HOST RESPONSE TO INFECTION

Distribution of immunoglobulin classes and IgG subclasses against a culture filtrate antigen of Burkholderia pseudomallei in melioidosis patients

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The class and subclass distribution of antibody response to the culture filtrate antigen (CFA) of Burkholderia pseudomallei was examined in the sera of 45 septicaemic and 17 localised melioidosis cases and 40 cases clinically suspected of melioidosis and the results were compared with those from high-risk and healthy control groups. The geometric mean titre index (GMTI) values for all classes and subclasses of immunoglobulins examined were higher for sera from the proven and clinically suspected melioidosis cases than for the control groups. However, the highest response in the three patient groups was that of IgG with GMTIs ranging from 219.4 to 291.6 and the lowest was for IgM with GMTIs of 22.5, 24.3 and 28.7. The IgA response was intermediate with GMTIs ranging from 119.2 to 170. The GMTIs were highest for IgG in septicaemic and localised infections and for IgA and IgM in localised infections. As regards IgG subclass distribution, IgG1 and IgG2 were the predominant subclasses produced against the CFA in contrast to IgG3 and IgG4, which were produced in low amounts. None of the sera from the control groups had any significant titres of antibodies.

Introduction

Melioidosis is an infectious disease of man and animals caused by a gram-negative saprophytic bacterium, Burkholderia pseudomallei. It presents in man in several forms ranging from inapparent infections, to wound infections, to acute septicaemia with a mortality rate of >60% [1]. Although much is known about the epidemiology, clinical manifestations and the course of the disease and its response to antimicrobial agents, the host immune response towards the infection has still not been completely defined [2–4]. Bacterial antigens secreted or shed into the body would be among the first candidate molecules encountered by the host immune system. Very little is known of the isotypes of antibody produced against these secreted antigens in the host micro-environment. Although cell-mediated immune processes are likely to be necessary in immunity against bacteria, bactericidal and phagocytosis-promoting antibodies may have an important role in the initial clearance of the bacteria.

The major class of immunoglobulin produced in normal human serum is IgG and of its four subclasses, IgG1 amounts to c. 65% of the total IgG [5, 6]. Certain antigens elicit immunoglobulin production restricted to only some of the three major classes, i.e., IgG, IgM, and IgA and perhaps some of the four IgG subclasses [7]. Selective deficiency of one or more of the IgG subclasses could increase susceptibility to recurrent infections or allow an existing infection to be prolonged [8, 9]. On the other hand, the selective increase in one or other class or subclass of antibody may aid the host in eliminating the existing infection and protect against recurrent infection or re-infection. Current knowledge on the antibody isotype distribution in melioidosis patients is poor and restricted to anti-lipopolysaccharide antibody response [10]. Further information on the antibody isotype distribution will aid in the development of a good assay system for diagnosis and monitoring the progression of the disease. The aim of the present study was to determine the immunoglobulin class and IgG subclass distribution against a B. pseudomallei culture filtrate antigen (CFA) in patients with melioidosis.

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Materials and methods

Bacterial strain

B. pseudomallei (CMS01859) isolated from the blood of a patient with meliodosis was used in this study.

Serum specimens

Sera were collected from 45 patients with septicaemic meliodosis, 17 patients with localised meliodosis and 40 patients with clinically suspected meliodosis, at the University Hospital, University of Malaya, Kuala Lumpur and various other hospitals in East and West Malaysia. All septicaemic and localised meliodosis cases were bacteriologically confirmed from blood culture or pus samples and serum samples were collected soon after diagnosis (2–3 days). All the clinically suspected meliodosis cases were diagnosed as fever of unknown origin and results were negative for other infections endemic in the areas such as tuberculosis, typhoid, leptospirosis, typhus and rickettsial fevers. Serum samples from these cases were collected c. 1–3 weeks after onset of illness. In addition, these cases had clinical features suggestive of meliodosis such as prolonged fever, non-healing wounds and ulcers in diabetic patients, community-acquired pneumonia, abscesses, cellulitis, empyema and pleural effusions but did not have specimens available for culture or the culture results were negative. Serum samples were also collected from 25 normal subjects from rural areas of Malaysia (high-risk group) and 15 blood donors from the Kuala Lumpur metropolitan area (healthy control group).

