Intestinal Antilectin Immunoglobulin A Antibody Response and Immunity to *Entamoeba dispar* Infection following Cure of Amebic Liver Abscess

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We followed 93 subjects with amebic liver abscess (ALA) and 963 close associate controls at 3-month intervals for 36 months to characterize intestinal and humoral antibody responses to the amebic galactose-inhibitable lectin and to determine whether immunity developed to *Entamoeba histolytica* or *Entamoeba dispar* infection following cure of ALA. We found that ALA subjects had a higher prevalence and level of intestinal antilectin immunoglobulin A (IgA) and serum anti-LC3 (cysteine-rich recombinant lectin protein) IgA and IgG antibodies, $P<0.01$ and $P<0.05$, respectively, compared to controls. The intestinal antilectin IgA antibody response was sustained over a longer time period in ALA subjects (71.8% remained positive at 18 months and 52.6% at 36 months, $P<0.001$ compared to 17.6% and 10.3% of controls, respectively). ALA subjects were highly immune to *E. dispar* infection throughout the study (0% infected at 6 and 36 months, compared to 6.5% and 4.9% of control subjects, respectively, $P<0.05$). Upon entry into the study, 6.3% of ALA subjects were infected with *E. histolytica*; the incidence of new *E. histolytica* infections in controls (as determined by culture) was too low (1.4%) to determine whether ALA subjects exhibited immunity to new infections. We found that stool cultures every 3 months markedly underestimated the occurrence of new *E. histolytica* infections, as 15.3% of controls seroconverted after 12 months of follow-up. Unfortunately, under the field conditions present in Durban, South Africa, enzyme-linked immunosorbent assay for detection of lectin antigen in stool yielded unreliable results. In summary, subjects cured of ALA exhibited sustained mucosal IgA antibody responses to the amebic galactose-inhibitable lectin and a high level of immunity to *E. dispar* infection. Determination of immunity to *E. histolytica* following cure of ALA will require the use of more sensitive and reliable diagnostic methods.

One of the major questions in amebiasis research is whether cure of invasive disease is followed by development of immunity to new intestinal infections and, thus, recurrence of disease. The enteric protozoan *Entamoeba histolytica* is one of the leading parasitic causes of death worldwide. Disease results from the parasite's ability to invade the colon, causing amebic colitis, or spreading via the portal venous system to the liver, resulting in formation of an amebic liver abscess (ALA). Amebic liver abscesses are more common in adult men and were thought to be fatal if untreated (7). A recent study in Hue, Vietnam, revealed that ALA is even more common than previously realized and may occur frequently in a subclinical manner (10). One large noncontrolled study reported that the rate of recurrence of amebic liver abscesses over 5 years in a high-risk population was less than expected compared to historical controls (14). In a cross-sectional study, the point prevalence of *Entamoeba* species intestinal infection was lower in subjects who possessed serum antiamebic antibodies (13).

The *E. histolytica* galactose-inhibitable lectin (12, 22, 26, 27) appears to have a crucial role in colonization of the gut and parasite invasion. The lectin mediates attachment of *E. histolytica* trophozoites to colonic mucins (11, 12), host epithelial cells and immune effector cells (22, 30). Galactose-inhibitable lectin binding is an absolute requirement for trophozoites to exhibit a lytic effect on host cells (25). The purified lectin in native and recombinant forms is a highly conserved antigen. In over 95% of samples obtained from hundreds of patients cured of amebic colitis or liver abscess studied worldwide, native lectin protein purified from a single cloned *E. histolytica* isolate is recognized by serum immunoglobulin G (IgG), IgM, and IgA antibodies (1, 3, 5, 6, 21, 32). The same has been found from subjects with noninvasive asymptomatic *E. histolytica* intestinal infection (28, 31).

Monoclonal antibodies raised to the lectin's carbohydrate-binding domain completely inhibit parasite binding to colonic mucins in vitro (11, 12), suggesting that intestinal antilectin IgA antibodies could prevent parasite colonization of the gut. In a prospective follow-up study of children in Bangladesh, there was a delay in the onset of *E. histolytica* intestinal infections when intestinal antilectin IgA antibodies were present (17). The lectin in native and recombinant form has been demonstrated to be efficacious as a subunit vaccine in the gerbil model of amebic liver abscess (24, 32).

