

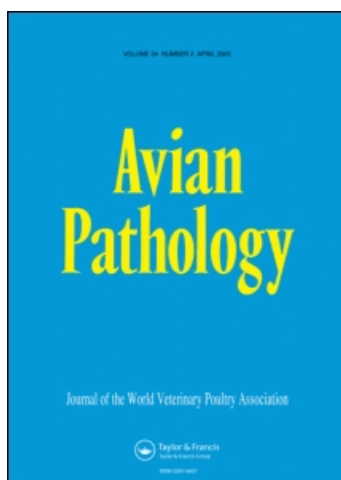
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Nele Bleyen <sup>a</sup>; Ellen Ons <sup>a</sup>; Maarten De Gussem <sup>b</sup>; Bruno M. Goddeeris <sup>a</sup>

<sup>a</sup> Division Gene Technology, Department Biosystems, Faculty of Bioscience Engineering, K.U. Leuven, Leuven, Belgium <sup>b</sup> Alpharma Animal Health Inc., Antwerpen, Belgium

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# Passive immunization against *Histomonas meleagridis* does not protect turkeys from an experimental infection

Nele Bleyen<sup>1\*</sup>, Ellen Ons<sup>1</sup>, Maarten De Gussem<sup>2</sup> and Bruno M. Goddeeris<sup>1</sup>

<sup>1</sup>Division Gene Technology, Department Biosystems, Faculty of Bioscience Engineering, K.U. Leuven, Kasteelpark Arenberg 30, 3001 Leuven, Belgium, and <sup>2</sup>Alpharma Animal Health Inc., Laarstraat 16, B-2610 Antwerpen, Belgium

Histomonosis or blackhead is a disease of gallinaceous birds, caused by the protozoan *Histomonas meleagridis*. As recent regulatory action has removed almost all drugs against this disease from the European market, the development of new prophylactics has become crucial. Identification of the protective immune mechanism would facilitate the choice and development of a vaccination strategy to prevent histomonosis. In this study, turkeys were either actively or passively immunized and were then challenged to assess the role of antibody-mediated immunity in the protection from this disease. Active immunization was performed either by experimental infection and treatment or by intramuscular injection with lysed *H. meleagridis*. Passive immunization was attempted by intraperitoneal administration of pooled, concentrated, neutralizing antisera from immunized donor animals to naïve turkeys. A significantly higher IgG response was observed after infection and treatment than after intramuscular injection, which in turn was higher than the responses of placebo and control birds. While active immunization of turkeys by intramuscular injection of dead *H. meleagridis* antigens appeared not to be protective against histomonosis, immunization by infection and treatment did induce protection. However, no significant level of protection could be observed in the passively immunized birds. These results suggest that serum antibodies to *H. meleagridis* may not be a key component in the protection against this parasite. It is, however, possible that the concentration of antibodies at the mucosal site is insufficient. Therefore, further investigation on mucosal immune responses is necessary.

## Introduction

Histomonosis or blackhead—a disease of gallinaceous birds, especially of turkeys—is caused by the protozoan parasite *Histomonas meleagridis*. The disease is characterized by necrotic foci in the liver, thickening and ulceration of the caecal wall and sulphur-coloured droppings. Mortality and morbidity may reach 100% in turkeys, while the disease is much less severe in chickens (McDougald, 1991). Due to recent regulatory action, a registered product for the prevention or treatment of blackhead is no longer available in the European Union. This has encouraged the search for alternative products, such as a vaccine against *H. meleagridis*.

In the past, it has been established that the parasite causes higher mortality in young turkeys than in adults, demonstrating the possibility of immunity against *H. meleagridis* in the older birds (Tyzzer, 1934). However, immunization by intramuscular or intravenous injections with diseased tissue emulsions failed to induce protection against histomonosis (Higgins, 1915; Tyzzer *et al.*, 1921). Recently, intramuscular vaccination with inactivated, cloned *H. meleagridis* was also unable to cause protection (Hess *et al.*, 2008). In contrast, attempts to vaccinate turkeys with *in vitro* attenuated *H. meleagridis* have led to partial or even complete protection (Tyzzer,

1933, 1934, 1936; Lund *et al.*, 1966; Hess *et al.*, 2008), although the effector mechanisms of this protective immunity remain unclear. In 1963, Clarkson investigated the protective value of antibodies against this parasite. He passively immunized naïve birds with antisera collected from infected and treated birds. His results suggested that antibodies do not protect birds from histomonosis, although only a semi-quantitative analysis of the immune response was conducted and specific antibody titres were not measured (Clarkson, 1963). Hence, the objective of the present study was to determine the strength and protective value of an antibody response against *H. meleagridis*.