Culture filtrate antigen (CFA)

B. pseudomallei was grown in protein-free Sauton’s medium supplemented with L-asparagine 0.4%. The culture filtrate antigen (CFA) was prepared and concentrated from the spent media of a 24-h culture of B. pseudomallei as described previously [11]. The concentrate was dialysed (12–14 kDa) overnight against 0.01 M phosphate-buffered saline (pH 7.2) and the protein concentration was estimated by the method of Lowry [12] after which sodium azide 0.05% was added. The preparations were stored at −20°C until use.

ELISA for immunoglobulin classes and subclasses

ELISAs for IgG, IgM, IgA and IgG subclasses were performed as described earlier [13, 14] with minor modifications. CFA concentration and dilution of conjugates were optimised by checkerboard titration. The CFA was applied at a concentration of 100 ng/well in 100 μl of 0.05 M Tris-buffered saline (TBS), pH 8.6, in MaxiSorp polystyrene flat-bottomed microtitration plates (Nunc, Denmark) and incubated overnight at 4°C. The wells were then blocked for 1 h at 37°C with bovine serum albumin (Sigma) 1% after which the test sera, serially double-diluted from an initial dilution of 1 in 25 in PBS supplemented with Tween 20 0.05% (PBS/T), were added to duplicate wells and incubated at 37°C for 2 h. Horseradish peroxidase-labelled goat anti-human IgG, IgM or IgA conjugate (Sigma) in PBS/T was then added and incubated at 37°C for 2 h. Tetramethylbenzidine substrate (Sigma) 0.05% in citrate phosphate buffer, pH 6, and H2O2 0.001% was used for detection of the enzyme reaction. The reaction was stopped after 10 min by the addition of 1 N HCl and the OD values were read with a multi-scan ELISA reader at a wavelength of 450 nm. An OD value of 0.05 above the nullified background (blank wells) OD was considered as reactive for specific antibodies and the highest reactive dilution was considered to be the endpoint titre. Samples that gave OD values greater than the mean OD + 3 SD of serum samples from healthy individuals at the cut-off dilution were considered as positive. The cut-off dilution was that which excluded all healthy controls. The geometric mean titre (GMT) for each patient group was calculated with SSPS software application, and as these values were very high, a geometric mean titre index (GMI) was calculated by dividing the GMT of each patient group by the GMT value of the normal control group. Serum from a patient with culture-positive meliodosis was used as the positive control in all plates and an inter-assay variation of <10% was considered acceptable.

For the detection of IgG subclass antibodies, the ELISA was performed as above except that a biotin-streptavidin system was used. After incubation of the CFA with the test sera, optimally diluted biotin-labelled mouse anti-human IgG1, IgG2, IgG3 and IgG4 monoclonal antibodies (MAbs) (Sigma) in PBS/T supplemented with bovine serum albumin 1% were added and incubated at 37°C for 2 h. Optimally diluted horseradish peroxidase-labelled streptavidin (Sigma) was added and incubated at 37°C for a further 2 h. The GMT and GMI values were calculated as described above.

ELISA for antigen-specific IgE

The ELISA for CFA-specific IgE antibody was performed as described by van Knapen et al. [15] with minor modifications. Briefly, the wells of microtitration plates were coated with CFA, blocked and stored as described above. For the assay, 100 μl of serially double-diluted sera (from 1 in 25 in PBS/T supplemented with fetal calf serum 2%) was added and allowed to react with the antigen at 37°C for 2 h. The plates were then washed three times with PBS/T and incubated further with optimal dilution of anti-human IgE-alkaline phosphatase (Sigma) in 0.05 μl TBS, pH 7.4, supplemented with fetal calf serum 2% for 2 h at 37°C. The wells were washed three times with TBS supplemented with Tween 20 0.05% and incubated with 5 mM p-nitrophenyl phosphate in sodium acetate buffer supplemented with 1 mM MgCl2 for a further 30 min at
37°C. The enzyme reaction was read at 405 nm with a multi-scan ELISA plate reader and a nullified OD value >0.05 was considered as reactive for antigen-specific IgE antibodies.

Results

The ELISA results are expressed as percentage positivity of the different classes and subclasses of antibodies in the various groups of the study population (Table 1). In the localised melioidosis group, the percentage positivity was found to be 100% for IgG and IgA and 85% for IgM. The septicemic group showed a positivity of 96% for IgG and 89% for IgA, but only 66% for IgM. The positivity rates for the clinically suspected cases were similar, 94% for IgG and 86% for IgA, but IgM was higher at 84%, similar to the localised infections.

