Anthelmintic activity of the white wormwood, *Artemisia herba-alba* against *Heterakis gallinarum* infecting turkey poults

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Accepted 3 May, 2011

*Artemisia herba-alba* is widely used in the Egyptian folk medicine as vermifuge. The objective of this study was to evaluate the proclaimed anthelmintic efficacy of crude aqueous extracts of *A. herba-alba*, ACEA, in comparison to albendazole, ABZ, against *Heterakis gallinarum* infecting turkey poults. 60, 1 day old large white turkey poults (males) were divided into four groups. Group 1 was neither infested nor treated. Groups 2, 3 and 4 were inoculated with 500 embryonated eggs of *H. gallinarum* at 1 day old. On day 25 post infestation and for three successive days, group 3 was treated with ABZ, 2.5% (20 mg/kg B. wt.) and group 4 was treated with ACEA (0.4 g/kg B. wt.) in drinking water. The whole experiment had been repeated three times. Seven days post treatments, ABZ and ACEA reduce egg output (97.31 and 97.78%, respectively), and worm burden of *H. gallinarum* (95.08 and 96.07%, respectively). The weight and feed conversion ratios were improved in group 4. Biochemical analysis and histopathological sections revealed the adverse effect of ABZ. ACEA is then considered as a good anthelmintic alternative therapy and recommended in the control of ascaridosis in poultry industry, since it is effective, safe, available and cheap.

Key words: Heterakis, herbal remedies, *Artemisia*, albendazole, turkey.

INTRODUCTION

Helminthiasis is frequent among Egyptian birds (Khater, 1993). Infection with intestinal roundworms have been estimated to cause production losses in the range of 10 to 20% due to impaired feed conversion, reduced growth and egg production, and increased mortality (Ikeme, 1971; Soulsby, 1982; Choudury and Das, 1993; Seddiek et al., 2007). The nematode *Heterakis gallinarum* (Movsessian and Pkhrikian, 1994) (family: Ascarididae) is cosmopolitan in domestic chickens and related birds. The worm (1 to 2 cm in length) lives in the cecum, where they feed on its contents. *H. gallinarum* is one of the most important nematodes of poultry due to its role in the epidemiology of histomoniasis (blackhead disease) caused by a flagellate protozoan, *Histomonas meleagridis*, causing a particularly serious disease in turkeys (Papini and Cacciuttoli, 2008). *H. gallinarum* infections linked to histomoniasis have been well documented in chicken (Homer and Butcher, 1991; Permin, 2003). Blackhead disease causes high mortality in turkeys, sometimes approaching 100% of a flock. In chicken, the mortality may be 10 to 20% with high morbidity, although many outbreaks pass unnoticed (McDougald, 2005).

The life cycle of *H. gallinarum* is simple and direct, similar to that of *Ascaridia galli* with a minimum prepatent period of 22 days under temperate climatic conditions (Lund and Chute, 1972; Movsessian and Pkhrikian, 1994). After ingestion of the infective eggs and hatching of eggs in the upper small intestine, the larvae reached the caeca at the end of 24 h post-infestation (PI). The larvae are embedded in the mucosal layer of the caeca.

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for a varying period of 3 to 12 days (Lund and Chute, 1972), and then the mature worms infest the lumen of blind caeca. Fertilization occurred and oviposition starts 22 to 25 days PI (Movsesian and Pkhrikian, 1994). Eggs of *H. gallinarum* contain a zygote when laid. They develop into the infective stage in 12 to 14 days at 22°C and can remain infective for four years in soil. Birds allowed to roam a barnyard usually are infected (Jansson et al., 2004; Roberts and Janovy, 2005).

If eaten by an earthworm, a juvenile may hatch and become dormant in the worm's tissues, remaining infective for at least a year. Since these nematodes do not develop further until eaten by a bird, an earth worm is a paratenic host (Roberts and Janovy, 2005). Diagnosis of *H. gallinarum* is based on faecal isolation of eggs or direct identification of adult worms in the intestine (Soulsby, 1982; Roberts and Janovy, 2005). *H. gallinarum* caused severe caecal alterations in the turkey poult characterized by necrosis, chronic typhlitis, haemosidrosis and nodular formation in the caeca (Menezes et al., 2003; Brener et al., 2006). The increasing prevalence of anthelmintic resistant strains of helminths (Walter and Prichard, 1985; Kaplan, 2004; Hoque et al., 2003; Borgsteede et al., 2007; Beech et al., 2011), drug residues in animal products (Kaemmerer and Butenkov, 1973; McKellar, 1997), and high cost of conventional anthelmintics have created an interest in studying medicinal plants as an alternative source of anthelmintics. The use of plant extracts as mendicants may alleviate these obstacles. An additional constraint in anthelmintic use comes from the consumer and the ever-increasing need for the drug-free production of foods (Harper and Makatouni, 2002) as they are not only natural products but may comprise new therapeutic molecules to which resistance has not yet developed.

Medicinal plants have been used as a source of remedies since ancient times. The ancient Egyptians were familiar with many medicinal herbs and were aware of their usefulness in treatment of various diseases. When the Arabs came to Egypt, Arabic medicine was practiced and the “art” of healing made use of all available knowledge gained from different civilizations such as the Persian, Chinese, Greek, as well as the Ancient Egyptian.

