Production of antibodies to staphylococcal superantigens in atopic dermatitis

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Abstract
Staphylococcal superantigens (SAG) are implicated in the inflammation of atopic dermatitis. As SAG mediated diseases may be modified by specific antibodies, the antibody response to SAG in atopic dermatitis was investigated. Immunoglobulin (Ig) G to staphylococcal enterotoxin A (SEA), staphylococcal enterotoxin B (SEB), and toxic shock syndrome toxin 1 (TSST-1) were measured by sandwich enzyme linked immunosorbent assay (ELISA) in 74 children with atopic dermatitis and 111 controls. Controls had detectable IgG to SEA, SEB, and TSST-1, which increased with age. Atopic dermatitis subjects had an increased response to SEB at 6 months to 2 years (76% v 42%) and 2 to 7 years (79% v 57%), and equivalent responses to SEA and TSST-1, compared to controls. It is suggested that increased responses to SEB relate to increased colonisation and hence exposure to superantigen producing staphylococcus in atopic dermatitis, and that inflammation of atopic dermatitis is not caused by an inability to make antibody to SAG.

Keywords: atopic dermatitis; staphylococcus; superantigen; antibody

Superantigens are particular antigens capable of inducing massive T cell activation caused by their ability to bind multiple T cell receptors in an antigen non-specific manner. Staphylococcal superantigens (SAG) are potent toxins produced by particular strains of Staphylococcus aureus. Stimulation of T cells by SAG is considered to trigger diseases such as Kawasaki disease and atopic dermatitis. Although there is some information on antibody responses to SAG in normal children, the production of immunoglobulin (Ig) G antibodies in atopic dermatitis has not been described. It is possible that subjects with atopic dermatitis would exhibit increased responses either because of their cutaneous colonisation or because of immune deviation to a Th2 (antibody producing) type response. Alternatively atopic subjects may exhibit reduced antibody responses, in addition to the previously demonstrated reduction in cell mediated immunity to SAG. A decreased capacity for antibody production could make children with atopic dermatitis more prone to superantigen producing S aureus colonisation. To examine these questions we measured serum IgG antibody to SEB, staphylococcal enterotoxin A (SEA), and TSST-1 from children with atopic dermatitis, from non-atopic children, and young adults.

Methods
SUBJECTS
Sera from 74 children with mild to moderate or severe atopic dermatitis, classified according to the criteria of Hanifin and Rajika, were collected from children having blood taken as part of the investigation and management of their atopic dermatitis. Ethics approval for experiments and blood collection from children with atopic dermatitis was obtained from the Royal Children's Hospital Research Foundation ethics committee. Samples were obtained from 44 boys and 30 girls (ratio male: female, 1.5:1). The mean IgE of the subjects was 4433 IU/ml (60–54 296 IU/ml). The majority of the samples came from Caucasian children (n = 61) with the remainder of samples from Asian children (n = 13). There were 33 children with atopic dermatitis aged 6–24 months, 24 aged 2–7 years, and 17 aged 8–16 years. Sera from 111 child and young adult controls were obtained from samples sent to our laboratory for evaluation of immunoglobulins, antinuclear antibodies, complement and/or antigliadin antibodies, where IgE was in the normal range for age: < 6 IU/ml (0–0.2
years), < 45 IU/ml (0.2–0.5 years), < 20 IU/ml (0.5–0.7 years), < 35 IU/ml (1–2 years), < 130 (2–4 years), < 150 IU/ml (4–7 years), < 200 IU/ml (> 7 years), and the requested test was negative. IgE was measured using Quanticlone IgE radioimmunoassay (Kallestad Diagnostics Inc, Chaska, Minnesota, USA) according to the manufacturer’s instructions, with a sensitivity of 5 IU/ml.

Age ranges were examined in groups: 6–24 months, 2–7 years, 8–16 years, and more than 16 years. Infants below 6 months were not included because of the possibility of persistent maternal antibody. Serum was stored at −20°C until assayed.

ELISA FOR TSST-1, SEB, SEA
TSST-1, SEB, and SEA antibodies were measured by sandwich enzyme linked immunosorbent assay (ELISA). High binding plates were coated with 100 µl/well of TSST-1, SEB, and SEA (Sigma Chemical Co, St Louis, Missouri, USA) at 1 µg/ml in carbonate buffer, pH 9.6, and incubated for 16 hours at 4°C. Plates were washed with phosphate buffered saline (PBS)/Tween, blocked with 1% gelatin/PBS/Tween for two hours and washed a further four times. Samples (100 µl/well) were added at 1:1000 (1% gelatin/PBS/Tween) and incubated at 37°C for 90 minutes. Biotinylated monoclonal antihuman IgG (Pharmingen, San Diego, California, USA) 100 µl/well (0.5 µg/ml) was added, incubated at room temperature for 45 minutes, washed, and avidin peroxidase diluted at 1:400 (100 µl/well) was then added and incubated at room temperature for a further 30 minutes. After final washing 100 µl/well of 3,3',5,5'-tetramethyl benzidine substrate (KPL Laboratories, Gaithersburg, Maryland, USA) was added and reaction stopped by addition of 100 µl/well of 2 M H₂SO₄. Colour was read at 450 nm (background 650 nm).