The titres for IgG, IgA and IgM were found to vary from 100 to 409,600 (Fig. 1). The GMTI values were calculated for each patient group. The values for both IgG and IgA were significantly higher than those recorded for IgM (p < 0.005) (Table 2). The GMTI for IgG was 292 for septicemic, 245 for localised infections and 219 for the clinically suspected cases. The GMTI values for IgA were intermediate, 150 for septicemic cases, 170 for localised melioidosis and 119 for the clinically suspected cases, whereas for IgM the values were as low as 23 for the septicemic cases, 24 for the localised melioidosis cases and 29 for the clinically suspected group. The low GMTI value of IgM antibodies was due to elevated levels of antigen-specific IgM antibodies in control groups, whose titres varied from 50 to 3200; however, the IgG and IgA antibody levels were <100 in the non-melioidosis groups. In summary, the mean levels of IgG, IgA and IgM in the melioidosis groups, which included the clinically suspected cases, were significantly higher than those in the high-risk and healthy control groups (p < 0.001, p < 0.005 and p < 0.01, respectively). Antigen-specific IgE antibody was not detected in either the melioidosis or the control sera (results not shown).

All four subclasses of IgG antibodies were detected in the sera of all melioidosis patients and the clinically suspected group. Among septicemic cases 87% had IgG1 and 93% had IgG2 and among those with localised infections 94% had IgG1 and 85.5% IgG2 (Table 1). The clinically suspected group had IgG1 and IgG2 in 96% and 94%, respectively. However, IgG3 was present in only 52% of septicemic, 53% of localised and 48% of clinically suspected cases and IgG4 was present in 60, 84 and 67.5% of cases, respectively. Levels of IgG3 and IgG4 were significantly lower than those of IgG1 and IgG2 (p < 0.05), and were not detected in the sera of either the high-risk or the healthy control groups.

The endpoint titres of IgG1 and IgG2 subclass antibodies varied from 100 to 409,600 (Fig. 2). The GMTI for IgG1 and IgG2 were higher in septicemic cases (117 and 69, respectively), as compared with localised infections (90 and 55, respectively) (Table 2). The endpoint titres for IgG3 and IgG4 were also lower and varied from 50 to 10,000, in all the three melioidosis groups. The GMTI values for IgG3 and IgG4 antibodies in septicemic, localised and clinically suspected cases were significantly higher (p < 0.05) than those in the control sera studied.

Discussion

The IgM and IgG responses to different antigens of R. pseudomallei in sera of patients with melioidosis have been examined by many workers and in general a high antibody level was observed [5, 16, 17]. Current knowledge on the antibody isotype distribution in melioidosis patients is restricted to anti-lipopolysaccharide antibody responses [10]. In the present study, class and subclass antibody responses to a CFA were analysed in melioidosis patients by ELISA. Serum samples from all three melioidosis groups had high levels of IgM, IgA and IgG classes of antibodies in 66–100% of cases as compared with the control groups. Interestingly, the highest percentages for all three major classes of immunoglobulins were found among sera from localised infections, with IgA and IgG in 100% and IgM in

<table>
<thead>
<tr>
<th>Patient group</th>
<th>Total no.</th>
<th>IgG (400)</th>
<th>IgA (800)</th>
<th>IgM (6400)</th>
<th>IgG1 (200)</th>
<th>IgG2 (200)</th>
<th>IgG3 (200)</th>
<th>IgG4 (200)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Septicaemic</td>
<td>45</td>
<td>96</td>
<td>89</td>
<td>66</td>
<td>87</td>
<td>93</td>
<td>52</td>
<td>60</td>
</tr>
<tr>
<td>Localised</td>
<td>17</td>
<td>100</td>
<td>100</td>
<td>85</td>
<td>94</td>
<td>88.5</td>
<td>53</td>
<td>84</td>
</tr>
<tr>
<td>Clinically suspected</td>
<td>40</td>
<td>94</td>
<td>86</td>
<td>84</td>
<td>96</td>
<td>94</td>
<td>48</td>
<td>67.5</td>
</tr>
<tr>
<td>Healthy controls</td>
<td>25</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

*Cut-off dilution for positivity (shown in parentheses) was the dilution which excluded all healthy controls. Samples giving OD values that were greater than the mean OD + 3 SD of healthy controls at the cut-off dilution were considered positive.
Table 2. Geometric mean titre indices of immunoglobulin classes and subclasses.

