ABSTRACT

Background: Continuous attempts are being made to develop new and more effective drugs for treatment of schistosomiasis. *Curcuma longa* is a medicinal plant that contains yellow pigments known as curcuminoids (curcumin, bidesmetoxi-curcumin and monodesmetoxi-curcumin), with curcumin as the highest component. The main biological effect depends on curcumin content. Curcumin has low bioavailability and poor-water solubility. Preparation of *C. longa* (CI) nanoemulsion was attempted to enhance solubility and bioavailability of curcumin.

Objective: The production of CI nanoemulsion, and study of its pharmacokinetic properties (namely solubility and intestinal bioavailability) and anti-helminthic effects on *Schistosoma mansoni* cercariae, schistosomules and adults.

Methodology: CI nanoemulsion was prepared from ethanol extract of crude powder of CI rhizome. Characterization was done by transmission electron microscope (TEM), testing solubility and intestinal bioavailability. Schistosomicidal effects were performed by *in vitro* assay.

Results: CI nanoemulsion showed increased solubility and bioavailability compared to ethanol extract of the tested plant; as well as time and dose dependent schistosomicidal effects on cercariae and 24h-old schistosomules of *S. mansoni*. In addition, it exhibited an optimal activity against the adult stage with decreased motor activity of the worms.

Conclusion: This study proves that CI nanoemulsion has enhanced solubility and bioavailability, as well as schistosomicidal activity.

Key Words: *Curcuma longa*, Nanoparticles, *Schistosoma mansoni*.

INTRODUCTION

Human schistosomiasis is a chronic debilitating disease caused by parasites of the genus *Schistosoma*. *Schistosoma* infections ranks just after malaria in terms of parasite-induced human morbidity and mortality. It is estimated that 779 million people are at risk of schistosomiasis. The number of active *Schistosoma* infections was estimated to be between 391 and 587 million people worldwide. Praziquantel and oxamniquine are the drugs available for the treatment of schistosomiasis. However, development of resistant strains for both drugs has been reported, leading to schistosomiasis treatment failure that highlighted the importance of developing new and more effective drugs for this disease. In this context, the past years have been marked by increasing search for anti-parasitic drugs from natural sources. Plants are the major source of biologically active compounds for the development of new treatments. In 2013, Abaza reviewed all herbs that were used in treatment of schistosomiasis including Chinese medicine, Carvacrol (essential oil of *Origanum vulgare*) obtained from pepperwort, Myrrh (oleo-gum resin from *Commiphora molmol*), artemisinin derivatives isolated from *Artemisia annua*, curcumin (*C. longa*), quinine and quinidine (*Cinchona officinalis*), garlic extract (*Allium sativum*), black seeds (*Nigella sativa*) and other several native plants from Brazil.

One of these plants is the rhizome of *Curcuma longa*, or turmeric (*Zingiberaceae* family) which is highly regarded as a universal remedy in herbal medicine, with a wide spectrum of pharmacological activities. Turmeric contains 69.4% carbohydrates, 6.3% protein, 5.1% fat, 3.5% minerals, and 13.1% moisture. The essential oil (5.8%) obtained by steam distillation possesses sesquiterpenes (53%), zingiberene (25%), a-phellandrene (1%), sabinene (0.6%), cineol (1%), and borneol (0.5%). Curcumin (3–4%) is responsible for the yellow color, and comprises curcumin I (94%), curcumin II (6%) and curcumin III (0.3%). Its benefits have been reported against gastrointestinal diseases, hepatic disorder, inflammation, rheumatism, sinusitis, anorexia, cough, anti-cancer, anti-diabetic, anti-oxidant, hypolipidemic, anti-microbial, anti-fertility, anti-venom. In addition, curcumin possesses
anti-parasitic activity against T. brucei[17], P. berghei[18], G. lamblia[19], L. donovani[20], and S. mansoni[21].