## Materials and Methods

**Turkeys and parasites.** All animal experiments were approved by the Ethical Committee for Animal Experiments of the K.U. Leuven, according to the international regulations (project number P05098). Commercial turkeys (breed BIG6) were obtained as 1-day-old poults and housed on litter in a room with high efficiency particulate arresting (HEPA)-filtered air. They had free access to food and water.

To immunize and challenge the turkeys, *H. meleagridis* strain mdc (Bleyen *et al.*, 2007) was cultured as described by McDougald & Galloway (1973). Microscopic analysis and a diagnostic polymerase

\*To whom correspondence should be addressed. Tel: +32 16 32 14 33. Fax: +32 16 32 19 94. E-mail: Nele.Bleyen@biw.kuleuven.be

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chain reaction (Bleyen *et al.*, 2007) were performed to confirm the presence of *H. meleagridis*.

**Experiment 1: analysis of protection after active immunization.** Thirty-four turkeys were randomly divided into four groups. Two groups acted as controls: an uninfected, non-immunized control (UNC, four birds) and an infected, non-immunized control (INC, 10 birds). Two groups of 10 turkeys were immunized against *H. meleagridis* either by infection and treatment (IC) or by intramuscular immunization with *H. meleagridis* antigens (IM).

At 21 days of age (0 days post immunization [d.p.i.]), group IC was infected via the cloaca with  $2 \times 10^5$  *H. meleagridis* protozoa, as described by McDougald & Fuller (2005). Six days later, as clinical signs were showing, the birds were treated with ronidazole 10 mg/kg (Tricho Plus, Oropharma, Maastricht, The Netherlands). Also at 0 d.p.i., group IM was immunized intramuscularly with *H. meleagridis* antigens. The antigens were prepared by homogenizing infected caeca and foci of the liver in 0.9% NaCl solution ("physiological saline") and lysing the cells ( $2 \times 10^5$  *H. meleagridis*/ml) by sonication. Protein concentrations were determined with a protein assay using bicinchoinic acid for the colourimetric detection and quantitation. (Pierce, Rockford, Illinois, USA). Each bird received 250 µg of protein, emulsified in Freund's incomplete adjuvant (Sigma, Steinheim, Germany). Three weeks later (20 d.p.i.), a booster immunization was administered to group IM in the same manner. Groups UNC and INC were not treated during this period of the experiment.

At 27 d.p.i., groups INC, IC and IM and were challenged via the cloaca with  $10^5$  *H. meleagridis* organisms. Ten days later (10 days post challenge), all animals were killed by decapitation for postmortem examination. Macroscopic liver lesions were scored as follows: 0 = normal; 1 = a few small foci (off-white and variable in appearance) visible on the surface of the liver; 2 = lesions covering one-half of the liver surface; 3 = necrotic lesions (often large) covering more than 50% of the liver surface; and 4 = death from histomonosis, with coalescing, necrotic lesions all over the liver surface. Lesions in the caeca were scored as follows: 0 = normal; 1 = a few scattered small lesions, but little or no thickening of the mucosal wall; 2 = yellow and foamy contents, lesions prominent but discreet and some bleeding and thickening of the mucosa visible; 3 = enlarged caeca, empty or filled with yellow caseous abnormal lumen contents, thickened walls and confluent lesions, entire caecum involved; and 4 = death from histomonosis, distended caeca with fragile walls and necrotic lesions.