It could create an herbal remedy export market and thereby create more jobs in the country. Egypt possesses an enormous diversity of plant resources that is useful for herbal remedies for humans and animals (Hifnawy et al., 2001; Khater, 2003; Shalaby and Khater, 2005; El Garhy and Mahmoud, 2007; Khater and Shalaby, 2008; Mobarak et al., 2008; Khater and Khater, 2009; Khater et al., 2009, 2011). Many *Artemisia* spp. (family: Asteraceae, formerly Compositae) have a prominent position in the herbal de-worming literature (El Garhy and Mahmoud, 2002; Iqbal et al., 2004; Caner et al., 2008; Urban et al., 2008; Tariq et al., 2009; Bashtar et al., 2011).

They also have a high value in several fields, as food plants (Benmansour et al., 1990; Fenardji et al., 1994; Benmansour and Taleb-Bendiab, 1998), anticoagulant (Naidoo et al., 2008; del Cacho et al., 2010), and antimalaria (World Health Organization, 2005; Enene et al., 2009) in medicine. The Romans used dried, unexpanded flower heads obtained from several species of the genus *Artemisia* in the first century, for the treatment of *Ascaris, Enterobius* and tapeworm infections and it became an important member of the European pharmacopoeia until the early 20th century (Mohamed et al., 2010).

The Anthelmentic effect of *Artemisia* species is caused by the sesquiterpene lactone, santonin (Rachkovskaya, 1978; Akhtar et al., 1982). *Artemisia herba-alba*, known as desert or white wormwood, known in Arabic as shih and in French as Armoise Blanche, is widely distributed in North Africa. It is used traditionally by the Egyptians as a vermifuge in addition to its other medical and veterinary uses (Saleh et al., 2006; Seddiek et al., 2007; Mobarak et al., 2008; Mohamed et al., 2010).

It has been used in folk medicine by many cultures since ancient times, used in Moroccan folk medicine to treat arterial hypertension and/or diabetes (Ziyat et al., 1997; Tahraoui et al., 2007; Zeggwagh et al., 2008). During an ethnopharmacological survey carried out among the Bedouins of the Negev desert, it was found that *A. herba-alba* relieved stomach disorders (Friedman et al., 1986). In addition to the previous benefits *A. herba-alba* induced hypoglycemic effect (Al-Waili, 1988; Marrif et al., 1995). Moreover, the aqueous extract and essential oil of *A. herba-alba* expressed antileishmanial activity against *Leishmania* major (Hatimi et al., 2001). Herbal tea from this species has been used as antibacterial, analgesic, antiinflammatory and hemostatic agents (Laid et al., 2008). The plant is used to bandage wounds, cure stomach-ache, and neuralgia and other pains when mixed with henna and diluted in water and applied to the head (Le Floch, 1983). Alternative treatments for gastrointestinal helminths in poultry (for example anthelmintic plants) have been investigated in several studies (Mali et al., 2007; Brito et al., 2009; Kosalge and Fursule, 2009; Parida et al., 2010).

However, scientifically validated data on the efficacy of herbal treatments against *H. gallinarum* remain scarce and mainly in vivo (Nagaich, 2000). As the people consume *A. herba-alba* to cure helminthic infections as per the literature, we attempted to investigate this medicinal plant for its claimed anthelmintic activity. The present study was designed to investigate the efficacy of treatments with aqueous cured extract of *A. herba-alba* (ACEA) and albendazole (ABZ), as a reference anthelmintic, on turkey poult infected with *H. gallinarum* eggs.

Therefore, several parameters have been evaluated such as fecal egg count reduction (FECR), worm count reduction (WCR), growth performance (body weight, body gain and food conversion ratio, FCR), biochemical,
and histopathological features.

MATERIALS AND METHODS

Parasite

*H. gallinarum* adults were obtained from the caeca (blind portion) of the freshly killed turkey poults (naturally infested) and washed several times in saline. Heterakis ova were obtained by gentle crushing of gravid female worms, with a small spatula, through a 150 μm sieve into small Petri dishes containing distilled water (2 to 3 mm in depth). Few drops of 2% formalin solution had been added to each Petri dish, and then incubated for 21 to 28 days at 28 ± 2°C to permit embryonation of the eggs (Oliver, 1953).

Experimental design

To study the effect of *A. herba-alba* on *H. gallinarum* infection, a total of 180 male turkey poults were used. The complete experimental procedure was repeated 3 times to obtain consistent results. In each experiment, 60, 1 day-old poults were divided into four groups consisting of 15 birds each. Birds were placed in wire-floored cages measuring 50 × 50 cm, with an independent supply of water and food and elevated approximately 50 cm above the litter. A total of 5 birds were placed in one cage and 3 cages were used for each group. During the experimental period, birds were given balanced commercial starter ration and water ad-libitum from 0 to 42 days of age. Animals in Groups 2, 3 and 4 were infected, intra crop through stomach tube, with a single dose of 500 embryonated eggs of *H. gallinarum* 1 day after hatching of turkey poults, according to Permin et al. (1997). Birds in Groups 1 and 2 were fed with food and water free of anthelmintic compounds (non-medicated control birds). Group 1 was considered as the negative control group (non-infected, non medicated) and Group 2 was taken into account as the positive control group (infected, non medicated).

25 days post infection, birds in Groups 3 and 4 (medicated birds) received anthelmintic compounds in drinking water for 3 consecutive days. Group 3 was treated with ABZ suspension 2.5% (Arabcomed Co, Egypt) in a dose of 20 mg/kg B. wt; whereas, Group 4 received crude aqueous extract of *A. herba-alba* (ACEA) in a dose of 0.4 g/kg B. wt. ACEA was prepared by using the soaking method of the shoots (leaves and stems).