However, curcumin is a low water soluble compound, of low bioavailability, which means that its beneficial active compound may not be fully absorbed by the body when it is digested[22]. Therefore, improvement of curcumin properties may be beneficial in enhancing its effects. One of the methods is by reducing particle size to a nano size. Nanoemulsions can be prepared by solubilizing the lipophilic bioactive components as the oil phase, and then homogenizing this phase with an aqueous phase containing a water-soluble emulsifier. The small size of droplets in nanoemulsion have different physicochemical and biological properties than in a standard emulsion. Aggregation and separation of particles are better attained when in nanoemulsion form due to the small droplet sizes[23,24]. Nanoparticles from curcumin can be prepared by different methods. One of them is a homogenization process, in which a heterogeneous solution classified as nanoemulsion is formed[25]. The preparation of Cl nanoemulsion in our study was attempted to enhance its solubility as well as stability especially for medical application. The droplet size, solubility and penetration of Cl nanoemulsion were observed and compared with that of an extract curcumin emulsion.

Based on the promising effects of curcumin, the purpose of this work is to evaluate the in vitro effect of Cl nanoemulsion on S. mansoni cercariae, schistosomules and adults.

MATERIALS AND METHODS

Type of study: Prospective comparative experimental study.

Preparation of Cl nanoemulsion: Cl nanoemulsion was prepared from ethanol extract of crude powder of C. longa rhizome purchased from the local market. The latter was prepared by mixing 200 g crude powder of rhizome of tested plant in 800 ml of 98% ethanol using high speed mixture. Liquid phase was evaporated at room temperature in petri dishes. The resultant powder was preserved in a sterile glass bottle at room temperature till used for preparing the nanoemulsion. Oil water Cl nanoemulsion was prepared by mixing oil phase of ethanol extract in aqueous phase containing 10% (v/v) Tween 80 (based on the oil phase) and maltodextrin (4% and 10%) dissolved in phosphate buffer solution (PBS) at pH 7 and stirring using high speed (20000 rpm) magnetic stirrer for at least 15 min to ensure complete hydration[26]. The final product was a stable emulsion containing 20% ethanol extract of curcumin.

Cl nanoemulsion characterization: Extract particles sizes were measured by using transmission electron microscope (JEM-1200EX, JEOL, Tokyo, Japan), at electron microscope unit, Faculty of Science, Ain Shams University. A drop of the nano-curcumin suspension (10 µL; 2 mg/ml) was placed on the 3 mm copper grid and allowed to dry. Solubility of both ethanol extract and nanoemulsion of tested plant were tested against different solvents namely acetone, ethanol, methanol, and cold distilled water. Cl nanoemulsion particles were mixed with the solvents in the ratio 1:1. The solution was well mixed and allowed to stand for 6 hours before observation to ensure complete dissolving. GIT bioavailability of different preparations was tested by measuring the extent of their penetration through a closed isolated loop of rabbit's jejunum 10 cm in length perfused in physiological Tyrod solution at 37°C in a 10 ml organ bath for 3 hours. Concentration of absorbed curcumin was measured at maximal absorption of curcumin (530 nm) using a spectrophotometer[27].

Assessment of drug effect on S. mansoni stages

Cercaricidal effect of Cl nanoemulsion: S. mansoni (John Bruce Egyptian strain) cercariae were obtained from infected Biomphalaria alexandrina snails obtained from the Schistosome Biological Supply Center (SBSC), Theodore Bilharz Research Institute (TBRI, Imbaba, Giza, Egypt). Cercariae were collected by exposing snails to light for 2 h. The cercariae (from 30 to 40) were incubated in 10 ml dechlorinated tap water. Cl nanoemulsion was added in final concentrations of 12.5, 25, 50 and 100 µg/ml, in petri dishes. Untreated cercariae were examined as a control. Cercariae were observed after 5, 30, 60, 90 and 120 min under a dissecting microscope (Olympus, Tokyo, Japan). The cercariae were considered dead when no movement was observed for at least 1 min of examination. All the experiments were repeated twice.