In Durban, South Africa, *E. histolytica* and *Entamoeba dispar* infections are highly endemic (16, 20). *E. dispar* is a distinct species that is morphologically identical to *E. histolytica* but is not known to cause disease (15). *E. dispar* trophozoites possess...
functional galactose-binding lectin molecules that are 85% homologous with the *E. histolytica* lectin (25) and have many common epitopes as determined by studies with murine monoclonal antibodies raised to the *E. histolytica* lectin (23). The purpose of our study was to characterize over time the human mucosal and humoral antilectin antibody responses and to determine whether intestinal immunity to infection exists following cure of invasive amebiasis. These findings provide information that is crucial for the development of an effective lectin-based amebiasis subunit vaccine.

We conducted a prospective cohort study of 93 subjects treated for ALA and 963 controls who were family members or closely associated neighbors. All subjects were enrolled prospectively and followed for at least 36 months. The demographics, risk factors for infection by *Entamoeba* species, and prevalence of infection with other intestinal parasites will be reported elsewhere.

**MATERIALS AND METHODS**

**Subject recruitment and study enrollment.** Subjects with ALA were recruited at King Edward VIII Hospital and other regional hospitals and clinics in the area around Durban, South Africa. Nurses fluent in Zulu obtained informed consent in English or Zulu. Control subjects included nuclear family members, individuals residing in the same household, and close neighbors living in the same environment. Control subjects were recruited by study nurses through contact with the index case; we sought at least 10 close associates controls for each index case. No criteria for age or gender were applied except age ≥16 years. All subjects provided blood by venipuncture, feces, and throat washings at entry into the study (1 week after commencing treatment of the ALA index case) and at 3-month intervals for a total of at least 36 months of follow-up. Over the duration of the study, only seven of the 100 family groups were lost to follow-up. At the first visit, study nurses filled out a detailed epidemiologic questionnaire based on an oral history obtained from each of the subjects enrolled in the study. The University of Minnesota and University of Natal’s institutional review boards for human subjects approved the consent form, questionnaire, and all aspects of the study.

Assays performed included fecal microscopy, stool culture, and zymodeme determination for *E. histolytica* or *E. dispar*, enzyme-linked immunosorbent assay (ELISA) for serum anti-LC3 (recombinant cysteine-rich section of the lectin heavy subunit) (32) IgA and IgG antibodies, ELISA for fecal antilectin and anti-LC3 IgA antibodies, and a monoclonal antibody-based antigen capture ELISA to detect *E. histolytica* and *E. dispers*-specific lectin antigen in feces (1, 2, 3, 5, 6).

**Stool culture.** Fecal samples were cultured in Robinson’s medium (29) for detection of *E. histolytica* and *E. dispers* parasites. Primary cultures were performed by adding a small piece of fecal material to a Biotex container containing an agar slope, to which was added 10 mg of starch, 4 drops of 20% erythromycin, and 100 ml of lactic acid, diluted to 2.5 liters with distilled water. For use, 100 ml of stock was diluted with 7.5 ml of 40% NaOH and 2.5% of bromothymol blue, adjusted to 1 liter with distilled water at pH 7.0, and autoclaved. After 24 h, the supernatant was removed, leaving the starch-fecal layer. The supernatant was replaced about 2/3 of the way up the slope with BRS medium (equal volume of BR and sheep serum incubated for 24 h at 37°C) diluted 1:4 with phthalate solution (10.2% potassium phthalate, 2% NaOH, pH 6.3). After 48 h of incubation at 37°C, a drop from the starch layer was mixed with phthalate solution (10.2% potassium phthalate, 2% NaOH, pH 6.3). After 24 h, the supernatant was removed, leaving the starch-fecal layer. The supernatant was replaced about 2/3 of the way up the slope with BRS medium (equal volume of BR and sheep serum incubated for 24 h at 37°C) diluted 1:4 with phthalate solution (10.2% potassium phthalate, 2% NaOH, pH 6.3). After 48 h of incubation at 37°C, a drop from the starch layer was mixed with phthalate solution (10.2% potassium phthalate, 2% NaOH, pH 6.3). After 48 h of incubation at 37°C, a drop from the starch layer was mixed with phthalate solution (10.2% potassium phthalate, 2% NaOH, pH 6.3).