The shoots at a dose of 0.4 g/kg B. wt. were soaked in a known volume of distilled water for 24 h (Marrif et al., 1995), then sieved (stock). The dose for one day was calculated using the formula: Dose = 0.4 × average weight of birds at the day of treatment × number of birds. Birds were starved overnight prior to treatment with the drug the following morning. The drug was dissolved in the drinking water and made available over a 6 to 8 h period. The clinical signs were recorded. After slaughtering, the intestinal tract was examined and PM lesions were recorded. Egg counts per gram (EPGs) were determined in excreta samples taken from each subgroup at days 25 and 32 of age (just before treatment and 7 days post treatment, respectively) to evaluate the degree of infestation using modified McMaster technique (Thienpont et al., 1986).

The numbers of adult worms were recorded in five sacrificed poults (per group) on the 25th day post-infestation (before treatment) as well as on the 7th day after treatment according to Permin and Hansen (1998). Feed conversion ratio (FCR) values were calculated weekly as the ratio of feed intake to weight gain. The mean weight gain (MWG) was calculated using the formula:

\[
MWG = \frac{(\text{mean final weight of live parakeets birds in a cage}) - (\text{mean initial weight of all parakeets birds in that cage}) + (\text{weight of dead parakeets birds})}{(\text{mean initial weight of all parakeets birds in that cage}) + (weight of dead birds)}
\]

The group feed conversion ratio (FCR) for the study period was calculated using the formula:

\[
FCR = \frac{\text{feed consumed per group (g)}}{(\text{weight gain of surviving birds} + \text{weight gain of dead birds})}
\]

Individual FCR were not calculated as animals were fed as a group. At day 32 of age, seven days post treatment, 5 birds per group were sacrificed for detection of post mortem lesions and getting liver specimens which immediately fixed in 10% neutral buffered formalin. Paraffin sections were stained with hematoxylin and eosin (H&E) and examined microscopically according to Bancroft et al. (1996).

Biochemical analysis

Blood samples were collected from the wing vein of 5 birds per group on the 7th day post treatment. Each sample was allowed to separate the serum and kept at -20°C till biochemical analysis. Several tests were performed for determination of the serum Alanine aminotransferase (ALT), Aspartate aminotransferase (AST) enzyme activity (Reitman and Frankel, 1957), total protein (Weichselbaum, 1946), albumin (Doumas, 1971), globulins (difference between total protein and albumin), serum uric acid (Haisman and Muller, 1977), and creatinine (Husdan and Rapaport, 1968).

Data analysis

The differences among experimental treatments were tested at P ≤ 0.05 by one-way ANOVA according to Duncan (1955) and Snedecor and Cochran (1969) using the SPSS program (SPSS v.11, SPSS 1986).

RESULTS

Once infected it took approximately 14 days for the birds to demonstrate clinical signs of infection. The observed clinical signs were depression (fluffed up feathers, appetite loss, irritability, and feather plucking), dullness, loss of appetite, emaciation and unthriftiness. The necropsy (on the 25th day post-infestation) revealed the presence of adult worms and inflammation of the caeca indicating typhilitis, thickening and nodular formation in the caecal mucosa. There were no mortalities among birds. Seven days post treatment, ABZ and ACEA treated groups were significantly effective (P ≤ 0.05) in reducing egg outputs, FECRs were 97.31 and 97.78%, respectively, when compared with that of the infected, non medicated group, 0% (Table 1).

In addition, the mean number of worm burden in the medicated groups (Groups 3 and 4) was significantly (P ≤ 0.05) decreased, WCRs were 95.08 and 96.07%, respectively, when compared with that of the positive control group (Table 2). Body weight, body weight gain and feed conversion ratio values till the age of 21 days did not differ among infected groups (Groups 2, 3 and 4). At the age of 6 weeks, such parameters in turkey poults treated with ACAE were significantly (P ≤ 0.05) improved...
when compared with those of the positive control and albendazole treated groups (Table 3). Among ABZ treated group, some biochemical parameters were significantly (P ≤ 0.05) altered when compared with those of the other groups (Groups 1, 2 and 4). Such alterations included reduction in the total protein, globulin and albumin levels and elevation of the levels of ALT, AST, creatinine and uric acid (Table 4).

Microscopically, male *H. gallinarum* infested the caeca of turkey pouls is very conspicuous with two unequal spicules (Figure 1). Histological sections of the liver of turkey pouls treated with ABZ revealed mild vacuolar degeneration of the hepatocytes and moderate numbers of mixed inflammatory cells in sinusoids (Figure 2). Whereas those of the treated group with ACAE indicated normal histological appearance of the hepatocytes (apparently healthy) and low numbers of inflammatory cells in sinusoids (Figure 3).

**DISCUSSION**

The infestation of turkey pouls with *H. gallinarum*, in this study, appeared to cause depression, dullness, emaciation, dehydration and lower locomotion. In addition, cross sections of cecum of infected turkey pouls indicated the presence of *H. gallinarum* and inflammatory reaction of the submucosa. Similar observations were reported (Brener et al., 2006).

