be inactive.

Effect of tribendimidine on adult *S. ratti*: dose-response relationship

Figure 1 shows the effect of tribendimidine on *S. ratti* rhabditiform larvae harvested from rat fecal samples as assessed by quantitative stool examination. In the first set of experiment (evaluating single 50, 100 and 200 mg/kg oral doses of tribendimidine) on 5 daily stool examinations, between 3300 (6 days post-exposure) and 10,000 (9 days post-exposure) rhabditiform larvae per gram of stool were estimated in the group of untreated and served as controls.

Statistical analysis

Statistical analyses were done with version 2.4.5 of Statsdirect statistical software (Statsdirect Ltd; Cheshire, UK). The effect of tribendimidine was assessed by comparing the mean number of *S. ratti* rhabditiform larvae in the stool and *S. ratti* adults and rhabditiform larvae in the intestine in the treatment group with the mean number of larvae and adults in the respective control group. The responses between the medians of the treatment and control groups regarding larvae and stool and larvae and adults in the intestines were analysed with the Kruskal-Wallis (KW) test. Differences in medians were considered to be significant at a significance level of 0.05.

Results

In vitro findings

Table 1 summarizes the effect of *S. ratti* L$_3$ after exposure to tribendimidine at different concentrations in vitro. *S. ratti* exposed to tribendimidine at 10 or 100 μg/ml contracted immediately and had a coiled shape appearance. The worms died within 24 h. No effect was observed with the lowest concentration (1 μg/ml) of tribendimidine: 66.6% of *S. ratti* were still active after an incubation period of 72 h. Ninety-six h post-incubation 50.4% of *S. ratti* showed no movement when exposed to this concentration, similar to the control group, where 60.0% of the worms were found to be inactive.

Effect of tribendimidine on adult *S. ratti*: dose-response relationship

Figure 1 shows the effect of tribendimidine on *S. ratti* rhabditiform larvae harvested from rat fecal samples as assessed by quantitative stool examination. In the first set of experiment (evaluating single 50, 100 and 200 mg/kg oral doses of tribendimidine) on 5 daily stool examinations, between 3300 (6 days post-exposure) and 10,000 (9 days post-exposure) rhabditiform larvae per gram of stool were estimated in the group of untreated and served as controls.

Statistical analyses were done with version 2.4.5 of Statsdirect statistical software (Statsdirect Ltd; Cheshire, UK). The effect of tribendimidine was assessed by comparing the mean number of *S. ratti* rhabditiform larvae in the stool and *S. ratti* adults and rhabditiform larvae in the intestine in the treatment group with the mean number of larvae and adults in the respective control group. The responses between the medians of the treatment and control groups regarding larvae and stool and larvae and adults in the intestines were analysed with the Kruskal-Wallis (KW) test. Differences in medians were considered to be significant at a significance level of 0.05.

Results

In vitro findings

Table 1 summarizes the effect of *S. ratti* L$_3$ after exposure to tribendimidine at different concentrations in vitro. *S. ratti* exposed to tribendimidine at 10 or 100 μg/ml contracted immediately and had a coiled shape appearance. The worms died within 24 h. No effect was observed with the lowest concentration (1 μg/ml) of tribendimidine: 66.6% of *S. ratti* were still active after an incubation period of 72 h. Ninety-six h post-incubation 50.4% of *S. ratti* showed no movement when exposed to this concentration, similar to the control group, where 60.0% of the worms were found to be inactive.

Effect of tribendimidine on adult *S. ratti*: dose-response relationship

Figure 1 shows the effect of tribendimidine on *S. ratti* rhabditiform larvae harvested from rat fecal samples as assessed by quantitative stool examination. In the first set of experiment (evaluating single 50, 100 and 200 mg/kg oral doses of tribendimidine) on 5 daily stool examinations, between 3300 (6 days post-exposure) and 10,000 (9 days post-exposure) rhabditiform larvae per gram of stool were estimated in the group of untreated and served as controls.

