Antigenaemia during acute graft versus host disease

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Abstract

Aims—Animal studies have shown that antigens present within the gut play an important role in the development of acute graft versus host disease (GvHD) following allogeneic bone marrow transplantation (BMT). In previous studies, inert sugars have been found to penetrate the small bowel mucosa after conditioning therapy for BMT; endotoxaemia can also occur during acute GvHD. Data on absorption of antigenic proteins across the gut following BMT in humans have not been presented as yet.

Methods—Six patients undergoing allogeneic BMT were studied to determine whether enteric ovalbumin absorption increased or endotoxaemia developed during acute GvHD.

Results—Three patients had minimal antigenaemia and no detectable endotoxaemia before receiving conditioning therapy. At the onset of acute GvHD, however, much higher ovalbumin concentrations were detected in those patients with severe antigenaemia. Serum concentrations of specific anti-ovalbumin IgG and IgA, or antiendotoxin IgM or IgG had no bearing on detectable IgG or IgM ovalbumin or endotoxin concentrations.

In five of six patients, small bowel permeability increased, as tested by the lactulose/mannitol sugar absorption test, but detectable ovalbumin absorption increased in only three of these and only two developed endotoxaemia.

Conclusions—Antigens present within the gut can cross the mucosal epithelium during acute GvHD, probably resulting in an enhanced immune response.

Keywords: Acute graft versus host disease, antigenaemia, endotoxaemia.

Enteric microbial antigens are thought to play a role in the development of acute graft versus host disease (GvHD) as has been shown in both animal experiments1–3 and clinical studies.4,6

Antigens are thought to penetrate the mucosa following damage caused by conditioning therapy and sensitise donor lymphocytes to host antigens, enhancing the graft versus host reaction. Conditioning chemotherapy can damage the bowel mucosa for up to four weeks following transplantation, with acute GvHD also compromising barrier function. Indeed, we have shown previously that endotoxaemia is not unusual during acute GvHD.8

Absorption of tiny amounts of antigenic protein occurs in normal individuals and is thought to play an important role in the development of tolerance to gut derived antigens.7 The absorption of antigenic proteins is not well understood. Here, we describe the absorption of ovalbumin and endotoxin across the human small bowel mucosa following allogeneic bone marrow transplantation (BMT) at onset of acute GvHD.

Methods

The study population comprised patients entering the Bone Marrow Transplant Unit at the University Hospital of Wales, Cardiff. Patient details are presented in table 1.

All patients received the same conditioning therapy comprising cyclophosphamide 60 mg/kg on two successive days followed by total body irradiation (1125 cGy given in five fractions over three days at a rate of 5 cGy/minute), were nursed in laminar flow rooms, and received gut decontamination therapy consisting of nystatin, amphotericin, colistin, and co-trimoxazole along with acyclovir. Initially, 18 patients were enrolled into the study but only six successfully completed the tests, almost entirely because of their refusal to repeat the ovalbumin test as this had a particularly bad taste and consistency. Lactulose/mannitol permeability tests and ovalbumin penetration studies were performed and plasma endotoxin/antiendotoxin antibody concentrations determined before administration of conditioning therapy, at onset of GvHD and one day thereafter.

OVALBUMIN PENETRATION STUDY

Ovalbumin is a large phosphoglycoprotein with a molecular weight of 45 000 and comprises 40% of total egg protein.

After an overnight fast, patients drank milk (50 ml) containing gamma irradiated egg powder (equivalent to 6 g ovalbumin) over 15 minutes. Blood samples were taken from a central
line before the test and subsequently at 30, 60, 120, 180, and 240 minutes after ingestion of the milk; the patients took nothing further by mouth during this period. Serum was prepared and stored in aliquots at −60°C until analysed.

DETECTION OF OVALBUMIN
Polyclonal rabbit antiserum was prepared following inoculation with ovalbumin (Sigma, Poole, Dorset, UK). This was purified and labelled with biotin. An enzyme linked immunosorbent assay (ELISA) plate (Falcon) was coated with ovalbumin antiserum (diluted 1 in 10,000) and incubated overnight at 4°C. The plate was washed three times in phosphate buffered saline (PBS)/TWEEN before the addition of 0-1% bovine serum albumin (BSA) in PBS over 15 minutes. After a further wash, a standard curve was constructed (ovalbumin 25 to 0-75 ng/ml) and the patient samples added. This was incubated for one hour and after further washing, anti-ovalbumin/biotin was added (diluted 1 in 2000) for one hour. Again after washing, peroxidase labelled Streptavidin (Sigma) was added. After one hour, a final wash was performed and ortho-phenylene diamine added. This reaction was stopped after 30 minutes with 2 M sulphuric acid and the optical density was then measured at 492 nm. All incubations were carried out at 37°C and all tests were performed in triplicate. The ELISA could detect 0-01 ng/ml ovalbumin with an intra-assay coefficient of variation of 5-9% (25 ng/ml) and 6% (1-5 ng/ml) and an interassay coefficient of variation of 5-7% (25 ng/ml) and 7% (1-5 ng/ml).