Pathological changes included congestion, hemorrhages and nodules with necrotic center in the caecum were noted. Alike lesions were recorded (Choudury and Das, 1993; Roberts and Janovy, 2005; Menezes et al., 2003; Brener et al., 2006). Because of the longevity of the eggs, 4 years, it is difficult to eliminate *H. gallinarum* form a domestic flock. Although adult chickens may affect a self-cure, infective eggs are available for the following spring, when new chicks hatch. Furthermore, as earth worms feed in contaminated soil, they accumulate large numbers of juveniles, which in turn cause massive infections in unlucky birds that eat them (Roberts and Janovy, 2005). Generally speaking, *H. gallinarum* is not highly pathogenic in itself. However, *H. meleagris*, is transmitted between birds within eggs of *H. gallinarum* (Long et al., 1987) leading to histomoniasis which cause necrosis of the caecal mucosa, swelling of the caecum and liver necrosis (Papini and Cacciuttoli, 2008).

Consequently, reducing the number of eggs in litter through treatment of anthelmintics, is a highly desirable feature which should be taken into account in the control of ascaridiasis that is characterized by high biotic potential and large number of eggs which may accumulate in deep letter houses (Davis and Jayner, 1955). Concerning conventional anthelmintics, our results indicated that ABZ was highly effective in controlling *H. gallinarum* as FECR was 97.31% and WCR was 95.08%. Analogues to our result, ABZ in a single dose of 20 mg/kg B. wt. is highly effective in the treatment of chickens for

### Table 1. Eggs per gram of faeces (EPG) of turkey pouls infested with *Heterakis gallinarum* invasive eggs and treated with albendazole suspension and *Artemisia herba-alba* aqueous extract.

<table>
<thead>
<tr>
<th>Age in days</th>
<th>Group 1 -ve control</th>
<th>Group 2 +ve control</th>
<th>Group 3 ABZ</th>
<th>Group 4 ACAE</th>
<th>LSD</th>
</tr>
</thead>
<tbody>
<tr>
<td>25 days</td>
<td>0.00 ± 0.00</td>
<td>150.00 ± 6.5</td>
<td>154.50 ± 2.61</td>
<td>153.42 ± 2.84</td>
<td>153.00*</td>
</tr>
<tr>
<td>32 days</td>
<td>0.00 ± 0.00</td>
<td>153.76 ± 2.66</td>
<td>4.25 ± 0.46</td>
<td>3.50 ± 1.68</td>
<td>4.25*</td>
</tr>
<tr>
<td>Production (%)</td>
<td>0</td>
<td>100</td>
<td>2.70</td>
<td>2.22</td>
<td></td>
</tr>
<tr>
<td>FECR (%)</td>
<td>0</td>
<td>97.31</td>
<td>97.78</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Eggs per gram faeces (No. X 1000); (Mean ± SE; n = 12); Data were analyzed by one way ANOVA. LSD: least significance difference among means at P ≤ 0.05. Means with different alphabetical superscripts in the same row are significantly different. ABZ: albendazole. ACAE: crude aqueous extract of *Artemisia herba-alba*. 25 days (just before treatment); 32 days (7 days post treatment). FECR= Reduction % of fecal egg count.

### Table 2. Worm burden of turkey pouls infested with *Heterakis gallinarum* invasive eggs and treated with albendazole suspension and *Artemisia herba-alba* aqueous extract.

<table>
<thead>
<tr>
<th>Age in days</th>
<th>Group 1 -ve control</th>
<th>Group 2 +ve control</th>
<th>Group 3 ABZ</th>
<th>Group 4 ACAE</th>
<th>LSD</th>
</tr>
</thead>
<tbody>
<tr>
<td>25 days</td>
<td>0.00 ± 0.00 b</td>
<td>67.20 ± 1.86 a</td>
<td>66.40 ± 0.81 a</td>
<td>66.20 ± 2.13 a</td>
<td>66.20</td>
</tr>
<tr>
<td>32 days</td>
<td>0.00 ± 0.00 c</td>
<td>61.00 ± 0.71 a</td>
<td>3.00 ± 0.32 b</td>
<td>2.40 ± 0.25 b</td>
<td>2.40</td>
</tr>
<tr>
<td>Production (%)</td>
<td>0</td>
<td>100</td>
<td>4.92</td>
<td>3.93</td>
<td></td>
</tr>
<tr>
<td>WCR (%)</td>
<td>0</td>
<td>95.08</td>
<td>96.07</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

(Mean ± SE; n = 5); Data were analyzed by one way ANOVA. LSD: least significance difference among means at P ≤ 0.05. Means with different alphabetical superscripts in the same row are significantly different. ABZ: albendazole. ACAE: crude aqueous extract of *Artemisia herba-alba*. 25 days (just before treatment); 32 days (7 days post treatment). WCR= Reduction % of worm count.
Table 3. Growth performance parameters of turkey poults infested with *Heterakis gallinarum* invasive eggs and treated with albendazole suspension and *Artemisia herba-alba* aqueous extract.