<table>
<thead>
<tr>
<th>Antibody Type</th>
<th>Geometric Mean Titre (log 10)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>IgM AHA</td>
<td>22.5</td>
<td>0.0012</td>
</tr>
<tr>
<td></td>
<td>20.45</td>
<td>0.0015</td>
</tr>
<tr>
<td></td>
<td>14.98</td>
<td>0.0004</td>
</tr>
<tr>
<td></td>
<td>11.68</td>
<td>0.0007</td>
</tr>
<tr>
<td></td>
<td>8.75</td>
<td>0.0006</td>
</tr>
<tr>
<td></td>
<td>6.7</td>
<td>0.0015</td>
</tr>
<tr>
<td></td>
<td>5.8</td>
<td>0.0008</td>
</tr>
<tr>
<td></td>
<td>4.12</td>
<td>0.0028</td>
</tr>
<tr>
<td></td>
<td>2.7</td>
<td>0.0024</td>
</tr>
<tr>
<td></td>
<td>1.94</td>
<td>0.0029</td>
</tr>
<tr>
<td>IgM AGP</td>
<td>28.7</td>
<td>0.0001</td>
</tr>
<tr>
<td></td>
<td>24.4</td>
<td>0.0003</td>
</tr>
<tr>
<td></td>
<td>19.2</td>
<td>0.0003</td>
</tr>
<tr>
<td></td>
<td>14.6</td>
<td>0.0003</td>
</tr>
<tr>
<td></td>
<td>10.7</td>
<td>0.0004</td>
</tr>
<tr>
<td></td>
<td>7.5</td>
<td>0.0005</td>
</tr>
<tr>
<td></td>
<td>5.5</td>
<td>0.0007</td>
</tr>
<tr>
<td></td>
<td>3.8</td>
<td>0.0015</td>
</tr>
<tr>
<td></td>
<td>2.7</td>
<td>0.0029</td>
</tr>
</tbody>
</table>

Antibody titre (log 10) values were calculated by dividing geometric mean titre (GMT) by the respective GMT value of healthy controls.

![Image of a diagram showing antibody titre indices for different antibody types.]

**Legend:**
- AHA: Anti-Hemolytic Antibody
- AGP: Anti-Globulin Antibody
- SC: Secretory Component
- LC: Light Chain
- CS: Unrelated Specific
- HRG: High Relapsing Group
- HC: Healthy Controls

Geometric mean titre indices were calculated by dividing geometric mean titre (GMT) by the respective GMT value of healthy controls.

**Note:** All data is presented as median values with significant differences indicated by asterisks.
who demonstrated a higher detection rate for IgG antibodies in 82% of cases on days 0–5 after admission. In contrast, IgM antibody was found in 60% of cases. Similarly among septicemic cases, IgG was found in 88% of cases and IgM in 65%. These findings lend further support to the fact that the detection of IgG to *B. pseudomallei* antigen was far superior to the detection of IgM for the early diagnosis of melioidosis.

In post-septicemic melioidosis patients under maintenance therapy, elevated levels of IgM antibodies were maintained throughout the observation period (unpublished observations). The incubation period and duration of illness among septicemic cases is not accurately known. However, among the localised infection group it may be possible to estimate the incubation period, especially in post-trauma sepsis and abscesses. The secretory antigens used in the present study consisted mainly of polysaccharides, which are thymus-independent antigens known to activate B cells to elicit low affinity IgM, and proteins (unpublished observations). These antigens, especially those of a polysaccharide nature, can persist for a long time in the lymphoid tissues and continue to stimulate newly maturing B cells in the production of IgM, which may then later be switched to other isotypes. At the same time these antigens could be continuously sequestered from intracellular organisms, thus priming the immune response.

In another study with lipopolysaccharide as the antigen, it was reported that IgM antibodies were detected only in localised melioidosis cases but not in septicemic cases [10]. Interestingly, the present study showed that patients with localised infections, which were superficial soft tissue infections, had the highest levels of IgA in addition to IgM antibodies. Although very little is known about the humoral response in skin, secretory IgA is known to be present in skin secretions. As B lymphocytes are rarely present in cutaneous tissue, it may be possible that IgA produced by B cells activated in the draining lymph nodes is transported back to the skin via the circulation [19].