Statistical analyses were done with version 2.4.5 of Statsdirect statistical software (Statsdirect Ltd; Cheshire, UK). The effect of tribendimidine was assessed by comparing the mean number of *S. ratti* rhabditiform larvae in the stool and *S. ratti* adults and rhabditiform larvae in the intestine in the treatment group with the mean number of larvae and adults in the respective control group. The responses between the medians of the treatment and control groups regarding larvae and stool and larvae and adults in the intestines were analysed with the Kruskal-Wallis (KW) test. Differences in medians were considered to be significant at a significance level of 0.05.
untreated control animals. No larvae were found in fecal samples obtained from rats treated with single 100 or 200 mg/kg oral doses of tribendimidine and 0.5 mg/kg ivermectin commencing 48 h post-treatment. While no larvae were found at 48 h and 96 h post-treatment with 50 mg/kg tribendimidine, a low mean of 100 larvae per gram of stool was estimated 72 h post-treatment (Figure 1). Untreated rats in the second set of experiment (assessing the activity of 12.5 and 25 mg/kg tribendimidine) passed between 1000 and 18,600 *S. ratti* rhabditiform larvae per gram of stool on days 5 to 9 post-exposure. Larvae were also present in the stools of rats treated with 25 and 12.5 mg/kg tribendimidine; the highest numbers of rhabditiform larvae were detected 72 h post-treatment, namely 1300 and 2900 larvae per gram of stool in rats treated with 25 and 12.5 mg/kg, respectively (Figure 2). However, treatment had a significant effect on larvae presence in stools (KW = 67.1, degree of freedom (df) = 2; \(P < 0.001\)).

The effect of tribendimidine against *S. ratti* rhabditiform larvae and adults in the intestine, as assessed by worm burden reduction, is summarized in Table 2. The untreated control rats harbored a mean of 1012 *S. ratti* rhabditiform larvae and 413 adult worms in their intestines. Tribendimidine given at doses of 50, 100 and 200 mg/kg resulted in complete elimination of larvae and adult worms.

For comparison, ivermectin administered at 0.5 mg/kg achieved a significant larval reduction (90.0%; KW = 4.58, \(P = 0.032\)) and a complete elimination of adult worms.

### Table 1. Observed immobility of *S. ratti* L3 after exposure in vitro to tribendimidine at three different concentrations.

| Drug                  | Drug concentration (\(\mu\)g/ml) | % of worms inactive (SD) after incubation for |
|-----------------------|---------------------------------|---------------------------------------------|
|                       |                                 | 5 min       | 1 h         | 2 h         | 24 h        | 48 h        | 72 h        | 96 h        |
| Control               | No treatment                    | 40.0 (28.5) | 55.0 (7.1)  | 36.7 (20.1) | 23.4 (23.2) | 26.7 (16.5) | 50.0 (28.6) | 60.0 (14.1) |
| Tribendimidine        | 1                               | 30.0 (26.7) | 0 (0)       | 33.4 (2.4)  | 23.4 (6.3)  | 5.0 (7.1)   | 33.4 (9.4)  | 58.4 (17.0) |
|                       | 10                              | 30.0 (28.6) | 48.3 (6.2)  | 70.0 (32.4) | 100 (0)     | -            | -            | -            |
|                       | 100                             | 36.6 (35.6) | 98.4 (2.4)  | 100 (0)     | -            | -            | -            | -            |

SD: standard deviation.

doi:10.1371/journal.pntd.0000136.t001
The second control group harbored 750 *S. ratti* adults and 2025 larvae in their intestines. Tribendimidine at 25 mg/kg produced a 91.4% reduction of *S. ratti* larvae and a complete reduction of adult worms. When tribendimidine was given at 12.5 mg/kg, the adult worm burden was reduced by 83.3% and the larval burden by 54.4%. There was a significant difference in the larval (KW = 6.83, df = 2, *P* = 0.041) and the adult worm (KW = 9.46, df = 2, *P* = 0.009) burden between these 2 treatments and the control groups.

**Effect of tribendimidine on juvenile *S. ratti***

Figure 3 shows the effect of tribendimidine given 1–3 days post-exposure on *S. ratti* rhabditiform larvae present in stool. The number of rhabditiform larvae in the control group increased from 700 on day 5 post-exposure to 6700 rhabditiform larvae per gram of stool 4 days later. A larval reduction ranging from 30.6% (6 days post-exposure) to 73.1% (9 days post-exposure) was observed in fecal samples of rats treated with 50 mg/kg tribendimidine on day 5 post-exposure.