<table>
<thead>
<tr>
<th>Time (days)</th>
<th>Growth performance parameters</th>
<th>Group 1 -ve control</th>
<th>Group 2 +ve control</th>
<th>Group 3 ABZ</th>
<th>Group 4 ACAE</th>
<th>LSD</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 day old</td>
<td>Body weight</td>
<td>65.67 ± 0.95</td>
<td>66.87 ± 0.96</td>
<td>45.20 ± 2.90</td>
<td>65.93 ± 0.90</td>
<td>NS</td>
</tr>
<tr>
<td></td>
<td>Body weight</td>
<td>239.33 ± 0.86</td>
<td>231.47 ± 1.03</td>
<td>233.83 ± 1.39</td>
<td>233.13 ± 1.38</td>
<td>5.40*</td>
</tr>
<tr>
<td>7 day old</td>
<td>Body weight</td>
<td>45.20 ± 2.90</td>
<td>171.53 ± 1.91</td>
<td>162.60 ± 1.57</td>
<td>164.87 ± 0.85</td>
<td>5.40*</td>
</tr>
<tr>
<td></td>
<td>FCR</td>
<td>1.31 ± 0.01a</td>
<td>1.33 ± 0.01a</td>
<td>1.32 ± 0.02a</td>
<td>1.32 ± 0.01a</td>
<td>NS</td>
</tr>
<tr>
<td>14 day old</td>
<td>Body weight</td>
<td>249.40 ± 1.81</td>
<td>231.47 ± 1.03</td>
<td>233.83 ± 1.39</td>
<td>233.13 ± 1.38</td>
<td>91.10*</td>
</tr>
<tr>
<td></td>
<td>Body gain</td>
<td>215.60 ± 2.36</td>
<td>162.60 ± 1.57</td>
<td>164.87 ± 0.85</td>
<td>166.13 ± 1.39</td>
<td>5.40*</td>
</tr>
<tr>
<td></td>
<td>FCR</td>
<td>1.35 ± 0.01b</td>
<td>1.58 ± 0.02a</td>
<td>1.58 ± 0.01a</td>
<td>1.57 ± 0.01a</td>
<td>0.23*</td>
</tr>
<tr>
<td>21 day old</td>
<td>Body weight</td>
<td>320.80 ± 3.03</td>
<td>191.20 ± 3.35</td>
<td>192.27 ± 2.68</td>
<td>193.93 ± 2.78</td>
<td>127.87*</td>
</tr>
<tr>
<td></td>
<td>Body gain</td>
<td>1.54 ± 0.02b</td>
<td>2.41 ± 0.03a</td>
<td>2.39 ± 0.03a</td>
<td>2.41 ± 0.02a</td>
<td>0.86*</td>
</tr>
<tr>
<td>28 day old</td>
<td>Body weight</td>
<td>1261.67 ± 5.18</td>
<td>757.47 ± 2.82</td>
<td>947.33 ± 3.18</td>
<td>951.27 ± 8.45</td>
<td>189.87*</td>
</tr>
<tr>
<td></td>
<td>Body gain</td>
<td>491.53 ± 4.67</td>
<td>200.47 ± 5.49</td>
<td>398.73 ± 7.69</td>
<td>401.13 ± 8.04</td>
<td>90.40*</td>
</tr>
<tr>
<td></td>
<td>FCR</td>
<td>1.55 ± 0.01c</td>
<td>2.43 ± 0.02a</td>
<td>1.87 ± 0.02b</td>
<td>1.87 ± 0.02b</td>
<td>0.32*</td>
</tr>
<tr>
<td>35 day old</td>
<td>Body weight</td>
<td>1814.13 ± 6.87</td>
<td>1049.47 ± 10.94</td>
<td>1406.00 ± 8.1</td>
<td>1415.33 ± 8.27</td>
<td>356.53*</td>
</tr>
<tr>
<td></td>
<td>Body gain</td>
<td>545.87 ± 5.99</td>
<td>290.60 ± 12.34</td>
<td>458.67 ± 8.02</td>
<td>465.40 ± 8.67</td>
<td>80.47*</td>
</tr>
<tr>
<td></td>
<td>FCR</td>
<td>1.57 ± 0.02d</td>
<td>2.58 ± 0.03a</td>
<td>1.92 ± 0.02b</td>
<td>1.81 ± 0.03c</td>
<td>0.11*</td>
</tr>
<tr>
<td>42 day old</td>
<td>Body weight</td>
<td>2754.67 ± 6.24</td>
<td>1392.00 ± 15.19</td>
<td>2235.33 ± 7.41</td>
<td>2292.34 ± 7.79</td>
<td>36.00*</td>
</tr>
<tr>
<td></td>
<td>Body gain</td>
<td>947.20 ± 4.80</td>
<td>323.27 ± 18.48</td>
<td>829.00 ± 9.24</td>
<td>867.33 ± 8.85</td>
<td>38.33*</td>
</tr>
<tr>
<td></td>
<td>FCR</td>
<td>1.61 ± 0.01d</td>
<td>2.41 ± 0.04a</td>
<td>1.88 ± 0.01b</td>
<td>1.70 ± 0.02d</td>
<td>0.10*</td>
</tr>
</tbody>
</table>

Mean ± SE, n = 15; Data were analysed by one-way ANOVA. ACAE: crude aqueous extract of *Artemisia herba-alba*. ABZ: albendazole; LSD: least significance difference among means at P ≤ 0.05, NS = non significant; Body gain (g); Means with different alphabetical superscripts in the same row are significantly different.