**Detection of serum anti-LC3 IgG and IgA antibodies by ELISA.** Detection of serum anti-LC3 IgG and IgA antibodies by ELISA was performed as described previously (17, 26, 32). Recombinant 52-kDa LC3 protein was purified as described by Song et al. (32); 96-well microtiter flat-bottomed polystyrene ELISA plates were coated with LC3 protein, and nonreactive sites were blocked with 1% bovine serum albumin. Serum samples were analyzed by ELISA at a 1:1,000 dilution for IgG and a 1:500 for IgA antibodies in phosphate-buffered saline-Tween (1% bovine serum albumin). Following incubation for 2 h at room temperature, alkaline phosphatase-conjugated goat anti-human IgG (Sigma) or IgA antibodies (ICN Biomedicals; Costa Mesa, Calif.) were utilized at 1:5,000 for IgG and 1:2,000 for IgA in phosphate-buffered saline-Tween (1% bovine serum albumin) for 2 h at room temperature. Reading the plates and correcting the results for nonspecific background binding were performed as described (28).

**Detection of fecal lectin antigen by ELISA.** The ELISA for detection of 170-kDa lectin antigen was performed as described (3). Briefly, 96-well flat-bottomed microtiter polystyrene ELISA plates (Costar, Corning, N.Y.) were coated with monoclonal antibody 3F4, which recognizes epitopes present in both *E. histolytica* and *E. dispers* lectin, or the 8C12 antibody, which is specific for epitopes present only in *E. histolytica* lectin (23). Feces were mixed in an equal volume of phosphate-buffered saline containing 2 mM phenylmethylsulfonyl fluoride (USB, Cleveland, Ohio). Fecal samples were added at 100 µl per well and incubated for 2 h at room temperature or overnight at 4°C. Alkaline phosphatase-conjugated antilectin monoclonal antibodies 8A3 (recognizing both *E. dispar* and *E. dispers*) and E. dispar lectin (ICN Biomedicals) were added at a 1:5,000 dilution and incubating in developing buffer for 2 h at room temperature. Plate reading with correction for nonspecific background was performed as described (28). Detection of fecal antilectin and anti-LC3 IgA antibodies by ELISA. Native E. histolytica galactose-inhibitable lectin protein (22) and recombinant LC3 protein (32) were purified as described and used in ELISA for detection of fecal antilectin and anti-LC3 antibodies (7). Briefly, flat-bottomed microtiter plates were coated with lectin protein (0.125 µg/well) and the nonreactive sites were blocked with 1% bovine serum albumin. Fecal samples were mixed with an equal volume of phosphate-buffered saline–2 mM phenylmethylsulfonyl fluoride and added at 100 µl/well for incubation at room temperature for 2 h or overnight at 4°C. Alkaline phosphatase-conjugated goat antihuman IgA antibodies (Sigma) were added at a 1:5,000 dilution in phosphate-buffered saline-Tween containing 1% bovine serum albumin, for incubation at room temperature for 2 h. Plate reading with correction of results for nonspecific background binding was performed as described (7).

**Treatment of data.** Assays to detect humoral (anti-LC3 IgG and IgA) or mucosal (focal anti-LC3 and antilectin IgA) antibody responses utilized a continuous optical density (OD) scale, with cutoffs for positivity of two standard deviations above a culture-negative control used as a laboratory standard. A subject was considered antibody positive if he had at least two consecutive positive findings at baseline or at follow-up. Subjects were spared or nonnormally distributed) were used to evaluate differences. It should be noted that while we enrolled ALA cases and close associate controls, the study design is essentially a prospective cohort, with exposure groups defined by disease status as a proxy for prior infection with *E. histolytica* at baseline (ALA versus control). Similarly, subjects could be scored by culture positivity at base-line or follow-up as antibody positive or negative.