**Experiment 2: specific antibody response to active immunization.** For the generation of immune serum, 49 turkeys were divided randomly into five groups. At 24 days of age, two groups—group HIC (10 birds) and—group HIM (10 birds)—received the same treatment as the IC and IM groups, respectively (see Experiment 1). A placebo group was inoculated via the cloaca with homogenized caeca and liver from an uninfected turkey (PIC, 10 turkeys), and another group was immunized intramuscularly (PIM, nine turkeys) with the same mixture. At 20 d.p.i., a booster immunization was administered to all groups in the same manner. The control group (CON, 10 turkeys) was neither vaccinated nor challenged. Blood samples were taken at 0, 6, 13, 20 and 27 d.p.i. (at necropsy). Serum was collected and stored at  $-20^\circ\text{C}$  for indirect immunofluorescence assays (IFA). Seven days after the booster was given (27 dpi), the birds were killed by decapitation, all blood was collected and necropsy was performed. Antiserum collected from each group (HIC, HIM, PIC and PIM) was pooled and used for antibody transfer to naïve turkeys (Experiment 3, see below).

To determine serum IgG titres by IFA, 25 µl of a 2-day-old culture of *H. meleagridis* ( $5 \times 10^5$ /ml) fixed in 4% paraformaldehyde were seeded onto the 6 mm wells of a Teflon-coated slide (Immuno-Cell, Mechelen, Belgium). They were air-dried and treated with blocking buffer (5% horse serum, 0.02% sodium azide in phosphate-buffered saline) for 30 min at  $37^\circ\text{C}$ . The slides were then incubated with two-fold dilutions of the test sera (in blocking buffer) for 30 min at  $37^\circ\text{C}$ . Finally, the slides were incubated with goat fluorescein isothiocyanate-labelled anti-turkey (IgG) antibodies (1/50 diluted in blocking buffer; KPL, Maryland, USA) for 30 min at  $37^\circ\text{C}$ . Fluorescence staining was examined microscopically at  $500\times$  magnification.

**Experiment 3: analysis of protection after passive immunization.** Antibodies from the pooled sera from Experiment 2 were precipitated in ammonium sulphate (40% wt./vol.) and the pellets were resuspended in 1/10 volume physiological saline. The antibody mixtures were subsequently dialysed extensively against physiological saline. Antibody titres were determined by IFA as described above.

Before antibody transfer, the pooled and concentrated antisera were evaluated for their ability to induce complement-mediated lysis of *H. meleagridis*. A 2-day-old culture was diluted in culture medium to a concentration of approximately  $1.5 \times 10^5$  *H. meleagridis*/ml. Serum collected from a *H. meleagridis*-negative turkey (confirmed by IFA) was used as the source of complement. Pooled antisera from the HIC and the HIM groups were tested, while that from the CON group was used as a control. Prior to use, these test samples were inactivated at  $56^\circ\text{C}$  for 30 min. The assay was performed in a sterile 96-well Cellstar culture plate (Greiner Bio-One, Wemmel, Belgium).

Each test reaction contained 50 µl *H. meleagridis* culture, 50 µl tested antisera (undiluted, and 1/10, 1/100 and 1/1000 diluted) and 50 µl complement. Control reactions included parasites plus complement (PC) and parasites plus physiological saline (PP). The concentration of parasites in each reaction mixture was determined, using a Neubauer counting chamber, at the beginning of the experiment, after incubation at  $37^\circ\text{C}$  for 1 h, for 3 h and after overnight incubation. The percentage reduction in live parasites was calculated as follows:

$$\% \text{ reduction} = \left[ \frac{\text{concentration}_{\text{start}} - \text{concentration}_{x \text{ hours after incubation}}}{\text{concentration}_{\text{start}}} \right] \times 100.$$

For the passive immunization, 35 turkeys were divided in six groups (see later Table 3). At 5 weeks of age, the birds were given an intraperitoneal injection of 4 ml concentrated antisera. The passively immunized groups, PASHIC and PASHIM, received antibodies from the HIC and HIM groups, respectively; while the placebo groups PASPIC and PASPIM received concentrated PIC and PIM antibodies, respectively. Groups POS and NEG were not immunized. Four hours after serum transfer, the birds in all groups except for the NEG group were challenged via the cloaca with  $3 \times 10^5$  *H. meleagridis*. Ten days later (10 days post transfer), all of the turkeys were killed by decapitation for postmortem examination of livers and caeca as described above. To track the antibody titres before and after passive immunization, blood was taken 1 day before ( $-1$  day post transfer), 1 day after serum transfer and at 10 days post transfer. Sera were collected and stored at  $-20^\circ\text{C}$  until analysis by IFA as described above.