Schistosomulicidal effect of Cl nanoemulsion: Schistosomes were prepared in vitro according to the method of Colley and Wikel[28]. Cercariae were subjected to shearing forces created by 10-14 passages through a 22-gauge hypodermic needle. The organisms obtained were concentrated simply by gravity sedimentation on ice. The sediment was re-suspended in RPMI-1640 medium containing 10% heat-inactivated fetal calf serum, 100 IU/ ml penicillin and 100 µg/ml streptomycin[29]. About 20-40 schistosomes were added to each well of sterile tissue culture plates with flat-bottomed wells and incubated overnight in 5% CO2, 100% humidity, at 37°C. After 24 h of incubation, Cl nanoemulsion was added to the culture medium to give final concentrations of 12.5, 25, 50, and 100 µg/ml. All experiments were performed in duplicates and examined for viability at 24 and 48 h post treatment. Some wells were maintained without addition of Cl nanoemulsion as control for non-specific death of schistosomes. The percentage of damaged schistosomes was measured by addition of trypan blue at a final dilution of 0.2%. Dark blue schistosomes that permitted penetration of the dye were considered damaged[30].

45
Schistosomicidal effect of Cl nanoemulsion: Swiss albino mice CD-1, weighing 18–22 g each, were obtained from SBSC, kept under environmentally-controlled conditions (temperature 25°C; humidity 70%; 12-h light and 12-h dark cycle) and acclimatized for one week before infection. The maintenance and care during experimentation of animals was compliant with international guidelines for the human use of laboratory animals. Adult worms were removed from portal and mesenteric veins of infected mice after 90 days[29], sexed and counted[30]. Six to ten mature worms including both sexes were cultured per well in 24-well plates containing RPMI medium at 37°C and 5% CO₂, immediately after animal perfusion to ensure their vitality. Cl nanoemulsion was added to obtain final concentrations of 12.5, 25, 50, 100 µg/ml. Negative control wells contained adults incubated in culture media. All experiments were performed in duplicates and read at 24, 48 and 72 h post treatment[30]. The effect of Cl nanoemulsion concentrations on S. mansoni adult worms were monitored every 24 h during 96 h for evaluation of their motor activity. The mortality rate was given a score of 0-3 (3=normal motility, 2=slow motility, 1=sluggish motility, 0=no motility)[30]. The worms were considered dead when no movement was observed for at least 2 min of examination[31].

Statistical analysis: The collected data were analyzed using SPSS version 16 software, data were presented as number and percentage. Goodness of fit test (one way chi square) and Z tests were used. P<0.05 was considered significant.

RESULTS
Transmission electron microscopy (TEM) of Cl nanoemulsion particles showed that the average particles size was 24 nm; Cl extract demonstrated particles with an average size distribution of 214 nm (Figures 1, 2). Cl nanoemulsion proved to be highly soluble in water, methanol, ethanol and acetone. The Cl extract dissolved in methanol, ethanol and acetone, but not water. The in vitro penetration of Cl nanoemulsion across the membrane cells was 20%, and that of Cl extract was 13%. Cl nanoemulsion showed 100% death of cercariae after 30, 60, 75 min for drug concentrations of 100, 50 and 25 µg/ml, respectively. No cercaricidal effect was noted with drug concentration 12.5 µg/ml, after 120 min incubation.

In vitro schistosomicidal effect of Cl nanoemulsion on adult worms was statistically significant (P<0.001), according to concentration and incubation period. The effect on their motility was variable after 24 h of incubation with concentrations of 50, and 100 µg. After 48 h of incubation with the same concentration there were variable effects on motility and death of worms. After 96 h of incubation with 100 µg of the nanoemulsion all worms were dead (Table 1). There was no statistical significant difference (P> 0.05) in its effect on males and females (Figure 3). Incubation of 24 h old schistosomules with different concentrations of nanoemulsion resulted in 100% mortality after 24 h incubation at a concentration of 100 µg. Other concentrations had statistically significant lower effects (P<0.001) (Figure 4).

