Immunoglobulin Subclass Distribution and Diagnostic Value of Leishmania donovani Antigen-Specific Immunoglobulin G3 in Indian Kala-Azar Patients

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Visceral leishmaniasis, or kala-azar, a fatal tropical disease, remains problematic, as early diagnosis is difficult and treatment often results in drug resistance and relapse. We have developed a sensitive enzyme-linked immunosorbent assay (ELISA), using leishmanial membrane antigenic extracts (LAg) to detect specific antibody responses in 25 untreated Indian visceral leishmaniasis patients. To investigate the pathogenetic significance of isotype markers in kala-azar, relative levels of specific immunoglobulin G (IgG), IgM, IgA, IgE, and IgG subclasses were analyzed under clinically established diseased conditions. Since LAg showed higher sensitivity for specific IgG than lysate, the immunoglobulin isotype responses were evaluated, with LAg as antigen. Compared to 60 controls, which included patients with malaria, tuberculosis, leprosy, and typhoid and healthy subjects, visceral leishmaniasis patients showed significantly higher IgG (100% sensitivity, 85% specificity), IgM (48% sensitivity, 100% specificity), and IgE (44% sensitivity, 98.3% specificity) responses. Low levels of IgA in visceral leishmaniasis patients contrasted with a 13-fold-higher reactivity in sera from patients with leprosy. Among IgG subclasses, IgG1, -3, and -4 responses were significantly higher in visceral leishmaniasis patients than in the controls. IgG2 response, however, was significantly higher (twofold) in leprosy than even visceral leishmaniasis patients. The rank orders for sensitivity (IgG = IgG1 = IgG3 > IgG4 > IgG2 > IgM > IgE > IgA) and specificity (IgM = IgG3 > IgE > IgG4 > IgG2 > IgG > IgG1 > IgA) for LAg-specific antibody responses suggest the potentiality of IgG3 as a diagnostic marker for visceral leishmaniasis.

MATERIALS AND METHODS

Study subjects. The subjects of the present investigation were 25 Indian patients with visceral leishmaniasis admitted to School of Tropical Medicine, Calcutta, India. These patients came from Bihar (eastern India), one of the main areas of endemicity. Diagnosis of these patients was confirmed parasitologically in the case of malaria, and sera were collected before treatment.

Preparation of antigen. Leishmania AG83, originally isolated from an Indian kala-azar patient, was cultured in vitro for antigen preparation as described earlier (1). Briefly, stationary-phase promastigotes, harvested after the third or fourth passage, were washed four times in cold phosphate-buffered saline (PBS) (pH 7.2) and resuspended in the same Tris buffer and sonicated in an ultrasonicator. The suspension was centrifuged at 4,390 × g for 30 min, and the supernatant containing the LAg was harvested and stored at −70°C until use. The amount of protein obtained from 1.0 g of cell pellet as assayed by the method of Lowry et al. (26), was 16 mg. The lysate used in this study was prepared from 5 × 10⁷ stationary-phase promastigotes per ml according to the method of Jaffe and Zalis (21). Protein concentration (5 mg/ml) was assessed as described above.

Enzyme-linked immunosorbent assay (ELISA). For serological studies, microtiter plates (Teraso) were coated overnight with 2 μg of lysate or LAg per well. For Leishmania-reactive immunoglobulin G (IgG), IgM, IgA, and IgE antibody determination, the antigen-coated plates were incubated with sera diluted 1:1,000-fold, and reacted with peroxidase-conjugated goat anti-human IgG, IgM, IgA, and IgE polyclonal antibodies (Sigma Immunochemicals) at a 1:5,000 dilution and developed with o-phenylenediamine dihydrochloride (1). For IgG subclass determination, human sera were reacted with mouse anti-human IgG1,
IgG2, IgG3, and IgG4 monoclonal antibodies (1:3,000 dilution; Sigma Immunochemicals). Bound antibodies were detected with peroxidase-conjugated goat anti-mouse IgG (1:5,000 dilution; Sigma Immunochemicals) (1).

