Mitochondrial Ribosome as the Target for the Macrolide Antibiotic Clarithromycin in the Helminth *Echinococcus multilocularis*

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The mitochondrial rRNA of the tapeworm species *Echinococcus multilocularis* carries an adenine at sequence position 2058 (numbering according to that for *Escherichia coli*) of the large-subunit rRNA (lsrRNA), while the nucleus-encoded rRNA, as determined in this study, is characterized by 2058G. This indicates a dichotomy in the drug susceptibilities of ribosomes: cytoplasmic ribosomes are predicted to be resistant to macrolide antibiotics, while mitochondrial ribosomes lack the most common chromosomal resistance determinant, lsrRNA 2058G. Upon incubation with the macrolide clarithromycin, the formation of vesicles from metacestode tissue was reduced in a dose-dependent manner. Electron microscopy revealed distinct morphological alterations both of the mitochondria and of the vesicle wall (e.g., loss of microtriches) in drug-treated vesicles. Adult worms lost their motility and displayed morphological changes (shortening and constriction of proglottids and the presence of vacuoles) upon incubation with clarithromycin. Our findings demonstrate that macrolides have distinct in vitro effects on *E. multilocularis*, endorsing the use of sequence-based in silico approaches for exploitation of available ribosomal drugs as anthelmintic agents.

*Echinococcosis* is a severe human disease caused by the larval (metacestode) stages of tapeworms belonging to the genus *Echinococcus*. Depending on the parasite species involved, different forms of echinococcosis are found, namely, cystic echinococcosis (causative agent *Echinococcus granulosus*), alveolar echinococcosis (causative agent *Echinococcus multilocularis*), and polycystic echinococcosis (causative agents *Echinococcus vogeli* and *Echinococcus oligarthrus*). Metacestodes develop in a tumor-like manner in different sites of the human body from oncospheres liberated from accidentally ingested worm eggs, which are shed in the feces of various species of carnivores that can act as definitive hosts for the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the species of carnivores that can act as definitive hosts for the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms. Cystic echinococcosis and alveolar echinococcosis are of special medical importance, with the adult, intestinal tapeworms.

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The effects of the drugs in vitro were assessed by counting the number of vesicles produced from metacestode tissue after 14 days (14) and by assessing the viability and morphology of adult worms after 2 days of drugs exposure.

Metacestode tissue was cut into small blocks, with the lengths of the edges being approximately 3 mm, and the blocks were washed twice in complete medium (minimum essential medium [MEM; Gibco, Basle, Switzerland] supplemented with 10% fetal calf serum, 2 mM glutamine, 100 U penicillin/ml, 100 μg streptomycin/ml, 0.25 μg amphotericin B/ml). Two blocks per well were placed in 12-well plates (Techno Plastic Products, Trasadingen, Switzerland) with 3 ml of culture medium per well. The culture medium consisted of a 1:1 mixture of MEM and fresh supernatants of human fibroblast cell (MRC-5) cultures growing in the same medium in T flasks. The plates were kept in an incubator at 37°C with 5% CO2, and 2 ml of medium per well was exchanged every 3 to 4 days. Four wells per treatment (control or drugs) were used, and these experiments were done three times.

Adult worms (approximately 20 per well) were seeded into 12-well plates and incubated as described above in 3 ml of the complete MEM supplemented with 12 mg/ml glucose. The effects of the drugs were assessed in triplicate assays in a single experiment after 2 days.

**Drugs.** Clarithromycin (Klacid) was from Abbott (Baar, Switzerland); hygromycin B, mebendazole, and praziquantel were from Fluka (Buchs, Switzerland). Stock solutions (10 mg/ml) of mebendazole were prepared in dimethyl sulfoxide; those of praziquantel were prepared in ethanol.

**lsrRNA sequences.** The peptidyltransferase region of the cytoplasmic lsrRNA was amplified by PCR with primers 22 and 27, which target conserved flanking regions (36), with *E. multilocularis* DNA isolated from metacestode material. The sequences of both strands were determined by direct sequencing with an ABI PRISM 310 sequencer (Applied Biosystems, Baar, Switzerland). The corresponding sequence of the mitochondrial lsrRNA was available (Fukunaga, unpublished data; GenBank accession no. AB018440).