DETERMINATION OF ANTIBODIES AGAINST OVALBUMIN
An ELISA plate was coated with ovalbumin (10 ng/ml) and incubated overnight at 4°C. After washing three times (PBS/TWEEN), pooled human serum (diluted 1 in 100 in 1% BSA) was added to four wells as a standard. The patients’ serum samples (diluted 1 in 10) were then added in doubling dilutions (1 in 10 to 1 in 5120) and incubated for one hour at 37°C. After a further wash, antihuman IgG-peroxidase (Sigma) (diluted 1 in 2000 in 0-1% BSA) was added to all wells for a further hour. A final wash was performed before the addition of orthophenylene diamine for 30 minutes; the reaction was stopped with 2 M sulphuric acid. Optical density was then measured at 492 nm. A similar method was used to estimate anti-ovalbumin IgA, but in this instance the pooled human serum was diluted 1 in 10 and antihuman IgA (diluted 1 in 2000; Sigma) was used. The titre was defined as that concentration of patient serum which equaled half the maximum of pooled human serum anti-ovalbumin activity.

LACTULOSE/MANNITOL TEST
After an overnight fast, the patients emptied their bladders and then drank a solution comprising 5 g mannitol, 5 g lactulose and 65 ml water over five minutes. Urine was then collected over five hours. The patients were allowed to eat and drink two hours after starting the test. The total volume of urine passed was recorded and a 20 ml aliquot was taken and stored at −20°C until analysis was performed using the methods of Laker. The per cent excretion of both lactulose and mannitol was measured and the results expressed as a ratio of lactulose to mannitol (normal range <0-02). Mannitol (a monosaccharide, molecular weight 182, radius 0-4 nm) is passively absorbed transcellularly through aqueous pores in the epithelial cell membrane. A decrease in the urinary excretion of mannitol represents epithelial cell loss and a reduction in the cell surface area of the bowel (normal range of excretion 8-40%). Lactulose (a non-digestible disaccharide, molecular weight 342, radius 0-54 nm) is passively absorbed through the intercellular tight junctions and the villous tip extrusion zone. An increase in lactulose excretion represents damage to the tight junctions (normal range of excretion <0-3%).

Expression of lactulose and mannitol excretion as a ratio means that changes in bowel transit time, gastric emptying or renal function do not alter the overall result as each of the probe molecules will be similarly affected.

PLASMA ENDOTOXIN ASSAY
Endotoxin was determined using the limulus chromogenic substrate assay kit (Quadratech, Surrey, UK), adapted for use on a sterile microtitre plate. Absorption at 405 nm was read on an ELISA plate reader (Labsystems) and results converted into pg/ml endotoxin by linear regression analysis from standard curves generated with known endotoxin concentrations in standard plasma. This assay was found to be sensitive to less than 5 pg/ml, specific and reproducible (interassay coefficient of variation <5%).

ANTIENDOTOXIN ANTIBODY ESTIMATION
Antiendotoxin antibody estimation was performed using ELISA. Endotoxin antigens from Escherichia coli EH100, Salmonella minnesota Re595, Salmonella typhimurium SL684, and Shigella flexneri (all from Sigma) were mixed together to form a cocktail which was made up at 1 mg/ml in pyrogen free water. The cocktail was then diluted to 20 mcg/ml in 0-05 M sodium bicarbonate (pH 9-8) containing 0-02 M magnesium chloride, and 100 mcg of this solution was added per well to alternate rows of a 96-well plate (Dynatech) and left to coat overnight at room temperature. After washing the plates in PBS, containing 0-1% (v/v) Tween 20, 0-5% (w/v) sodium azide and 0-02 M magnesium chloride, remaining binding sites were blocked by post-coating in 1% BSA in pyrogen free PBS for one hour at 37°C. Patients’ serum was added to the plates in duplicate at a 1 in 100 dilution in wash buffer containing 4% polyethylene glycol 6000 (w/v) and 1% BSA. The test serum samples were compared with a control serum made from a pool of over 100 normal blood donors, matched for age and sex, which was added at dilutions from 1 in 25 to
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Figure 1  Serial serum ovalbumin concentrations before administration of conditioning therapy.

1 in 1600 in the same buffer. After incubation and washing, alkaline phosphatase conjugated antihuman IgG or IgM (Sigma) (diluted 1 in 500 or 1 in 1000) in the above buffer was added at 100 mcg per well. Finally, 100 mcg of 1 mg/ml substrate (p-nitrophenyl phosphate) (Sigma) were added per well in sodium carbonate/bicarbonate buffer (pH 9-8). The reaction was stopped with 50 mcg per well sodium hydroxide (1 N) and the colour read on a spectrophotometer at 405 nm (Labsystem Multiscan). Absorbances for the test serum samples were converted to a percentage of the control sera (100%).

Results

Before receiving conditioning therapy, none of the patients had detectable endotoxaemia, all had a normal lactulose/mannitol permeability test (table 2), but three patients had very low serum ovalbumin concentrations as early as 30 minutes after ingestion persisting for at least four hours (fig 1). Acute GvHD developed on the following days post-transplant: case 1 on day 17; case 2 on day 8; case 3 on day 10; case 4 on day 22; case 5 on day 10; case 6 on day 14.