Fig. 1: C. longa extract particles by TEM: Average diameter distribution of extract nanoparticles, 214 nm.size of 214 nm.

Fig. 2: C. longa nanoemulsion particles by TEM: Average diameter of nanoemulsion particles 24 nm.
### Table 1: *In vitro* schistosomicidal effect of C. longa nanoemulsion on adult worms

<table>
<thead>
<tr>
<th>Tested group</th>
<th>Incubation period</th>
<th>No of worm</th>
<th>Adult schistosomes activity No. (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Normal Score=3</td>
</tr>
<tr>
<td>No treatment</td>
<td>24 h</td>
<td>9</td>
<td>9 (100%)</td>
</tr>
<tr>
<td></td>
<td>48 h</td>
<td>9</td>
<td>9 (100%)</td>
</tr>
<tr>
<td></td>
<td>72 h</td>
<td>9</td>
<td>9 (100%)</td>
</tr>
<tr>
<td></td>
<td>96 h</td>
<td>9</td>
<td>9 (100%)</td>
</tr>
<tr>
<td>100 µg</td>
<td>24 h</td>
<td>15</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>48 h</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>72 h</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>96 h</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>50 µg</td>
<td>24 h</td>
<td>13</td>
<td>10 (76.9%)</td>
</tr>
<tr>
<td></td>
<td>48 h</td>
<td>3 (23.1%)</td>
<td>7 (53.8%)</td>
</tr>
<tr>
<td></td>
<td>72 h</td>
<td>2 (15.3%)</td>
<td>5 (38.5%)</td>
</tr>
<tr>
<td></td>
<td>96 h</td>
<td>0 (0%)</td>
<td>5 (38.5%)</td>
</tr>
<tr>
<td>25 µg</td>
<td>24 h</td>
<td>13</td>
<td>13 (100%)</td>
</tr>
<tr>
<td></td>
<td>48 h</td>
<td>8 (61.5%)</td>
<td>2 (15.4%)</td>
</tr>
<tr>
<td></td>
<td>72 h</td>
<td>1 (7.7%)</td>
<td>8 (61.5%)</td>
</tr>
<tr>
<td></td>
<td>96 h</td>
<td>0 (0%)</td>
<td>5 (38.5%)</td>
</tr>
<tr>
<td>12.5µg</td>
<td>24 h</td>
<td>6</td>
<td>6 (100%)</td>
</tr>
<tr>
<td></td>
<td>48 h</td>
<td>6 (100%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td></td>
<td>72 h</td>
<td>0 (0%)</td>
<td>6 (100%)</td>
</tr>
<tr>
<td></td>
<td>96 h</td>
<td>0 (0%)</td>
<td>6 (100%)</td>
</tr>
<tr>
<td><strong>Statistical analysis</strong></td>
<td><strong>P value</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>24 h: X²= 46.8</td>
<td>0.012 (S)</td>
<td>0.08 (NS)</td>
</tr>
<tr>
<td></td>
<td>48 h: X²= 64.8</td>
<td>0.03 (S)</td>
<td>&lt;0.001 (S)</td>
</tr>
<tr>
<td></td>
<td>72 h: X²= 77.07</td>
<td>&lt;0.001 (S)</td>
<td>0.01 (S)</td>
</tr>
<tr>
<td></td>
<td>96 h: X²= 95.4</td>
<td>&lt;0.001 (S)</td>
<td>0.028 (S)</td>
</tr>
</tbody>
</table>

*P <0.001 (S)*

*S= Significant, NS= Non significant*
**DISCUSSION**

Curcumin is the major constituent in the rhizome of *C. longa* (Zingiberaceae). It is a naturally occurring compound that exhibits several biological activities\(^{[23]}\). Curcumin was active *in vitro* against schistosomes\(^{[21,33]}\), however, no study with curcumin nanoparticles has been tested against *Schistosoma* species. This study presents the first attempt of preparation and evaluation of nanoparticles from *C. longa* extract by homogenization method.