**Statistical analysis.** All statistical analyses were performed using the SAS 8.2 software (SAS Institute, Cary, North Carolina, USA). Lesion scores, antibody titres in the sera at each time point and mortality rates were compared using Kruskal–Wallis analysis to detect significant differences between multiple groups, and the Wilcoxon rank sum test was used to compare differences between each pair of groups.  $P \leq 0.05$  was considered to be significant.

## Results

**Experiment 1: analysis of protection after active immunization.** During immunization by infection and treatment, two birds of the IC group died of histomonosis (at 8 d.p.i. and 21 d.p.i.). After challenge, none of the immunized birds of the IC group and only one bird of the intramuscularly immunized IM group died (8 days post challenge). Six birds of the infected, non-immunized control INC group died of histomonosis, three at 9 days post challenge and three at 10 days post challenge. There were no significant differences between the mortality rates of the uninfected UNC group and those of the immunized and challenged IC and IM groups, while the mortality of the challenged but non-immunized INC group differed significantly from all other groups (Table 1).

In the liver, significant differences were seen between the mean lesion scores of the non-immunized INC group

**Table 1.** Experiment 1: mortality and mean lesion scores of groups of turkeys after active immunization and challenge with *H. meleagridis*

Group <sup>a</sup>	Total <sup>b</sup>	Immunization	Challenge <sup>c</sup>	Mortality (%) <sup>d</sup>	Mean lesion scores <sup>e</sup>	
					Caeca	Liver
UNC	4	–	–	0 <sup>A</sup>	0.0 <sup>A</sup>	0.0 <sup>A</sup>
INC	10	–	+	60 <sup>B</sup>	2.8 <sup>B</sup>	2.9 <sup>B</sup>
IC	8	Infection/treatment	+	0 <sup>A</sup>	1.1 <sup>C</sup>	0.1 <sup>A</sup>
IM	10	intramuscular	+	10 <sup>A</sup>	2.2 <sup>B</sup>	2.2 <sup>B</sup>

Different uppercase superscript letters (within a column) indicate a statistical difference ( $P \leq 0.05$ ). <sup>a</sup>UNC, uninfected, non-immunized control; INC, infected, non-immunized control; IC, immunized by infection and treatment; IM, immunized by intramuscular injection with *H. meleagridis* antigens. <sup>b</sup>Number of turkeys in each group on the day of challenge. <sup>c</sup>Each group, except for the UNC group, was challenged with  $10^5$  *H. meleagridis*. <sup>d</sup>Mortality by 10 days post challenge: three INC birds died 9 days post challenge, three died 10 days post challenge; one IM bird died 8 days post challenge. <sup>e</sup>Mean lesion scores on the day of autopsy. Birds that died of histomonositis were given a lesion score of 4 per organ.

and the IC and UNC groups and between the IM group and the IC and UNC groups. No significant differences were observed between mean liver lesion scores of the INC and the IM group, or between the IC and the uninfected control UNC group. The mean caecal lesion score of the IC group was significantly lower than those of the INC and IM groups, but significantly higher than that of the UNC group. There were no statistical differences between the mean caecal lesion scores of the IM and INC groups, which were both significantly higher than those of the UNC and the IC groups. The mean caecal lesion scores of the surviving birds of the INC and the IM groups were however 1 and 2, respectively, while their mean liver lesions were 1.25 and 2, respectively (Table 1).

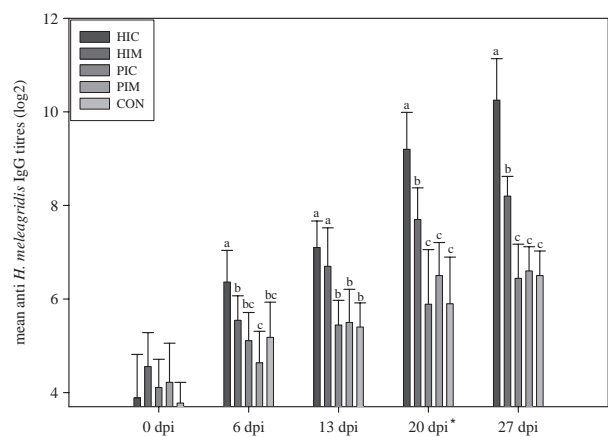
**Experiment 2: specific antibody response to active immunization.** Analysis of the serum IgG response against *H. meleagridis* using IFA with fixed *H. meleagridis* as antigen showed that the parasites fluoresced intensely. Without primary incubation with antisera, no fluorescence could be detected, indicating the absence of autofluorescence or non-specific fluorescence.