**Analytic methods.** For continuous data, differences of distributions between groups were evaluated with Wilcoxon rank-sum tests. Accordingly, tests results are presented with median levels of these variables, along with associated 25%-75% interquartile ranges. Contingency table analysis was used to compare proportions (yes/no) between groups: chi-square or Fisher’s exact tests (for data that were sparse or nonnormally distributed) were used to evaluate differences. It should be noted that while we enrolled ALA cases and close associate controls, the study design is essentially a prospective cohort, with exposure groups defined by disease status as a proxy for prior infection with *E. histolytica* at baseline (ALA versus control). Similarly, subjects could be scored by culture positivity at baseline or follow-up as antibody positive or negative. Results for all tests were aggregated at baseline or follow-up for all subjects for comparison between ALA and their control subjects.
RESULTS

Results from 36 months of follow-up are available for all subjects who completed the study, 93 ALA and 963 control subjects. Data are aggregated by baseline and follow-up periods. The average age of the ALA subjects was 41.5 years; the gender distribution was 83% male and 17% female. The control subjects were 25% male and 75% female, with an average age of 36.5 years.

Intestinal and serum antilectin antibody responses. The prevalence in ALA subjects and close associate controls of intestinal antilectin IgA antibodies at baseline (0 to 6 months) and follow-up (9 to 36 months) is illustrated in Fig. 1. ALA subjects had a higher prevalence of intestinal antilectin IgA antibodies at each time period studied compared to controls (85.7%, 67.2%, 56%, and 52.6% positive ELISAs versus 16.3%, 17.6%, 11.8%, and 10.3% in controls, respectively) \((P < 0.001\) for each time period; Fig. 1). Use of the purified native galactose-inhibitable \(E.\ histolytica\) lectin protein was more sensitive in ELISA for detection of antigen-specific fecal IgA antibodies than the recombinant LC3 protein \((P < 0.01; 85.7%\) compared to 55.2% positive at baseline). The prevalence of intestinal antilectin IgA antibodies decreased over time in ALA and control subjects, compared to baseline values \((P < 0.01)\) and \((P < 0.05)\), respectively. Seventy-two percent of the ALA subjects positive for intestinal antilectin IgA antibodies at baseline remained positive at 18 months, compared to only 37.5% of comparable controls \((P = 0.007)\). By 30 to 36 months, 25.9% of ALA subjects who were positive on entering the study had persistence of intestinal antilectin IgA antibodies, compared to 10.8% of controls \((P < 0.038)\).

At baseline the prevalence of serum anti-LC3 IgA antibodies was greater in ALA subjects than controls \((93.4%\) compared to 69.9% of controls, \(P < 0.001\), Fig. 2A). Serum anti-LC3 IgA antibodies remained elevated in ALA subjects compared to controls at each interval of follow-up \((85.5%, 67.2%, and 57.9%\) compared to 13.6%, 8.4%, and 6.3%, respectively, \(P < 0.001\) for each, Fig. 2A). There was an increase in serum anti-LC3 IgA antibodies in controls only at 9 to 18 months \((P < 0.001)\); in ALA cases there was a decrease in the prevalence of anti-LC3 IgG antibodies during each follow-up period \((P < 0.001)\) compared to the previous period.

FIG. 1. Prevalence of a positive ELISA for intestinal antilectin IgA antibodies in subjects cured of ALA (light bars) and controls (dark bars). The prevalence of antilectin IgA antibodies was greater in cases than controls at baseline (0 to 6 months) and during each follow-up period \((P < 0.0001,\) for each). There was a significant decrease in the prevalence of antilectin IgA antibodies between baseline and follow-up intervals in ALA cases \((P < 0.01\) for each comparison) and in close associate controls (0 to 6 and 9 to 18 months compared to 21 to 36 months, \(P < 0.05\)).

FIG. 2. Prevalence of serum anti-LC3 IgA (A) and IgG antibodies (B) for subjects cured of ALA (light bars) and controls (dark bars). The prevalence of a positive test for each antibody studied was greater in ALA subjects than controls at baseline and during each follow-up period \((P < 0.001)\) for each. The prevalence of serum anti-LC3 IgA antibodies increased in controls only at 9 to 18 months \((P < 0.001)\); in ALA cases there was a decrease in the prevalence of anti-LC3 IgG antibodies during each follow-up period \((P < 0.001)\) compared to the previous period.