**Table 2.** Effect of ivermectin and tribendimidine (different doses) against adult *S. ratti* harbored in rats.

| Treatment    | Dose (mg/kg) | No. of rats cured* investigated | Mean larval burden (SD) | Mean adult worm burden (SD) | Total larval burden reduction (%) | KW   | *P*-value | Total adult worm burden reduction (%) | KW   | *P*-value |
|--------------|--------------|--------------------------------|-------------------------|---------------------------|----------------------------------|------|-----------|-----------------------------------|------|-----------|
| Control 1a   | No treatment | 0/4                            | 1012 (640)              | 413 (284)                 | -                                | -    | -         | -                                 | -    | -         |
| Control 2b   | No treatment | 0/4                            | 2025 (1001)             | 750 (387)                 | -                                | -    | -         | -                                 | -    | -         |
| Tribendimidine 12.5b | 0/4            | 925 (846)                     | 125 (96)                | 54.3                      | 6.83                             | 0.041| 83.3      | 9.46                              | 0.008|           |
| 25b          | 0/4           | 175 (150)                     | 0                       | 91.4                      | 100                              | 0    | 100       | 100                               | 10.02| 0.016     |
| 50b          | 4/4           | 0                              | 0                       | 100                      | 14.61                            | 0.002| 100       | 10.28                             | 0.016|           |
| 100b         | 4/4           | 0                              | 0                       | 100                      | 100                              | 0    | 100       | 100                               | 0    |           |
| 200b         | 4/4           | 0                              | 0                       | 100                      | 100                              | 0    | 100       | 100                               | 0    |           |
| Ivermectin  2.5a | 4/4           | 150 (150)                     | 0                       | 90                       | 4.58                             | 0.032| 100       | 100                               | 4.0  | 0.045     |

*number of rats without *S. ratti*.
*a first experiment.
*b second experiment.

SD: standard deviation.

KW: Kruskal-Wallis.

doi:10.1371/journal.pntd.0000136.t002
1 post-exposure. In rats treated with tribendimidine at 50 mg/kg on day 2 post-exposure, a reduction of rhabditiform larvae ranging between 50.0% (7 days post-exposure) and 83.6% (9 days post-exposure) was observed. Finally, no rhabditiform larvae were found in 5 consecutive stool samples from rats treated with 50 mg/kg of tribendimidine on day 3 post-exposure. Administration of tribendimidine to rats harboring tissue stages of *S. ratti* had a significant effect on the presence of larvae in stool (KW = 14.1; df = 3, P = 0.002).

Table 3 summarizes observed larvae and adult worm burden reductions in the intestines of rats following treatment with tribendimidine. Administration of a single 50 mg/kg oral dose of tribendimidine on day 1 post-exposure to *S. ratti*-infected rats showed no effect on the intestinal larval and adult parasite load. Treatment of infected rats 48 h post-exposure with a single 50 mg/kg oral dose of tribendimidine resulted in larvae and adult worm burden reductions of 41.0–61.5%. Finally, when tribendimidine (50 mg/kg) was given 72 h post-exposure, a 98.9% reduction of larvae in the intestines was observed. There was a significant difference between the number of larvae (KW = 9.65, P = 0.021) and adult worms (KW = 6.29, P = 0.098) recovered from the intestines of treated and non-treated control rats.

**Table 3.** Effect of a single 50 mg/kg oral dose of tribendimidine against immature stages of *S. ratti* harbored in rats.

| Treatment | Treatment day post-exposure (stage) | No. of rats cured*/investigated | Mean larval burden (SD) | Mean adult worm burden (SD) | Total larval burden reduction (%) | Total adult worm burden reduction (%) |
|-----------|-----------------------------------|-------------------------------|------------------------|-----------------------------|---------------------------------|--------------------------------------|
| Control   | No treatment                      | 0/4                           | 2375 (1190)            | 150 (58)                    | -                               | -                                    |
| Tribendimidine | 1 (L3 larvae in lungs or in cranial or nasal cavities)* | 0/4                           | 2800 (2546)            | 225 (263)                   | 0                               | 0                                    |
|           | 2 (some preadolescent stages in intestine)* | 0/4                           | 1400 (716)             | 50 (58)                     | 41.1                            | 61.5                                 |
|           | 3 (some preadolescent stages in intestine)* | 0/4                           | 25 (50)                | 50 (58)                     | 98.9                            | 61.5                                 |