Statistical analysis. All data comparisons were tested for significance by using Student’s t test; P values of < 0.05 were considered significant. The lower limit of positivity (cutoff) was determined by the mean of healthy controls + 2 standard deviations (13, 14).

RESULTS

Serum IgG specificity for L. donovani lysate and LAg. Reactivities of serum IgG antibodies of kala-azar patients to the parasite lysate were compared to those of LAg. At a 1:1,000 dilution of sera, 20 of 25 patients were positive for the lysate, with IgG absorbance values ranging from 0.319 to 0.493 (Fig. 1a). Reactivity with LAg, however, resulted in 100% sensitivity, with significantly higher IgG absorbance values (1.517 to 2.066; Fig. 1b). Serum specimens from patients with diseases such as malaria, typhoid, tuberculosis, and leprosy were negative for the lysate, and only 1 of 12 serum samples from normal control individuals analyzed was found to be positive (Fig. 1a). Conversely, 1 of 12 healthy controls and 1 of 15 malaria patients were positive, while all typhoid and tuberculosis serum specimens were negative for LAg (Fig. 1b). The highest cross-reactivity was observed with sera from leprosy patients (seven of eight samples). However, the mean ± standard deviation of IgG absorbance (0.403 ± 0.176) of these specimens was just above the cutoff value of 0.276 and significantly lower than the mean IgG response observed with kala-azar patient sera. Since antibody reactivities of sera from kala-azar patients with LAg were higher than with lysate and 100% sensitive, Ig subclass distribution analysis was restricted to LAg.

LAg-specific serum Ig antibodies. Antibody reactivities of IgG, IgM, and IgE of sera from patients with visceral leishmaniasis with LAg were significantly higher than those of normal controls and those of patients with other diseases such as malaria, typhoid, tuberculosis, and leprosy (P < 0.05; Table 1). The IgA reactivity with LAg was, however, predominant in sera from patients with leprosy, with titers 13-fold higher than those even of patients with visceral leishmaniasis.

<table>
<thead>
<tr>
<th>Serogroup</th>
<th>IgG</th>
<th>IgM</th>
<th>IgA</th>
<th>IgE</th>
<th>No. of patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>VL</td>
<td>1.839 ± 0.143&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.878 ± 0.455&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.039 ± 0.084</td>
<td>0.083 ± 0.043&lt;sup&gt;c&lt;/sup&gt;</td>
<td>25</td>
</tr>
<tr>
<td>HC</td>
<td>0.094 ± 0.091</td>
<td>0.250 ± 0.295</td>
<td>0.051 ± 0.93</td>
<td>0.022 ± 0.035</td>
<td>12</td>
</tr>
<tr>
<td>MAL</td>
<td>0.085 ± 0.100</td>
<td>0.140 ± 0.130</td>
<td>0.002 ± 0.007</td>
<td>0.003 ± 0.004</td>
<td>15</td>
</tr>
<tr>
<td>TYP</td>
<td>0.031 ± 0.037</td>
<td>0.158 ± 0.205</td>
<td>0.091 ± 0.162</td>
<td>0</td>
<td>10</td>
</tr>
<tr>
<td>TB</td>
<td>0.110 ± 0.078</td>
<td>0.315 ± 0.281</td>
<td>0.158 ± 0.189</td>
<td>0</td>
<td>15</td>
</tr>
<tr>
<td>LEP</td>
<td>0.403 ± 0.176&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.249 ± 0.246</td>
<td>0.518 ± 0.312&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.006 ± 0.011&lt;sup&gt;c&lt;/sup&gt;</td>
<td>8</td>
</tr>
</tbody>
</table>