The mean anti-*H. meleagridis* antibody titres of the immunized groups (HIC and HIM) were significantly higher than those of the placebo (PIC and PIM) and control (CON) groups, from 6 d.p.i. (HIC) and from 13 d.p.i. (HIM) onwards (Figure 1). From 6 d.p.i., significantly higher anti-*H. meleagridis* antibody titres were found in group HIC compared with group HIM—except for 13 d.p.i., when no significant differences could be detected between them. Throughout the entire experiment, no significant differences were observed between the placebo and the control groups.

**Experiment 3: analysis of protection after passive immunization.** Before antibody transfer, the pooled, concentrated antisera were evaluated for their bioactivity by examining their ability to induce complement-mediated lysis of *H. meleagridis*. The results of this test are presented in Table 2. Incubation of *H. meleagridis* with complement and the undiluted HIC or HIM concentrated serum pools resulted in a reduction in living parasites after 1 h. Moreover, after 3 h and after overnight incubation, growth of *H. meleagridis* was impeded by 100% in both samples. This reduction was detectable at serum dilutions of 1/10 and 1/100 but not at 1/1000. When incubating the parasites and the comple-

ment with the pooled CON sera or without test sera, no reduction and even growth (indicated in Table 2 by the negative values) could be observed.

The anti-*H. meleagridis* antibody titres as determined by IFA are shown in Figure 2. One day before passive serum transfer (–1 day post transfer), no statistical differences in antibody titres could be detected. One day after transfer (1 day post transfer), the titres of the passively immunized groups (PASHIC and PASHIM) increased respectively 9.8-fold and 7.5-fold compared with the titres at –1 day post transfer, and were significantly different from the titres of the other groups. The birds were challenged via the cloaca 4 h after passive serum transfer. On the day of necropsy (10 days post transfer), the titres of groups PASHIC and PASHIM were still significantly different from those from groups PASPIC, PASHIM and POS. The antibody titres from all challenged groups were significantly different from group NEG at 10 days post transfer.



**Figure 1.** Experiment 2: serum antibody responses of groups of turkeys after active immunization with *H. meleagridis*. Turkeys immunized at 0 d.p.i., with a booster at 20 d.p.i. (indicated by the asterisk). HIC, immunized by intraoocel infection with *H. meleagridis*; HIM, immunized intramuscularly with *H. meleagridis* antigens; PIC, immunized by intraoocel injection with *H. meleagridis*-negative liver and caecal homogenate; PIM, immunized as PIC but intramuscularly; CON, non-immunized control. Titres measured by indirect immunofluorescence. Different lowercase letters indicate significant differences between treatments at each time point.

**Table 2.** Lytic activity of anti-*H. meleagridis* antibodies after different incubation times

Sample <sup>b</sup>	Reduction in growth of <i>H. meleagridis</i> by serum antibodies after incubation (%) <sup>a</sup>		
	1 h	3 h	Overnight
<b>HIC</b>			
Undiluted	88	100	100
1/10	75	50	50
1/100	75	0	-50
1/1000	33	-67	-67
<b>HIM</b>			
Undiluted	67	100	100
1/10	50	38	-25
1/100	50	50	50
1/1000	-50	-50	-100
<b>CON</b>			
Undiluted	0	0	-100
1/10	-43	-43	-186
1/100	-43	-43	-114
1/1000	0	0	-150
<b>Controls</b>			
PC <sup>c</sup>	-71	-71	-186
PP <sup>d</sup>	-40	-100	-200

<sup>a</sup>% reduction = [(concentration<sub>start</sub> - concentration<sub>x hours after incubation</sub>)/concentration<sub>start</sub>] × 100. A negative value indicates growth, a positive value indicates reduction. <sup>b</sup>Reaction containing parasites, complement and (un)diluted test sera from HIC, HIM or CON groups. HIC, immunized by intracloacal infection with *H. meleagridis*; HIM, immunized intramuscularly with *H. meleagridis* antigens; CON, non-immunized control group. <sup>c</sup>Control reaction containing parasites and complement. <sup>d</sup>Control reaction containing only parasites.