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**Fig. 2.** Scattergram showing anti-CFA IgG subclass titres in different study groups. SC, septicemic melioidosis; LC, localised melioidosis; CS, clinically suspected melioidosis; HRG, high-risk group; HC, healthy controls. a, IgG1; b, IgG2; c, IgG3; d, IgG4.
Previous reports have shown that, in general, only IgG and IgA antibodies activate neutrophils for mediator release and induction of phagocytosis of bacteria coated with antibodies of these classes [20]. IgG and IgA, but not IgM [21], were also found to induce the release of significant levels of β-glucuronidase by neutrophils. Human IgM is known to activate the classical complement pathway, whereas IgA and IgE activate the alternative complement pathway [22]. Bacterial antigens have also been reported to elicit IgE production either during infection or immunisation and the presence of IgE has been correlated with disease severity in other bacterial infections [20–24]. However, the current study failed to detect antigen-specific IgE antibodies in meliodosis patients, including those with fulminant sepsis, demonstrating the absence of allergenic antibody-specific epitopes in the CFA used in this study.

Although the hierarchy of levels of different immunoglobulin types was found to be IgG $>$ IgM $>$ IgA in both meliodosis and clinically suspected groups, the GMTI values demonstrated a different pattern, i.e., IgG $>$ IgA $>$ IgM. The difference in the hierarchy is due to the elevated level of reactive IgG as compared with IgG or IgA in sera of healthy individuals. This elevated IgM level in healthy controls may be due to continuous exposure to the organism in the environment. This observation also indicates that elevated levels of IgM antibodies are produced in meliodosis patients due to the presence of IgM-specific epitopes in the extra-cellularly secreted products and surface molecules of B. pseudomallei.

Analysis of IgG isotypes detected that IgG1 followed by IgG2 were the predominant subclasses involved in the humoral immune response to B. pseudomallei infection in man. The hierarchy of anti-CFA IgG subclass antibodies was IgG1 $>$ IgG2 $>$ IgG3 $>$ IgG4. This finding reflects the preferential response of IgG1 and IgG2 to protein and carbohydrate epitopes of this antigen. It has been reported that carbohydrate antigens tend to induce immunoglobulin isotype-restricted responses to IgM and IgG2 [22, 25] and that protein antigens induce IgG1 and IgG3 [26]. In human immunisation studies with pneumococcal vaccines, IgG2 has been found to be the predominant immunoglobulin subclass produced to polysaccharide antigens [18]. In another study, Ho and co-workers demonstrated a low IgG subclass response to O-PS I (isopseudomallei) of B. pseudomallei [10], but predominant levels of IgG1 and IgG2 subclass antibodies in sera of all patient groups studied, which was similar to the findings in the present study. An IgG3 response was seen only in survivors of septicaemic infection, while IgG4 was not detectable. Although the role of different subclasses of IgG antibodies in human meliodosis is not understood, human IgG1, IgG2 and IgG3 are known to activate the classical complement pathway. IgG1 and IgG3 are also known to bind to mononuclear cells, of which macrophages and monocytes have been implicated as the major cell population involved in cell-mediated immunity [27]. Phagocytosis is promoted when the cells bind with the opsonised micro-organisms through specialised receptors for IgG1 and IgG3 and C3b complement [28]. To this end it has been reported that sera from meliodosis patients do indeed mediate phagocytic killing by polymorphonuclear leucocytes, and the killing effect was enhanced by complement [10], giving credulity to the possible role of the subclasses of IgG in the immune response to B. pseudomallei infection.

In summary, this study has demonstrated that all immunoglobulin classes, except IgE, and IgG subclasses were present in the immune response mounted to secretory antigens of B. pseudomallei, and that IgG1 and IgG2 are the predominant subclasses produced against CFA. However, the reasons for the inefficiency of these antibodies in killing or clearing the pathogen in patients in vivo despite the elevated level are unclear. Further work is needed to determine the actual role of elevated IgG1 and IgG2 antibody levels in meliodosis cases. The findings also demonstrate the reliability and sensitivity of secretory antigens and IgG antibody detection for the diagnosis of meliodosis in an endemic area.

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