**Electron microscope.** Cysts that had developed in culture for 14 days were fixed with 2.5% glutaraldehyde in 100 mM Na/K-phosphate, pH 7.4, at 4°C for 1 h, washed in Na/K-phosphate, postfixed with 2% osmium tetroxide in Na/K-phosphate at 4°C for 1 h, and embedded in Epon. Sections of 1 μm were stained with toluidine blue for light microscopy; sections of 50 to 60 nm from areas of interest were stained with uranyl acetate and lead citrate and examined in a Cm 12 electron microscope (Philips, Eindhoven, The Netherlands) equipped with a slow-scan charge-coupled-device camera (Gatan, Pleasanton, CA).

**Nucleotide sequence accession number.** The part of the lsrRNA gene of the cytoplasmic ribosome containing the peptidyltransferase region obtained in this study has been submitted to GenBank and can be found under accession no. AY615426.

**RESULTS**

**Sequence analysis of large-subunit rRNA.** Part of the lsrRNA gene of the cytoplasmic ribosome containing the peptidyltransferase region was amplified by PCR and sequenced (604 bp; GenBank accession no. AY615426), and the sequence was compared to the corresponding published sequence of the mitochondrial gene (GenBank accession no. AB018440). The presence of a guanine at lsrRNA position 2058 (numbering according to that for *Escherichia coli*) of the nucleus-encoded rRNA is expected to provide resistance to macrolides, while mitochondrial ribosomes carry an adenine at this position, which is predictive of susceptibility.

**In vitro investigations.** To study the effect of clarithromycin on *E. multilocularis* in vitro, metacestode tissue was incubated with the antibiotic; and vesicle formation was recorded and compared with that in the presence of two other drugs, hygromycin B and mebendazole, used as controls. Hygromycin B is a protein synthesis inhibitor with universal activity against archae, prokaryotes, and eukaryotes and is expected to inhibit both the cytoplasmic and the mitochondrial ribosomes (24). Mebendazole, a benzimidazole carbamate derivative, is an established anthelminthic compound that interferes with the microtubulus system of parasitic cells.

Upon exposure to clarithromycin, the vesicle formation of *E. multilocularis* was reduced in a dose-dependent manner (Fig. 1).
1). Whereas 41 vesicles were produced in the absence of the drug, 6 (14.6% of control) were recorded in the presence of 100 μg/ml clarithromycin and 21 (58.5%) were recorded in the presence of 10 μg/ml clarithromycin. In cultures containing hygromycin (100 μg/ml) or mebendazole (10 μg/ml), an average of one (2.8%) vesicle was produced.

The effect of clarithromycin on adult worms was compared with those of hygromycin B and praziquantel. Praziquantel is an established anthelmintic compound that causes severe spasms and paralysis of the worm’s muscles. Compared to the untreated adult worms (Fig. 2A), which had a high motility, worms incubated for 2 days in 10 μg/ml clarithromycin had a...
similar motility but differed morphologically by being slightly swollen. A dramatic effect was observed in worms incubated in 100 μg/ml clarithromycin: their motility was highly reduced, they appeared shortened and darker, the proglottids were deeply constricted, and vacuoles were present (Fig. 2B). Hydroxyurea and praziquantel had similar, although more pronounced, effects on morphology compared to that of 100 μg/ml clarithromycin; and the worms were motionless.

**Electron microscopy.** Electron microscopy was used to study alterations at the subcellular level. These studies revealed distinct morphological and ultrastructural alterations of metacestode tissue from in vitro cultures after cocultivation with clarithromycin (Fig. 3). The germinal layer of control vesicles consisted of several layers of cells containing mitochondria with distinct cristae and of a distinct tegument and numerous microtriches protruding into the acellular laminated layer (Fig. 3A and B). After incubation with clarithromycin (100 μg/ml) for 14 days, the vesicles were smaller and had laminated layers of various thicknesses (Fig. 3C). The cells lining the laminated layers were thin and often detached. They contained condensed mitochondria and mitochondria with indistinct cristae (Fig. 3C and D). Microtriches and the tegument were largely lacking (Fig. 3D).