All challenged groups showed lesions in the examined organs (Table 3), but no significant differences between the mean lesion scores of the challenged groups could be found. For the unchallenged NEG group, no lesions could be detected in the liver or caeca, resulting in a significant difference in mean lesion scores between this group and the challenged groups.

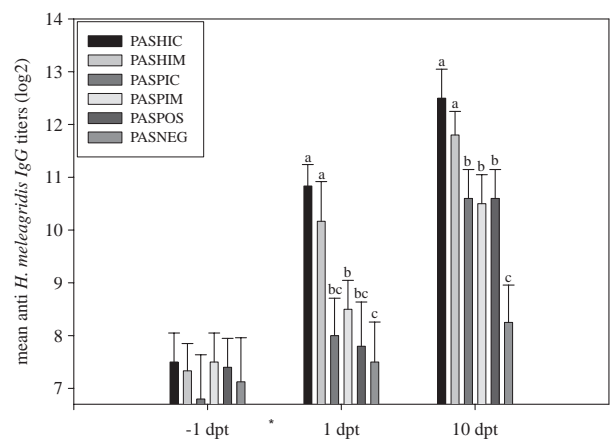
## Discussion

In the present study, an anti-*H. meleagridis* serum antibody response was detected after immunization by infection and treatment and after intramuscular immunization with *H. meleagridis* antigens. Immunity induced by intramuscular injection of dead *H. meleagridis* antigens appeared not to be protective against histomonos. In contrast, turkeys actively immunized by infection followed by treatment were protected against a *H. meleagridis* challenge. However, transfer of anti-*H. meleagridis* antibodies to naïve turkeys did not protect them against challenge via the cloaca with  $3 \times 10^5$  *H. meleagridis* organisms.

Several earlier studies encouraged us to investigate the protective value of the serum antibodies against *H. meleagridis*. Tyzzer (1934) suggested that protective immunity could be age-related, as older birds are more likely to be immune to this disease. Furthermore, a degree of protection against intracloacal challenge with *H. meleagridis* and against an oral challenge with *H. meleagridis*-contaminated *Heterakis gallinarum* eggs was detected after young turkeys were vaccinated with attenuated cultures of *H. meleagridis* (Tyzzer, 1933, 1934, 1936; Lund *et al.*, 1966). Recently, cloned attenuated *H. meleagridis* has been shown to induce protection against an intracloacal challenge (Hess *et al.*, 2008). However, examination of the immune response was not

included in these studies so it was not possible to link this protection to a systemic or mucosal immune response.

In 1963 Clarkson performed a passive immunization experiment to determine whether antibodies could protect turkeys against *H. meleagridis*, and suggested that they could not or could not completely induce protection. However, he used only a semi-quantitative analysis of the immune response (Clarkson, 1963),



**Figure 2.** Experiment 3: serum antibody responses of groups of turkeys after passive immunization and challenge with *H. meleagridis*. \*Transfer of immune serum and challenge (4 h later) with *H. meleagridis* at 0 days post transfer (dpt). PASHIC, passively immunized with HIC antibodies; PASHIM, passively immunized with HIM antibodies; PASPIC, received antibodies from PIC; PASPIM, received antibodies from PIM; POS, non-immunized challenged group; NEG, non-immunized non-challenged group. Titres measured by indirect immunofluorescence. Different lowercase letters indicate significant differences between treatments at each time point.