With the onset of acute GvHD, high serum ovalbumin concentrations were detected in three patients (fig 2) which seemed to correlate with lactulose excretion. Although patients with high ovalbumin concentration had higher grades of acute GvHD, it was not related to the degree of gut GvHD (tables 2 and 3).

There was no correlation between the detection of serum ovalbumin and the number of days that the test was performed following BMT. Indeed, case 4 was tested on day 23 following BMT and was found to have detectable ovalbumin concentrations. As all patients received exactly the same dose of chemotherapy, resulting gut damage probably does not account for the differences in detectable serum ovalbumin. Indeed, the gut is morphologically normal by 21 days unless acute GvHD has developed.1 2

Serum antiovalbumin IgA and IgG levels did not change between administration of conditioning therapy and the onset of acute GvHD. There was no correlation between serum antiovalbumin immunoglobulin concentration and ovalbumin penetration (tables 2 and 3).

Before conditioning therapy, two patients had detectable ovalbumin concentrations but this was no longer evident following the onset of acute GvHD. Two patients with detectable ovalbumin also developed endotoxaemia. Endotoxaemia was no longer detectable 24 hours after the onset of acute GvHD (data not shown). Again, the development of endotoxaemia did not seem to correlate with endotoxin antibody levels (table 3).

Table 2  Peak serum ovalbumin concentrations, antiovalbumin specific IgG and IgA levels and lactulose/mannitol excretion, and plasma endotoxin/antiendotoxin antibody levels before administration of conditioning therapy

<table>
<thead>
<tr>
<th>Patient No.</th>
<th>Peak Ovalbumin (ng/ml)</th>
<th>Serum Antiovalbumin</th>
<th>Plasma endotoxin (pg/ml)</th>
<th>Serum Antiovaltoxin</th>
<th>Mannitol excretion (%)</th>
<th>Lactulose excretion (%)</th>
<th>L/M ratio</th>
<th>L/M ratio</th>
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<tbody>
<tr>
<td>1</td>
<td>1/8</td>
<td>IgG 1/8 IgA 1/64</td>
<td>&lt;5</td>
<td>43 2 29 1 13 2</td>
<td>0 07</td>
<td>0 005</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>Undetected</td>
<td>1/128 1/8</td>
<td>&lt;5</td>
<td>88 4 60 5 18 2</td>
<td>0 06</td>
<td>0 003</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3/8</td>
<td>1/2 1/256</td>
<td>&lt;5</td>
<td>137 43 5 0 9</td>
<td>0 01</td>
<td>0 011</td>
<td></td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>2/0</td>
<td>1/16 1/64</td>
<td>&lt;5</td>
<td>62 7 48 9 0 7</td>
<td>0 026</td>
<td>0 004</td>
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<tr>
<td>5</td>
<td>Undetected</td>
<td>1/128 1/256</td>
<td>&lt;5</td>
<td>22 4 38 1 19 8</td>
<td>0 05</td>
<td>0 002</td>
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</tr>
<tr>
<td>6</td>
<td>Undetected</td>
<td>1/16 1/4</td>
<td>&lt;5</td>
<td>66 4 40 3 20 6</td>
<td>0 08</td>
<td>0 004</td>
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L,M, lactulose:mannitol.

Table 3  Peak serum ovalbumin concentrations, antiovalbumin IgG and IgA levels and lactulose/mannitol excretion, and plasma endotoxin/antiendotoxin antibody levels at the onset of GvHD with the grade of GvHD shown at the time of testing

<table>
<thead>
<tr>
<th>At onset of acute GvHD</th>
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<tbody>
<tr>
<td>Patient No.</td>
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<tr>
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</tr>
<tr>
<td>1</td>
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We have previously shown that endotoxaemia is readily detectable during acute GvHD.8 Experimental evidence suggests that endotoxin can induce acute GvHD perhaps by the release of secondary cytokines such as tumour necrosis factor.19

Any antigen that penetrates the mucosa should be presented by antigen presenting cells to T4 helper cells leading to the release of antigen specific antibodies by plasma cells thus neutralising the antigen.16 During acute GvHD, antigen presenting cell and gut plasma cell numbers are reduced, particularly in severe disease,17 and therefore the effector response to antigen penetration is impaired. In our study, however, there seems to be no simple correlation between antigen specific antibody levels and antigen penetration, confirming previous reports.20

During acute GvHD, there seemed to be a time lag before ovalbumin could be detected compared with that observed before transplantation. Some patients who had detectable concentrations before transplantation did not have acute GvHD at that time. All patients tested during acute GvHD were taking high dose corticosteroids which can reduce the passive absorption of bacterial antigens from the gut, including endotoxin, and may similarly affect ovalbumin absorption in patients with minimal mucosal damage.21

In conclusion, acute GvHD can result in antigens which usually reside in the gut gaining access to the systemic circulation resulting in an enhanced immune response.

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