<sup>a</sup> Absorbance values (means ± standard deviations).
<sup>b</sup> VL, visceral leishmaniasis; HC, healthy controls; MAL, malaria; TYP, typhoid; TB, tuberculosis; LEP, leprosy.
<sup>c</sup> Significantly different (P < 0.05) from the value for healthy control group as calculated by Student’s t test.
Antigen-specific distribution of serum IgG subclasses. Since the analysis of sera from patients with kala-azar revealed high levels of IgG antibody response to LAg, the IgG subclass specificity was further examined. The results demonstrate that serum samples from all 25 visceral leishmaniasis patients were positive for IgG1, IgG3, and IgG4 antibodies, whereas 21 of 25 had antibodies of the IgG2 subclass (Fig. 2). All these IgG subclasses showed significantly higher reactivity with LAg than normal controls with the dominance of IgG1, in agreement with a previous report (38). While LAg-specific reactivities of all the IgG subclasses were minimal with sera from patients with other diseases, sera from patients with leprosy showed significant levels of antibody responses. Seven of eight samples tested had IgG1 and IgG2 subclasses, and four had IgG4. Moreover, the mean IgG2 response in sera from patients with leprosy was twofold higher even than that of patients with visceral leishmaniasis (Fig. 2). Surprisingly, however, samples of neither leprosy nor any disease other than kala-azar had LAg-specific IgG3 subclass antibodies. Table 2 summarizes the sensitivity and specificity of IgG, IgM, IgE, and IgG subclasses of sera from patients with visceral leishmaniasis.

### DISCUSSION

We found that while a high proportion of Indian kala-azar patients have elevated levels of anti-LAg IgG, IgM, IgE, and IgG subclass antibodies, IgG1, IgG1, IgG3, and IgG4 were present in sera from all the patients, with IgG3 being specifically associated with this disease. Although investigations in murine models of *Leishmania major* and *L. donovani* infections clearly demonstrate a Th-2/IL-4/IgG1 relationship with disease progression, and a Th-1/IFN-γ/IgG2a relationship with resistance and protective immunity (1, 2, 4), such a relationship in humans is not fully understood. An association between antibody isotypes, cytokine profiles, and pathogenesis has been made for some diseases such as leprosy (12, 20), AIDS (6, 27), lymphatic filariasis (25), onchocerciasis (32), and malaria (28). In American cutaneous leishmaniasis, strong cell-mediated immunity and the predominance of IgG1, IgG2, and IgG3 iso-types in localized cutaneous and mucocutaneous leishmaniasis have been linked with Th-1 reactivity, whereas IgG4 subclass antibody response in sera of diffuse cutaneous leishmaniasis patients has been correlated with a Th-2 cell response (7, 9, 29, 35). Cytokine analysis of human visceral leishmaniasis suggests that the Th-2 response will be stronger than the Th-1 response during the active phase of the disease (8, 16, 23, 40). Investigations of IgG subclass response during disease show significant stimulation of all the IgG subclasses in Sudanese patients, with higher levels of IgG3 and IgG4 than IgG1 (13). Conversely, Venezuelan patients have a dominant IgG1 response followed by IgG4 (38). Indian kala-azar patients also showed a predominant IgG1 subclass antibody response, but the levels of IgG3, IgG4, and IgG2 were also significant. These subclasses of human IgG are endowed with unique biological and functional properties, including their response to different types of antigens (22). The elicitation of IgG1, IgG3, and IgG4 antibodies in kala-azar sera may be due mostly to the presence of protein antigens, and the elicitation of IgG2 antibodies in kala-azar sera may be due mostly to the presence of carbohydrate antigens, as reported for viral, bacterial, and parasitic infections (6, 12, 20, 25, 32). Induction of IgG1 and IgG2 is

### TABLE 2. Percent sensitivity and specificity of IgG, IgM, IgA, IgE, and IgG subclasses in visceral leishmaniasis patients

<table>
<thead>
<tr>
<th>Antibody</th>
<th>Sensitivity (%)</th>
<th>Specificity (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>IgG</td>
<td>100</td>
<td>85.0</td>
</tr>
<tr>
<td>IgM</td>
<td>48</td>
<td>100.0</td>
</tr>
<tr>
<td>IgA</td>
<td>4</td>
<td>80.0</td>
</tr>
<tr>
<td>IgE</td>
<td>44</td>
<td>98.3</td>
</tr>
<tr>
<td>IgG1</td>
<td>100</td>
<td>83.3</td>
</tr>
<tr>
<td>IgG2</td>
<td>84</td>
<td>86.6</td>
</tr>
<tr>
<td>IgG3</td>
<td>100</td>
<td>100.0</td>
</tr>
<tr>
<td>IgG4</td>
<td>100</td>
<td>91.6</td>
</tr>
</tbody>
</table>