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indicates that *E. dispar* infection can induce a transient intestinal but not humoral antilectin IgA antibody response.

Intestinal and serum ELISA results were analyzed as a continuous variable by comparison of OD values at equal dilutions. At baseline (0 to 6 months), ALA subjects had markedly higher ELISA OD values at the 25th, 50th, 75th, and 90th percentiles for intestinal antilectin IgA and serum anti-LC3 IgA antibodies, compared to ELISA-positive control subjects assayed at identical dilutions of feces or serum (P < 0.0001 for each percentile, Fig. 4). The same was true when ELISA OD levels for serum anti-LC3 IgG antibodies were compared between ALA subjects and ELISA-positive controls (P < 0.001, respectively, for each quartile, data not shown). During the entire 36-month follow-up period, antibody-positive ALA subjects had higher median OD values for all of the antiamebic antibodies studied, compared to ELISA-positive control subjects (P ≤ 0.04 for each, Table 1).

![Figure 3](image-url)

**FIG. 3.** Percentage of ALA (light bars) and controls (dark bars) individuals ever having a positive test over the entire 36 months of the study. ALA subjects had a higher cumulative positive percentage for each antibody studied compared to controls, P < 0.01. Of interest, the cumulative percentage of control subjects with fecal antilectin IgA antibodies was greater than for either serum anti-LC3 IgA or IgG antibodies (P < 0.005).

**TABLE 1.** Comparison of OD values between ELISA-positive ALA subjects and close associate controls during 36 months of follow-up.

<table>
<thead>
<tr>
<th>ELISA</th>
<th>Median OD</th>
</tr>
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<tbody>
<tr>
<td>Intestinal antilectin IgA</td>
<td>0.657*</td>
</tr>
<tr>
<td>Intestinal anti-LC3 IgA</td>
<td>0.123*</td>
</tr>
<tr>
<td>Serum anti-LC3 IgG</td>
<td>0.641†</td>
</tr>
<tr>
<td>Serum anti-LC3 IgA</td>
<td>0.42†</td>
</tr>
</tbody>
</table>

* †, P < 0.05 compared to close associates; †, P < 0.01 compared to close associates.

Occurrence of *E. histolytica* and *E. dispar* intestinal infections. By stool culture and zymodeme determination, 6.3% of ALA subjects were infected with *E. histolytica* during the baseline period (0 to 6 months), compared to only 1.2% of controls (P = 0.001, Table 2). The prevalence of *E. histolytica* infection over time was at 10.0%, 5.9%, and 3.1% in ALA subjects during follow-up, compared to 2.6%, 0.7%, and 1.0% of control subjects, respectively (P = 0.005, Table 2). Over the duration of the study, by culture and zymodeme determination, 15.6% of ALA subjects and 3.5% of controls were infected with *E. histolytica* (P = 0.001, Table 2). In ALA subjects there were seven *E. histolytica* infections detected after 6 months; six of seven were found at 9 to 18 months and only one was found thereafter (Table 2). The same pattern was evident in control subjects; 13 to 16 new infections were detected at 9 to 18 months and only three thereafter (Table 2). In controls, there was no correlation between being positive for intestinal antilectin IgA antibodies or serum antilectin IgA or IgG antibodies and the occurrence or nonoccurrence of new *E. histolytica* infections (P > 0.1 for each).

Studies of *E. dispar* intestinal infection revealed that at baseline (0 to 6 months), none of the 85 ALA subjects who submitted adequate fecal samples were infected with *E. dispar*, compared to 6.5% of controls (P = 0.007, Table 3). During the follow-up period, only 4 of 81 ALA subjects became transiently infected with *E. dispar* compared to 149 of 713 control subjects (4.9% compared to 20.9%, P < 0.05, Table 3). At the end of the prospective study, 30 to 36 months, none of the ALA subjects were infected with *E. dispar*, compared to 6.5% of controls (P < 0.05, Table 3). Among the controls, there was no correlation between having a positive serum or fecal antibody test at baseline and immunity against the occurrence of a new *E. dispar* infection (P = 0.80).