In this work, we extracted crude powder of *C. longa* rhizome using ethanol, which possesses predominate polar and minor non polar properties\(^{[34]}\). The end product of ethanol extraction of tested plant has a very low yield (20 g out of 1 kg of crude *C. longa* powder). The method employed in preparation of nanoemulsion improved the hydrophilic properties of the tested plant by transforming water insoluble crude extract into nanoemulsion\(^{[23]}\). Tween 80 which is a polyethoxylated sorbitin acted as emulsifier. It converted particles of tested plant into homogenously dispersed micelles. Tween 80 contains a lipophilic polysorbate group that faces the lipophilic core of curcumin micelle, and a polyoxyethylene hydrophilic group that faces hydrophilic solvents\(^{[30]}\). The role of multodextrin in improving solubility of Cl nanoemulsion is probably attributed to its hygroscopic properties. Magnetic stirring broke *C. longa* particles down into nanoparticles.

The present study revealed marked improvement of GIT bioavailability of Cl nanoemulsion compared to ethanol extract. This may be attributed to increase solubility of tested product into aqueous media of intestinal lumen. Giacomini and Sugiyama\(^{[36]}\) demonstrated that extremely water soluble and fat soluble substances are difficult to pass through a cell membrane. This was attributed to the poor absorption of the former through lipid bilayer of cell membrane, and the latter was attributed to low solubility in aqueous media of biological fluids because drug particles pass biological membranes in soluble form.

Method used in our study for nanoparticles preparation is simple, easy and inexpensive. The formation of spherical particles in the nanosize range was confirmed by transmission electron microscopy, which exhibited an average diameter of 24 nm (Figure 2) in contrast to 214 nm of the crude *C. longa* extract (Figure 1). Luz *et al.*\(^{[26]}\) succeeded in producing curcumin nanoparticles with an average size of 108 nm by incorporating curcumin into poly lactic-co-glycolic acid (PLGA) nanospheres using a nanoprecipitation technique. The characters of Cl nanoparticles are different from crude extract being highly soluble in water, ethanol, methanol and acetone. The crude extract did not dissolve in water but dissolved in ethanol and methanol and acetone. This confirms the experiment done by Jusnita *et al.*\(^{[25]}\) who showed that curcumin dissolves in acetone and ethanol.

---

Fig. 3: Schistosomicidal effect of Cl nanoemulsion on adult males and females *S. mansoni*. No statistical significant difference was detected (*P* > 0.05).

Fig. 4: Effect of Cl nanoemulsion on 24 hrs-old schistosomules. Data was statistically significant (*P* < 0.001).
The absorption and bioavailability of the hydrophobic active compound in the body would be enhanced by reducing the size of the molecule. Therefore, rapid penetration of small sized medicine particles occurs and raises the efficiency of the medicine\(^\text{[37]}\). The bioavailability of lipophilic components proved to be greater in nanocurcumin than conventional emulsions\(^\text{[38]}\). Our experiment demonstrated the in vitro penetration of nanocurcumin extract with a molecular diameter size lower than 100 nm as compared to the crude extract. Penetration of the curcumin extract across the membrane cell was 13.0% whereas the nanoemulsion penetrated by 20.0%. This indicated a greater penetration of nanocurcumin extract compared to the crude extract, and proved that the smaller molecular size would enhance the penetration across a cell in the body. Another experiment on transport through cells of the skin also showed that application of small size droplets would intensify transport of the active molecule\(^\text{[39]}\). According to He et al.\(^\text{[40]}\), food proteins stabilize nanoparticles used for delivering water insoluble drugs. This applies to using curcumin in nanoemulsion form as a delivery system to provide a large interfacial surface area for their absorption. In addition, Li and McClements\(^\text{[41]}\) observed that the small droplet size of the lipid phase increases the diffusivity of the cell.