*Calculations were done with respect to the ELISA cutoff values of 0.276, 0.840, 0.237, 0.092, 0.102, 0.108, 0.132, 0.095 optical density for IgG, IgM, IgA, IgE, IgG1, IgG2, IgG3, and IgG4, respectively.*
IFN-γ dependent, and IgG3 and IgG4 depend on IL-4 and are down regulated by IFN-γ (15, 24). The elevation of IFN-γ in kala-azar patients (23, 40) and the strong reactivity of IgG1 during disease appear to be consistent with the above observations. Their presence, however, fails to control the infection. The absence of IL-2, a Th-1 mediator, suggests a lack of Th-1 response during disease (8, 11). Stimulation of serum IgG3 and IgG4 during infection, together with the expression of cytotoxins such as IL-10 and IL-4 (16, 40), which are also responsible for the upregulation of these IgG subclasses, provides further evidence in support of a Th-2 cell response in determining the outcome of the disease. One explanation for the presence of IgG1 in kala-azar patients may be due to IFN-γ derived from alternative cell sources such as natural killer cells and γδ T cells (10, 39).

LAg-associated serological responses of patients with diseases other than kala-azar were observed to be maximal for leprosy for all isotypes except IgM, IgE, and IgG3. In contrast to a previous report of low reactions of leprosy sera with soluble extracts of Leishmania promastigote antigen for all isotypes (38), LAg gave strong reactions with IgG, IgA, IgG1, and IgG2 and low reactivity with IgG4. Further, reactions with IgA and IgG2 were 13- and 2-fold higher, respectively, than even kala-azar patient sera. Leprosy sera show reactivity with lipoarabinomannan B (LAM), a carbohydrate component of Mycobacterium leprae, through IgG2 and IgG4 and rarely with IgG3 (12). Phenolic glycolipid (PGL-1), another cell wall carbohydrate of M. leprae, reacts strongly with IgA (31) and IgG1 (12) antibodies in leprosy sera. While it is not understood how antibodies in leprosy sera react with LAg, these observations point to cross-reacting epitopes of LAM and PGL-1 in Leishmania LAg.

Amongst all the Ig isotypes and IgG subclasses studied, only IgG3 showed 100% sensitivity and specificity for LAg in visceral leishmaniasis patients. Hence, IgG3 antibody may be a more specific marker for this disease than IgG, which shows low cross-reactivity with other diseases and significant reactivity with leprosy. Moreover, we have found that although there is a decline in the levels of IgG and its subclasses after successful treatment, the decrease is maximal in IgG3 (data not shown), suggesting that IgG3 may be a useful tool for diagnosis as well as for the prognosis of visceral leishmaniasis. IgG3 elevation during leishmaniasis was reported earlier (13, 29, 34), and high specificity and sensitivity for IgG3 have been found in Sudanese visceral leishmaniasis patients (13). Better sensitivity and specificity for IgG3 observed in our studies may be due largely to the specificities of the antibodies to the antigen studied (35) in addition to ethnic variation and differences in parasite genotypes. The significance of IgG3 specificity in visceral leishmaniasis is not clearly understood. IgG3 in malaria is associated with recovery from the fatal disease (33), and skewing of the response toward the IgG3 subclass is merozoite receptor-bearing lymphocytes (36). However, the protective role of leishmaniasis-specific antibodies in human visceral leishmaniasis is still controversial. In conclusion, our serological data demonstrate the potentiality of LAg as an important antigen in the diagnosis of the outcome of infection with Leishmania donovani, with IgG3 as a marker for the identification of individuals with visceral leishmaniasis.

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REFERENCES