We determined that culture of fecal samples at 3-month intervals was not sensitive enough to detect the majority of new *E. histolytica* infections. By sero-conversion criteria, with serum anti-LC3 IgG antibodies, 15.3% of asymptomatic controls had a new *E. histolytica* infection during 12 months of follow-up (baseline compared to months 9 to 18), yet culture and zymodeme criteria detected only 1.0% as having a new infection (P > 0.01).

All fecal samples were subjected to a monoclonal antibody-based ELISA for detection of *E. histolytica* and *E. dispar*-specific lectin antigen, a research assay that we applied successfully in a number of studies encompassing hundreds of amebic infections in Cairo, Egypt (3, 6, 7). However, we found that the antigen detection ELISA results for all subjects over
the duration of the study did not correlate with stool culture and zymodeme determination (only 6 of 32 positive cultures for *E. histolytica* and 22 of 91 for *E. dispar* had true positive ELISAs, Table 4), with clinical group, or with seropositivity (data not shown). Therefore, under the field conditions and analysis as performed in this study, antigen detection ELISA technology was found to be unreliable.

**DISCUSSION**

We now report the first large prospective, controlled cohort study of *E. histolytica* and *E. dispar* intestinal infection following cure of ALA. We evaluated 93 subjects starting at 1 week following cure of ALA and 963 controls (immediate family members and/or neighbors). Subjects were evaluated at 3-month intervals thereafter with collection of feces and blood, for a total of 36 months of follow-up.

There was a high prevalence (>85%) of intestinal antilectic IgA antibody responses in ALA subjects, which unexpectedly persisted in over 50% of subjects for 18 months after treatment. The intensity of the intestinal antibody response as measured by ELISA OD readings was greater in ALA subjects than in antibody-positive controls during any time period. To rule out sample coding errors, we utilized a criterion of at least two positive culture results to identify an infected subject. A recently published study genotyped a subset of *Entamoeba* species isolated from our study subjects and revealed that, except for two rare transient exceptions, the same isolate was found over time in each individual studied, even for up to six positive cultures over 3 years (34). Therefore, in our study use of a duplicate positive criterion should not significantly mask the occurrence of new infections.

Following cure of ALA, subjects were highly immune to intestinal infection by *E. dispar*. There are no luminal amebicidal agents available for use in South Africa; therefore, we expected a high incidence of *E. histolytica* and *E. dispar* intestinal infection among ALA subjects upon entry into the study. Stool cultures revealed a lower than anticipated (16) prevalence of new *E. histolytica* infections in controls, making it impossible to determine whether ALA subjects, once cleared of their original infection, were immune to new *E. histolytica* infections by comparison to controls. Use of seroconversion criteria revealed that the rate of new asymptomatic *E. histolytica* infections in controls was actually sevenfold higher than suggested by the stool culture data. However, seroconversion criteria cannot be applied to ALA subjects, as almost all (>93%) were seropositive upon entry into the study.

Control subjects with intestinal antilectic IgA antibodies had no history of ALA or colitis; mucosal antibody responses were most likely due to a relatively recent asymptomatic *E. histolytica* or *E. dispar* intestinal infection. *E. histolytica* infections are well documented to produce humoral antilectic antibody responses (28, 32). In addition, intestinal antilectic IgA antibodies were more prevalent than humoral antilectic IgA or IgG antibodies and positively associated with new *E. dispar* infections, suggesting that *E. dispar* can induce a mucosal but not a humoral immune response. We were able to conclude that the low levels of intestinal antilectic IgA antibodies present in ELISA-positive control subjects are not sufficient to provide immunity to new *E. dispar* infections.