**DISCUSSION**

In this paper, we demonstrate for the first time that an antibacterial agent, the macrolide clarithromycin, can have a direct effect on a helminth organism, *E. multilocularis*. Antibacterial drugs have previously been shown to have promise for the treatment of infections with filarial nematodes, but in an indirect manner, by acting on endosymbiotic bacteria (15). In addition, the aminoglycoside antibiotic paromomycin was used to treat human intestinal infections with taeniid tape-worms (5, 30), but the mode of action against *Taenia* is not known.

As predicted by in silico analysis, the target of clarithromycin in *E. multilocularis* is the mitochondrial translational machinery. Indeed, we observed dramatic changes in the ultrastructure of mitochondria, where the loss of cristae and condensation (Fig. 3D) become apparent upon cocultivation with this drug. However, morphological alterations of mitochondria have also been observed upon incubation with other drugs. Albendazole, a benzimidazole component with activity on tubulin formation, resulted in rounded mitochondria, which were abnormally increased in size and which contained altered cristae (16). The presence of aberrant mitochondria was recently reported as an effect of nitazoxanide against *E. multilocularis* metacestode tissue in vitro, but no detailed information was provided (32). Hence, the mitochondrial alterations that we observed do not prove that clarithromycin directly affects the mitochondrial ribosome. However, in a comparable study with the amoebal species *Acanthamoeba castellanii*, where genetic analyses predicted the resistance of the cytoplasmic ribosome and the susceptibility of the mitochondrial ribosome to macrolides, the fortuitous isolation of a macrolide-resistant mutant and its genetic characterization showed that the expected point mutation 2058A→G of the mitochondrial lsrRNA was associated with drug resistance (no encystment of the amoebae) and, hence, with a lack of ultrastructural changes to the mitochondria upon cocultivation with clarithromycin (21).

The other structural changes observed in the *E. multilocularis* vesicles upon incubation with clarithromycin (e.g., loss of microtriches and separation of the germinal and the laminated layers) seem to be rather nonspecific effects, as they have also been observed in experiments with benzimidazoles or nitazoxanide (16, 32).

Rather surprising findings were the rapid (within 2 days) and distinct effects of clarithromycin (100 μg/ml) on adult worms (highly reduced motility, swelling, vacuolization, constriction of proglottids) which were comparable to the effects of antibiotic or anthelmintic compounds with unrelated modes of action (hydroxyurea B, praziquantel). The disruption of energy production in the mitochondria by clarithromycin appears to be a fast event, and the alternative, low-efficiency cytosolic respiratory metabolism in tapeworms (23) does not seem to have the capacity to compensate. Again, the similar changes induced by the different drugs may simply reflect a state of stress in the parasite that stems from the disruption of different cell organelles.

Clarithromycin showed potent anthelmintic activity in vitro, but whether such an effect will extend to the in vivo situation remains to be investigated. Of crucial importance is whether the drug would penetrate into larger metacestode tissues. In our in vitro experiments, by using metacestode material cut into small blocks, this possible barrier was not of concern. Furthermore, we would expect that macrolides act only in a parasitostatic manner and not in a parasitocidal manner, according to the results of experiments done with *A. castellani* (21), which revealed that the drug-induced effects were nearly perfectly reversible. Hence, macrolides might be promising agents for the treatment of alveolar echinococcosis, particularly because of their synergistic actions with other existing drugs (benzimidazoles, nitazoxanide).

The experiments described in this study have considerable relevance for the search for drugs endowed with anthelmintic activities. Our observations endorse the use of sequence-based in silico approaches for the possible exploitation of the available ribosomal drugs as anthelmintic agents. Ultimately, elucidation of the mechanisms governing the selectivities and the specificities of ribosomal drugs should provide ample opportunities for the development of anthelmintic agents by rational drug design.

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**REFERENCES**