The actual schistosomicidal activity of Cl nanoemulsion was evaluated in our study. A clear in vitro schistosomicidal effect on cercariae, schistosomules, and adults of S. mansoni was demonstrated. It caused death of 100% of cercaria, schistosomules, adult male and adult female worms with 100 μM at 30 min, 24 h, 48 h and 72 h, respectively. In the negative controls (RPMI medium), there were no dead cercariae or adult worms. Schistosomes infect their hosts by aquatic cercariae, which actively invade their host’s skin epidermis. Schistosomes survival fully depends on the cercarial invasion success. Many plant extracts were investigated for laboratory anti-cercarial activity. Marston and Hostettman\(^\text{[42]}\) reported seventeen effective compounds against cercariae. Lima et al.\(^\text{[43]}\) reported effectiveness of one single compound. However, these compounds were not active at lower concentrations or at shorter exposure times against cercariae. Chen et al.\(^\text{[44]}\) using pure curcumin reported effective low concentrations against cercariae. In our study, Cl nanoemulsion showed potential toxicity against cercariae indicating its successful use as a new water soluble cercaricidal agent.

The results showed that both male and female parasites are susceptible to Cl nanoemulsion. Magalhães et al.\(^\text{[21]}\) showed that curcumin caused 100% mortality of parasites at a concentration of 50 μM after 24 h, and no difference was observed between males and females. In contrast, it has been described in our study as well as in another study\(^\text{[45]}\) that male worms of S. mansoni are more susceptible than female worms. On the other hand, De Araújo et al.\(^\text{[3]}\) using nanoemulsion of a new schistosomicidal drug (BphEA) showed that male worms moved slowly at the end of 48 h whereas all the female worms died. The recorded decreased worm motility produced by Cl nanoemulsion has been described to be a result of inhibition of smooth muscle proliferation as well as inhibition of histamine induced smooth muscle contraction\(^\text{[46]}\). This is attributed to improvement of penetration of Cl nanoemulsion particles through parasite tegument, a result of increased passage of hydrophilic pits in schistosomal tegument and enhanced diffusion of nanoparticles. This occurs as a result of the increased solubility of tested product in biological media of the parasite. Claudineide et al.\(^\text{[46]}\) demonstrated that the schistosomal tegument consists of a double layer that encircles a smooth muscle layer and attaches to the suckers. The outer layer is glyocalx that contains hydrophilic pits. The inner one consists of lipid bilayer similar to human plasma membrane. The tegument exerts a regulatory role in passage of nutrients and excretion of lactic acid and metabolic wastes resulting from glycolysis in the parasite. In this context, the tegument represented an important target for anti-helmenthic drugs. In our study the female adult worm which has a thinner tegument and muscle layer showed resistance to the schistosomicidal effect of the tested plant. This contradictory result may be explained by the interaction of female hormonal factors on anti-proliferative and apoptosis promoting effect of tested plant.

In conclusion, curcumin appears as a new potential candidate drug against schistosomiasis. We successfully applied nanoemulsion preparation against different life stages of S. mansoni in vitro. Further study is needed to emphasis its effect in vivo. The employment of nanotechnology may offer a safe, effective and cheap treatment.

**Author Contribution:** NSM Aly conceived the study, wrote the manuscript, and shared AH Hussein in the study design. Besides, AH Hussein analyzed the data, HT Emam prepared the extract, and all authors performed the experiments and revised the manuscript.

**Financial support and sponsorship:** Nil.

**Conflicts of interest:** There are no conflicts of interest.

**REFERENCES**

3. Steinmann P, Keiser J, Bos R, Tanner M, Utzinger J. Schistosomiasis and water resources development:


C. longa nanoemulsion effect on S. mansoni Aly et al.