We suggest that the immunity to *E. dispar* infection found in our prospective longitudinal study of ALA patients is due to the high levels of intestinal antilectic IgA antibodies present. Haque et al. (17) reported that children previously treated for amebic colitis had a delay in acquisition of new *E. histolytica* intestinal infections and that this relative immunity correlated

**TABLE 2. Prevalence of *E. histolytica* infection in ALA cases and close associate controls***

<table>
<thead>
<tr>
<th>Group</th>
<th>No. infected/no. in group (% infected) at:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0-6 mo</td>
</tr>
<tr>
<td>ALA cases</td>
<td>5/80 (6.3)*</td>
</tr>
<tr>
<td>New infections</td>
<td>6/8</td>
</tr>
<tr>
<td>Controls</td>
<td>9/737 (1.2)*</td>
</tr>
<tr>
<td>New infections</td>
<td>13/19</td>
</tr>
</tbody>
</table>

* Includes only those subjects who had samples collected at baseline and during follow-up. *, P < 0.05 for prevalence of *E. histolytica* infection in ALA cases compared to controls. †, P < 0.05 for prevalence of *E. histolytica* infection in controls 9 to 18 months compared to 21 to 27 and 30 to 36 months of follow-up.

**TABLE 3. Prevalence of *E. dispar* infection in ALA cases and close associate controls***

<table>
<thead>
<tr>
<th>Group</th>
<th>No. infected/no. in group (% infected) at:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0-6 mo</td>
</tr>
<tr>
<td>ALA cases</td>
<td>47/719 (6.5)</td>
</tr>
<tr>
<td>New infections</td>
<td>56/74</td>
</tr>
</tbody>
</table>

* Includes only those subjects who had samples collected at baseline and during follow-up. *, P < 0.05 for lower prevalence of *E. dispar* infection in ALA cases compared to controls.
with the presence of intestinal antilectin IgA antibodies. Of interest, this immunity was observed only during the first five months of follow-up (17). In a subsequent study, Haque et al. (18) found that intestinal antilectin IgA antibodies were detectable for an average of only 31 days; this is in marked contrast to our findings following either ALA or during follow-up of intestinal IgA in antibody-positive asymptomatic controls.

Given the 85% homology in lectin amino acid sequence and shared epitopes between the E. dispar and E. histolytica lectins (23), we propose that immunity to E. dispar may be a surrogate marker for immunity to E. histolytica. Unfortunately, the insensitivity of stool cultures every 3 months for E. histolytica infection (4) and the in vitro adherence-inhibitory activity of murine fecal antilectin IgA antibodies (8), it is likely that intestinal antilectin IgA antibodies are mediating the immunity observed.

A commercially available ELISA (DiaTech Labs) for detection of lectin antigen has been utilized successfully (19). The conditions for performance of the ELISAs in this report are not identical to the commercial assay; in addition, the field conditions in South Africa resulted in a substantial delay (hours) between collection and processing of samples. We found an unacceptably low rate of correlation of lectin antigen positive ELISA results with positive stool cultures for E. histolytica and E. dispar, and therefore we could not apply antigen detection technology in this study. Previously, we successfully performed hundreds of fecal antigen detection ELISAs on samples collected in Cairo, Egypt, with excellent correlation to culture results (3, 4, 6). However, other investigators have found a similar lack of sensitivity and specificity with earlier generations of the DiaTech assay when applied to field conditions in the tropics (E. Tannich and T. F. H. G. Jackson, personal communications). It is important to emphasize that use of the DiaTech assay per the manufacturer’s instructions was not performed in this study. Studies utilizing PCR for detection of parasite ribosomal DNA confirm that stool culture withzymodeme determination underestimated the incidence of E. histolytica but not E. dispar infection (9).

In summary, we found that subjects cured of ALA have a high prevalence and level of both intestinal and serum IgA antibodies directed against the amebic galactose-inhibitable adherence lectin. Both E. histolytica and E. dispar contain functional galactose binding lectin molecules with multiple shared epitopes (23, 25). Cure of ALA is followed by a high level of immunity to E. dispar intestinal infection for the entire 36 months of our study. Therefore, our data and those of others (17, 18) indicate that mucosal antilectin IgA antibodies may mediate immunity in adults to new intestinal infections by E. dispar or E. histolytica. Use of a more sensitive and specific diagnostic test (PCR) rather than use of stool culture and zymodeme determination will allow us to more directly address this hypothesis. Numerous strategies have been developed for use of the E. histolytica galactose-inhibitable lectin as a subunit amebiasis vaccine, especially to elicit protective intestinal antilectin IgA antibodies (33). Clearly, if antilectin IgA antibodies have a role in human immunity to E. histolytica intestinal infection, as indicated by this study, such vaccine strategies should continue to be actively pursued.

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