*H. gallinarum* and *A. galli* as its efficacies for controlling larvae and adult burden were 98.9 and 94.9% for *H. gallinarum* and 98.2 and 100% for *A. galli*, respectively (Tucker et al., 2007). Phenothiazine is effective against *H. gallinarum, in vitro* (Oliver, 1953). Unfortunately, the increasing prevalence of anthelmintic resistant strains of helminths (Walter and Prichard, 1985; Kaplan, 2004; Beech et al., 2011), for example albendazole resistance in gastrointestinal nematode parasites (Hoque et al., 2003; Borgsteede et al., 2007) has been developed. Many great research challenges and prospects for the identification of new, safe and environmentally acceptable anthelmintics such as medicinal plants
were effective against and ethanolic extracts of A. galli charantia Sapindus trifoliatus, Butea frondosa gallinarum treating methanolic extract of homeostasis which is essential for the development of the nitrate generation which could interfere in local (Mohamed et al., 2010) which is capable of reducing death (John et al., 2009).

proteins in the gastrointestinal tract of the host animal or polyphenolics and related constituents (Mohamed et al., 2010) effect, such as santonin (Khafagy et al., 1971) and eight extract (CAE) (Iqbal et al., 2004).

infested sheep with treatment of different species of nematodes in naturally 2011). Lower FECR (67.2%) has been recorded after against chickens (Seddiek et al., 2012) and 97.31 and 97.78%, respectively, and worm burden was significantly reduced (95.08 and 96.07%) in both medicated groups (ABZ and ACEA, respectively) than that of the positive control groups. Similar to our results, complete reduction of egg production (100%) has been recorded for ACAE against A. galli infecting chickens (Seddiek et al., 2007) and Artemisia cina against Moniezia spp. infecting sheep (Bashtar et al., 2011). Lower FECR (67.2%) has been recorded after treatment of different species of nematodes in naturally infested sheep with Artemisia brevifolia crude aqueous extract (CAE) (Iqbal et al., 2004). A. herb-alba possesses several constituents that induced anthelmintic effect, such as santonin (Khafagy et al., 1971) and eight polyphenolics and related constituents (Mohamed et al., 2010). Synthetic phenolic anthelmintics interfere with the energy generation in the helminth parasites by uncoupling the oxidative phosphorylation. Another possible mechanism of action is that they bind to free proteins in the gastrointestinal tract of the host animal or to glycoprotein on the cuticle of the parasite and cause death (John et al., 2009).

Additionally, plant possess “antioxidant activity” (Mohamed et al., 2010) which is capable of reducing the nitrate generation which could interfere in local homeostasis which is essential for the development of helminthes (Borba et al., 2010). Similar to our results, methanolic extract of Caesalpinia cristaa Linn. seeds and piperazine (200 mg/kg) are equieffective, in vivo, in treating A. galli infecting chickens (Javed et al., 1994). Several anthelmintic plants are effective in vitro, such as A. sativum has shown anthelmintic action against H. gallinarum and A. galli (Nagaich, 2000); Carica papaya, Sapindus trifoliatu, Butea frondosa and Momordica charantia were more effective against A. galli than piperazine hexahydrate (Lal et al., 1976); and aqueous and ethanolic extracts of Morinda citrifolia fruit (noni) were effective against A. galli (Brito et al., 2009). Use of A. galli and Rallietina species as a suitable model for screening of anthelmintic drug was advocated earlier (Kaushik et al., 1974). Furthermore, several assays were performed in vitro using adult earthworm (Phereetima posthuma) for preliminary evaluation of anthelmintic activity, owing to its anatomical and physiological resemblance with the intestinal roundworm parasites, such as Ascaris lumbricoids, of human beings (Dash et al., 2002; Shivkumar and Kumar et al., 2003). Some herbs show in vitro anthelmintic activity against P. posthuma, tapeworms (Rallietina spiralis) and roundworms (A. galli) such as the aqueous extract of Thespesia lampas (Cav.) roots (Kosalge and Fursule, 2009) and crude extracts and fractions of Pterospermum acerifolium (Parida et al., 2010). In addition, the crude alcohol and aqueous extracts of the seeds of Cleome viscosa. Linn. shows anthelmintic activity against P. posthuma and A. galli (Mali et al., 2007). Santonin present in A. herb- alba (Khafagy et al., 1971) induces an anthelmintic effect. The vermifuge effect of santonin substance, prepared from the Artemisia spp., induced changes in the musculocutaneous sac (cuticle, hypoderm and muscle cells) of the worm through its direct action on muscle cells of the worm resulting in complete relaxation of its muscular layer leading to its expulsion to outside (Rachkovskaiia, 1978).

Akhtar et al. (1982) reported that the percentage reduction in EPG counts in the calves naturally acquired Neoascaris vitulorum and treated with 15 mg/kg of santonin on the seventh day, these values were 100, 100 and 99.7% in moderate, high and heavily infected calves, respectively. Both piperazine and santonin were associated with some side effects like diarrhea, restlessness, etc. Santonin has an efficacy similar to piperazine given at the 88 mg/kg dose level for the treatment of ascariasis in buffalo calves. Low concentrations of santonin are reported to have a selective toxic action on the ganglion located in the nerve ring of Ascaris spp. (Sollman, 1957). Against other nematodes, such as Oxyuris spp and cestodes, santonin is not effective (Steinegger and Hänsel, 1972). With regard to weight performances, the present study

Table 4. Biochemical parameters of turkey poults infested with Heterakis gallinarum invasive eggs and treated with albendazole suspension or Artemisia herba-alba aqueous extract.