**Table 3.** Experiment 3: passive immunization and challenge against *H. meleagridis*

Group	Total <sup>a</sup>	Antibodies <sup>b</sup>	Challenge <sup>c</sup>	Deaths <sup>d</sup>	Mean lesion scores <sup>e</sup>	
					Caeca	Liver
PASHIC	5	HIC	+	0	3.0 <sup>A</sup>	2.8 <sup>A</sup>
PASHIM	6	HIM	+	1	3.0 <sup>A</sup>	2.3 <sup>A</sup>
PASPIC	5	PIC	+	0	2.7 <sup>A</sup>	2.2 <sup>A</sup>
PASPIM	6	PIM	+	0	2.8 <sup>A</sup>	2.3 <sup>A</sup>
POS	5	/	+	0	3.0 <sup>A</sup>	2.5 <sup>A</sup>
NEG	8	/	–	0	0.0 <sup>B</sup>	0.0 <sup>B</sup>

Different uppercase superscript letters (within a column) indicate a statistical difference ( $P \leq 0.05$ ). <sup>a</sup>Number of turkeys in each group. <sup>b</sup>Group from Experiment 2 providing the antibodies. <sup>c</sup>Each group, except for NEG, was challenged with  $3 \times 10^5$  *H. meleagridis*.

whereas in the present study actively and passively immunized and challenged turkeys were examined on a quantitative basis, to determine the protective value of antibodies against this parasite.

In Experiment 1, the degree of protection was analysed after actively immunizing turkeys. Following challenge of turkeys immunized by infection and treatment (IC group), the caeca of the birds showed signs of infection, although the parasites did not appear to migrate to the liver or cause death. This clearly showed that turkeys are protected against *H. meleagridis* after infection and treatment. A second group of birds (group IM) was immunized by intramuscular injection with dead *H. meleagridis* antigens derived from infected livers and caeca. This choice of immunizing material should generate antibodies against all *in vivo* stages as the parasite appears in different stages within its host (Lee *et al.*, 1969; Honigberg & Bennett, 1971) but it resulted in significantly higher mean lesion scores of both the liver and caeca than those of the IC and the negative control group—although, as far as we could judge the mortality at 10 days post challenge, only one of 10 birds died of histomonosis. Furthermore, the mean lesion scores of the surviving birds were higher in the IM group than in the INC (infected non-immunized control) group, suggesting that immunity induced by dead antigens is not protective against histomonosis. These findings are supported by the results of Hess *et al.* (2008).

In a second active immunization experiment (Experiment 2) a strong anti-*H. meleagridis* serum antibody response was produced after experimental infection and recovery, while intramuscular immunization resulted in a slightly lower, but still significant, antibody response. The booster immunization resulted in both groups in a subsequent significant increase in parasite-specific serum antibodies.

In the passive immunization Experiment 3, the sera from the second experiment were concentrated and transferred to naïve animals. After transfer, the titres of the anti-*H. meleagridis* immunized groups (PASHIC and PASHIM) increased respectively 9.8-fold and 7.5-fold, resulting in higher mean antibody titres than those of the originating groups (HIC and HIM). However, despite the successful transfer of anti-*H. meleagridis* antibodies, no protective effect was observed, at least not against an intraoocyst challenge with  $3 \times 10^5$  parasites.

It is possible that the antibody titres of the concentrated pooled sera for passive immunization were not high enough to result in protection. This is considered

unlikely because group IC of Experiment 1 was protected after being immunized in the same manner as the HIC group (Experiment 2), although these birds had lower antibody titres at the day of challenge than the titres after passive immunization. *In vitro* analysis of the bioactivity of the specific antisera showed that the antibodies transferred to the naïve animals could still activate the lysis of *H. meleagridis* but this activity decreased at a dilution of 1/100 and ceased at 1/1000, which could indicate that the anti-*H. meleagridis* activity of the antibodies was not strong enough to induce protection.

Another consideration is that the serum antibody titres of passively immunized groups might no longer have been high enough to be protective when the infection reached its systemic phase. Indeed, the transferred antibodies peaked in the serum approximately 1 day after injection (preliminary experiment; data not shown) and declined thereafter. However, the titres of the passively immunized groups remained readily detectable and significantly higher than those of the placebo groups. Thus, a certain level of protection should have been seen.

Lastly, and most importantly, the challenge dose might have been too high to simulate a natural challenge. A lower challenge dose might have generated differences in mean lesion scores, thus revealing the protective value of the transferred antibodies.

Despite these explanations, the most logical and direct conclusion of our study is that systemic immunity by serum antibodies against *H. meleagridis* is not, or is not primarily, responsible for protection against histomonosis. Instead, mucosal immune responses might be more important in the prevention of this disease and deserve further investigation.

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