*number of rats without *S. ratti*.
*Harder et al. (2001) [25].
Kruskal-Wallis (KW) testing difference in total larval burden reduction (KW = 9.66, df = 3, P = 0.021) and adult worm burden reduction (KW = 6.29, df = 3, P = 0.098).
SD: standard deviation.
doi:10.1371/journal.pntd.0000136.t003
Our in vitro studies revealed that worms incubated in the presence of 100 μg/ml tribendimidine died within 2 h. The worms had a coiled appearance. It has been suggested that tribendimidine, which, similar to amidantel is biotransformed into p-(1-dimethylamino ethylimino) aniline (S.H. Xiao, pers. comm.), acts as agonist at the level of the acetylcholine receptor [21]. The rapid onset of action of tribendimidine is supported by our in vivo studies since no larvae were found in stools of rats treated with at least 50 mg/kg of tribendimidine 48 h post-treatment. A previous investigation showed that tribendimidine acted similarly rapidly when it was administered to mice infected with the intestinal trematode *E. caproni*. Scanning electron microscopic investigations revealed that severe damage of the tegument already occurred 2 h after drug administration and 8 h post-treatment the majority of worms had been expelled [12].

Concluding, we have documented in vitro and in vivo activities of tribendimidine against *S. ratti*. Our findings warrant further investigations, which is justified as follows. First, discovery and development of novel anthelmintic drugs in general [22,23] and strongylolidal drugs in particular, is limited. Hence, over the past decade only a few compounds have been examined in the *S. ratti*-rat model [24,25]. Second, tribendimidine has recently been registered in China as an anthelmintic drug, and it might thus be deployed as an additional control tool against major helminth infections [26]. Efforts are ongoing to pursue registration of tribendimidine in a 1st tiered regulatory agency so that the drug could eventually be integrated in global helminth control programs. Since many helminth infections show large geographical overlaps, it will be important to monitor the effect of tribendimidine on concomitant infections of different nematodes and trematodes. We are currently in the process of examining the effect of tribendimidine against *S. stercoralis* in people co-infected with this parasitic roundworm and other nematodes and trematodes.

**Acknowledgments**

We thank Professor Xiao Shuhua for providing a sample of tribendimidine to carry out the in vitro and in vivo studies reported here. We thank Professor Thomas A. Smith for expert help with the statistical analysis.

**Author Contributions**

Conceived and designed the experiments: JU JK. Performed the experiments: JU JK. Analyzed the data: JK. Contributed reagents/materials/analysis tools: YE. Wrote the paper: JU JK.