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Group 1 -ve control</th>
<th>Group 2 +ve control</th>
<th>Group 3 ABZ</th>
<th>Group 4 ACAE</th>
<th>LSD</th>
</tr>
</thead>
<tbody>
<tr>
<td>ALT (IU/L)</td>
<td>18.80±0.31&lt;sup&gt;b&lt;/sup&gt;</td>
<td>18.15±0.27&lt;sup&gt;b&lt;/sup&gt;</td>
<td>21.10±0.29&lt;sup&gt;a&lt;/sup&gt;</td>
<td>19.04±0.30&lt;sup&gt;b&lt;/sup&gt;</td>
<td>5.94*</td>
</tr>
<tr>
<td>AST (IU/L)</td>
<td>217.58±0.96&lt;sup&gt;b&lt;/sup&gt;</td>
<td>218.25±2.16&lt;sup&gt;b&lt;/sup&gt;</td>
<td>262.40±2.33&lt;sup&gt;a&lt;/sup&gt;</td>
<td>221.70±2.42&lt;sup&gt;b&lt;/sup&gt;</td>
<td>42.26*</td>
</tr>
<tr>
<td>Total protein (gm/dl)</td>
<td>4.34±0.20&lt;sup&gt;a&lt;/sup&gt;</td>
<td>4.35±0.21&lt;sup&gt;a&lt;/sup&gt;</td>
<td>3.40±0.17&lt;sup&gt;b&lt;/sup&gt;</td>
<td>4.86±0.13&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.78*</td>
</tr>
<tr>
<td>Serum albumin (gm/dl)</td>
<td>1.62±0.06&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.75±0.06&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.59±0.06&lt;sup&gt;b&lt;/sup&gt;</td>
<td>1.16±0.14&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.50*</td>
</tr>
<tr>
<td>Serum globulin (gm/dl)</td>
<td>2.72±0.07&lt;sup&gt;b&lt;/sup&gt;</td>
<td>2.60±0.05&lt;sup&gt;b&lt;/sup&gt;</td>
<td>2.81±0.08&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.70±0.07&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.89*</td>
</tr>
<tr>
<td>Creatinine (IU/L)</td>
<td>0.08±0.01&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.07±0.005&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.11±0.009&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.07±0.004&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.02*</td>
</tr>
<tr>
<td>Uric acid (mg/dl)</td>
<td>3.40±0.21&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.59±0.22&lt;sup&gt;b&lt;/sup&gt;</td>
<td>5.40±0.35&lt;sup&gt;a&lt;/sup&gt;</td>
<td>3.70±0.18&lt;sup&gt;b&lt;/sup&gt;</td>
<td>1.60*</td>
</tr>
</tbody>
</table>

Mean ± SE, n = 5; Data were analysed by one-way ANOVA. ACAE: crude aqueous extract of Artemisia herba-alba. ABZ: albendazole; LSD: least significant difference among means at P ≤ 0.05. Means with different alphabetical superscripts in the same row are significantly different.
indicated that infection with *H. gallinarum* significantly reduced body weight, body gain and increased FCR values highlighting the detrimental effect of the infection with this parasite on the performance of pouls.

Similarly, reduction of growth performance in chickens of 2 weeks PI had been recorded (Choudury and Das, 1993). ABZ and ACEA treatments exerted beneficial effects by significantly improving body weight gain and feed conversion ratio values compared to the positive control group. Moreover, ACEA significantly (*P* ≤ 0.05) enhanced growth performance of turkey pouls than that of ABZ. Comparable to our results, *A. herba-alba* improved the growth performance of chickens infested with *A. galli* (Seddiek et al., 2007) and chickens fed on ration contaminated with aflatoxin-B1 (Mobarak et al., 2008). In addition to the anthelmintic effect of the used herb, it improved weight performances indirectly because of its other biological activities. *A. herba-alba* improves the general health of infected turkeys because it possess antibacterial (Yashphe et al., 1979; Juteau et al., 2002; Laid et al., 2008; Elturbi et al., 2011), anti-oxidant (Aniya et al., 2000; Juteau et al., 2002; Kim et al., 2003; Kadri et al., 2011), and antifungal activities (El-Shayeb and Mabrouk, 1984; Saleh et al., 2006).

Moreover, *A. herba-alba* is used for digestive disorders, abdominal pain, colic and liver failure (Le Floc’h, 1983). Moreover, santonin induces significant antipyretic, anti-inflammatory effect and inhibit granuloma formation (Al-Habbi et al., 1994). Ideally, plants should provide, besides the anthelmintic effect, an alternative source of nutrition for animals of *A. herba-alba* is also suggested to be important as a fodder for sheep and for livestock in the plateau regions of Algeria where it grows abundantly (Benmansur et al., 1990; Fenardji et al., 1994; Benmansour and Taleb-Bendiab, 1998).

The liver of turkey pouls treated with ABZ showed vacuolar degeneration in the hepatocytes indicating the toxic effect of ABZ on the liver cells, meanwhile the liver of turkey pouls treated with ACEA showed no degenerative changes in hepatocytes which seem apparently healthy. The histopathological results ensured that ACEA has no adverse effect on hepatocytes. Biochemical analysis in the present study indicated the side effect of treatment with albendazole, for example reduction in the total protein and albumin levels and elevation of the levels of serum ALT and AST enzymes, when compared with that of ACEA medicated group. This may be due to side (toxic) effect of ABZ on the liver cells. Similar results were recorded in human (Choi et al., 2008) and in rat (Abd El-Rahman et al., 1999). Our results indicated that ACEA has no side effect on the liver cells. *A. herba-alba* has hepatoprotective effect (Aniya et al., 2000; Israpil et al., 2002) and it enhances bilirubin clearance (Mobarak et al., 2008). The adverse effect of ABZ on kidney functions has been confirmed by the elevated levels of creatinine and uric acid. Such elevation was not recorded for ACEA. A comparable result has been recorded for *A. herba-alba* (Marrif et al., 1995). A single dose of ABZ is safe and no adverse effects were observed on bird appearance, behavior, apparent appetite and weight gain (Tucker et al., 2007). Although ABZ is one of the most important antiparasitic drugs with high margin of safety, some unwanted side effects cannot be ignored (Abd El-Rahman et al., 1999). The adverse effect of ABZ in Group 3 on the liver and kidney functions and tissue may be due to the use of ABZ for somewhat longer time, to imitate what farmers in Egypt do to ensure that all birds get treated, than that recommended by the producer (20 mg/kg B. wt. for 3 successive days instead of a single dose).

The increase of ALT and AST enzymes was obtained in rat given ABZ in a dose of 400 mg/kg B. wt. as a single dose (Abd El-Rahman et al., 1999). Moreover, birds of the order Columbiformes, such as pigeon and dove, are susceptible to toxicosis after ABZ and fenbendazole (FBZ) administration (Howard et al., 2002). Regarding the nematocidal activity of *A. herba-alba*, ACEA “eradicate” intestinal infection with *Enterobius vermicularis* within 3 days in all 10 patients treated (Al-Waili, 1988). Ascarididae from hogs and ground worms were killed by the oil of *the Libyan* *A. herba-alba* in a short time (Callegari and Rossi, 1939, 1940). The powdered shoots of *A. herba-alba* expressed anthelmintic activity against experimental haemonchosis in Nubian goats which manifested by the

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**Figure 1.** Posterior end of *H. gallinae* adult worm (male), ventral view, infesting the caecum of turkey pouls showing two unequal spicules.
Figure 2. Liver of turkey poults infected with *H. gallinae* and treated with albendazole suspension (2.5%) showing mild vacuolar degeneration (arrow) of the hepatocytes and moderate numbers of mixed inflammatory cells (arrow head) in sinusoids stained with hematoxylin and eosin (X 400).

Figure 3. Liver of turkey poults infected with *H. gallinae* and treated with *A. herba-alba* extract showing normal histological appearance of the hepatocytes and low numbers of inflammatory cells (arrow head) in sinusoids stained with hematoxylin and eosin (X 400).
absence of eggs in the faeces or adult worms in the abomasum at necropsy and of significant lesions in the tissues of the goats and return of the concentrations of serum ammonia, sodium, potassium, total protein, creatinine and aspartate aminotransferase (GOT) to normal (Idris et al., 1982). The leaf extract of A. herba-alba was the most effective among twenty Jordanian plant species against two species of root-knot nematodes, in vitro (Al-Banna et al., 2003). In addition to its nematicidal activity, A. herba-alba induces pesticidal activity as it “induced” larvicidal activity against Culex pipiens mosquito, insecticidal activity against houseflies Musca domestica L., and rodenticidal activity against white mice Mus musculus (Hifnawy et al., 2001).

Furthermore, extracts of A. herba-alba are highly effective against arthropods of agricultural importance, such as Tetranynchus cinnabarinus mites (Azaiiez et al., 2007), Bemisia tabaci (Gennadius), Aphis gossypii (Glover) and Thrips tabaci (Lindman) (Soliman, 2006, 2007), Acanthoscelides obtectus (Coleoptera: Bruchidae) (Tani et al., 2008), A. obtectus, responsible for green beans rot (Derwich et al., 2009) and cotton leafworm Spodoptera littoralis (Biosd.) larvae (Hifnawy et al., 2001). Due to narrow therapeutic window (safety index) of and toxicity of A. herba-alba, the crude drug santonin is no longer used (Reynolds and Prasad, 1982; Tyler et al., 2001). Fortunately, A. herba-alba contains very few amount of santonin, 0.99% w/w (Khafagy, 1971). Like our result that indicated safety of ACEA, it (85 mg/kg) induces hypoglycemic effect and doses did not cause any acute toxicity or behavioral changes in rabbits (Iriadam et al., 2006) and the crude C. crista powder appears to be potent and safer than its methanol extract on the basis of the side effects observed (Javed et al., 1994). It is highly advisable that pre- and posttreatment helmintih levels be determined in flocks to monitor the levels of parasitism suffered by the birds and to ensure effective removal of parasites by herbal intervention. Efficacies below 90% are not considered therapeutic in the evaluation of anthelmintics for effectiveness (Yazwinski et al., 2003). Impacts of using medicinal plants for GI parasite control can be measured by: increased weight gain, improved FCR, decreased host mortality, reduced use of commercial anthelmintics, decreased EPG and reduced L3 larvae counts in coprocultures (Ketzis et al., 2006). In the present study, it was observed that ACEA was effective and well comparable with the standard drug, ABZ.

Conclusion

A. herba-alba induced anthelmintic effect as it reduced egg shedding and worm burden in the infected birds, in a similar manner to that of albendazole. The herbal extract produced significantly improved FCR over the other infected groups (positive control and albendazole treated group) and had no adverse effect on liver and kidney of treated poult. Taking together all these findings, we suggest that A. herba-alba could be used for controlling heterakid infection as an alternative to standard anthelmintic drugs. Attempts for the isolation and characterization of the active constituents responsible for such activities are currently under progress. Further studies are necessary to understand the exact mechanism of action.

ACKNOWLEDGMENTS

The authors thank Dr. Moustafa, Sh., Head of the Department of Pathology, Faculty of Veterinary Medicine, Benha University, Egypt and Dr. Soliman, A.S., chief researcher of pathology, Animal health research institute, for their cooperation in performing the histopathological part.

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