**References**

1. Bethony J, Brooker S, Albonico M, Geiger SM, Loukas A, et al. (2006) Soil-transmitted helminth infections: ascariasis, trichuriasis, and hookworm. Lancet 367: 1521–1532.
2. Steinmann P, Zhou XN, Du ZW, Jiang JY, Wang LB, et al. (2007) Occurrence of *Strongyloides stercoralis* in Yunnan province, China, and comparison of diagnostic methods. PLoS Negl Trop Dis 1: e75. doi:10.1371/journal.pntd.0000075.
3. Boulderware DR, Stauffer WM, Hendel-Paterson BR, Rocha JL, Sert RC, et al. (2007) Maltreatment of *Strongyloides* infection: case series and worldwide physicians-in-training survey. Am J Med 120: e1–8.
4. Keiser PB, Numan TB (2004) *Strongyloides stercoralis* in the immunocompromised population. Clin Microbiol Rev 17: 208–217.
5. Hirata T, Kishimoto K, Kinjo N, Hikama A, Kinjo F, et al. (2007) Association between *Strongyloides* stercoralis infection and biliary tract cancer. Parasitol Res 101: 1345–1348.
6. Utzinger J, Keiser J (2004) Schistosomiasis and soil-transmitted helminthiasis: common drugs for treatment and control. Expert Opin Pharmacother 5: 263–285.
7. Shikata K, Zaha O, Niimura S, Ikema M, Nakamura H, et al. (1992) Long term eradication rate of mebendazole therapy for strongyloidiasis. Kansenshogaku Zasshi 66: 354–359.
8. Satoh M, Kokazae A (2006) Treatment strategies in controlling strongyloidiasis. Expert Opin Pharmacother 5: 2293–2301.
9. Fox LM (2006) Ivermectin: uses and impact 20 years on. Curr Opin Infect Dis 19: 580–583.
10. Xiao SH, Wu HM, Tanner M, Utzinger J, Wang C (2005) Tribendimidine: a promising, safe and broad-spectrum anthelmintic agent from China. Acta Trop 94: 1–14.
11. Xiao SH, Wu ZX, Zhang J, Wang C (2007) Clinical observation on 899 children infected with intestinal nematodes and treated with tribendimidine enteric coated tablets: a phase IV clinical trial. Chin J Parasitol Para Dis (in press).
12. Keiser J, Xiao SH, Utzinger J (2006) Effect of tribendimidine on adult *Echinostoma caproni* harbored in mice, including scanning electron microscopic observations. J Parasitol 92: 850–8516.
13. Keiser J, Xiao SH, Choller J, Tanner M, Utzinger J (2007) Evaluation of the in vivo activity of tribendimidine against *Schistosoma mansoni*, *Fasciola hepatica*, *Clonorchis sinensis* and *Opisthorchis viverrini*. Antimicrob Agents Chemoter 51: 1096–1098.
14. Garcia LS (2001) Diagnostic medical parasitology. Washington D.C.: ASM Press. pp 791.
15. Ren HN, Cheng BZ, Zhuang ZN (1987) Experimental therapeutic efficacy of a new anti-hookworm drug, tribendimidine. Chin J Parasitol Para Dis 3: 262–264.
16. Tobata-Kudo H, Kudo H, Tada I (2005) *Strongyloides ratti*: chemokinesis of glycolytic enzyme- and lectin-treated third-stage infective larvae in vitro. Parasitol Int 54: 147–152.
17. Rajasekariah GR, Deb BN, Dhage KR, Bose S (1986) Response of laboratory-adapted human hookworm and other nematodes to ivermectin. Ann Trop Med Parasitol 80: 615–621.
18. Grove DI (1983) The effects of 22,23-dihydroavermectin B1 on *Strongyloides ratti* and *S. stercoralis* infections in mice. Ann Trop Med Parasitol 77: 405–410.
19. Mojon M, Saura C, Rousse N, Tran Manh Sung R (1987) Albendazole and thiabendazole in murine strongyloidiasis. J Antimicrob Chemother 19: 79–85.
20. Grove DI (1982) *Strongyloides ratti* and *S. stercoralis*: the effects of thiabendazole, mebendazole, and cambendazole in infected mice. Am J Trop Med Hyg 31: 469–476.
21. Tomlinson G, Albuquerque CA, Woods RA (1985) The effects of amidantel (BAY d 8815) and its deacylated derivative (BAY d 9216) on *Caenorhabditis elegans*. Eur J Pharmacol 113: 253–262.

22. Horton J (2003) Human gastrointestinal helminth infections: are they now neglected diseases? Trends Parasitol 19: 527–531.
23. Keiser J, Utzinger J (2007) Advances in the discovery and development of trematocidal drugs. Expert Opin Drug Discov (in press).
24. von Samson-Himmelstjerna G, Harder A, Schnieder T, Kalbe J, Mencke N (2000) In vivo activities of the new anthelmintic depsipeptide PF 1022A. Parasitol Res 86: 194–199.
25. Harder A, von Samson-Himmelstjerna G (2001) Activity of the cyclic depsipeptide emodepside (BAY 44-4400) against larval and adult stages of nematodes in rodents and the influence on worm survival. Parasitol Res 87: 924–928.
26. Hotez PJ, Bethony J, Bottazzi ME, Brooker S, Diemert D, et al. (2006) New technologies for the control of human hookworm infection. Trends Parasitol 22: 327–331.