All standards and samples were performed in duplicate. Optimal concentrations of coating superantigen and detection antibody were determined by set up checkerboard experiments. Pooled adult sera (n = 20), aliquoted and stored at −70°C, was used to construct a standard curve using doubling dilutions from 1:250. Arbitrary units of antibody were defined as pool 1:1000 = 100 laboratory units (LU). Concentrations of the samples (1:1000) were then calculated from the standard curve.

Interassay variability was 9%, 5.6%, and 9.6% for TSST-1, SEB, and SEA assays, respectively. An uncoated pair of wells was included in all plates to assess non-specific binding. Lower limit of detection for each assay was calculated using Biomek software, and final lower limit of detection for TSST-1, SEB, and SEA assays was calculated by mean (2 SD) of lower limit of detection. These were 2.4 LU, 8.7 LU, and 2.1 LU for SEB, TSST-1, and SEA, respectively.

STATISTICS
Comparison of atopic and non-atopic age antibody concentrations were performed using Mann-Whitney non-parametric U test. Comparisons of detectable antibody production were performed using χ² test.

RESULTS
CONTROL NON-ATOPIC CHILDREN
A significant proportion of control children had detectable antibodies to one or more of SEA, SEB, and TSST-1. The presence of these antibodies increased with age. Detectable antibodies to SEB were found in 42%, 57%, 78%, and 100% of subjects in the age groups 6–24 months, 2–7 years, 8–16 years, and more than 16 years, respectively (fig 1). Detectable antibodies to SEA rose from 78–95% (fig 2) and to TSST-1 rose from 69–100% (fig 3) over this age range. In subjects less than 16 years old antibodies to SEA and TSST-1 were more frequent than antibodies to SEB (p < 0.005, p < 0.005, respectively). Young children (6–24 months) were capable of significant antibody production, with titres up to 10-fold higher than adult pool in some cases. There was a trend for higher SEB antibody concentration (geometric mean) with increasing age. For all three superantigens the geometric mean of antibody was highest in those over 16 years.

There was no relation between TSST-1, SEB, and SEA IgG concentrations for each individual—that is, those children who made...
significant quantities of antibody to one superantigen did not tend to make more antibody to the other superantigens. The children with no antibody to all three superantigens were equally distributed in terms of age (data not shown).

### Table 1 Percentage of detectable antibodies to SEB, SEA, and TSST-1 in subjects with atopic dermatitis (atopic) and non-atopic controls

<table>
<thead>
<tr>
<th>Age</th>
<th>SEB Control</th>
<th>Atopic</th>
<th>SEA Control</th>
<th>Atopic</th>
<th>TSST-1 Control</th>
<th>Atopic</th>
</tr>
</thead>
<tbody>
<tr>
<td>6–24 months</td>
<td>42 67</td>
<td>78 70</td>
<td>69 70</td>
<td></td>
<td>75 83</td>
<td>95 82</td>
</tr>
<tr>
<td>2–7 years</td>
<td>57 79</td>
<td>86 83</td>
<td>75 83</td>
<td></td>
<td>93 82</td>
<td>100 82</td>
</tr>
<tr>
<td>8–16 years</td>
<td>78 76</td>
<td>93 82</td>
<td>93 82</td>
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<td>&gt; 16 years</td>
<td>100 –</td>
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**Discussion**

Our findings demonstrate that young infants are capable of producing antibody to SAG and indicate that exposure to SAG occurs frequently in the first years of life. As superantigens are proteins, this finding is consistent with previous reports that vigorous responses to other proteins are well established in the first year of life. It is not yet clear whether all infants have the capacity to make antibodies upon superantigen exposure. The finding that children with undetectable IgG to all three superantigens were equally distributed in age suggests that certain individuals may be hyporesponsive to SAG proteins in a non-age dependent manner.

The prevalence of specific antibodies to SAG in normal adults has been reported to range from 30–90%. There are several studies reporting the prevalence of SAG antibodies in normal children. In a survey of 168 children from Utah, 94% of TSST-1 producing *S aureus* carriers and 33% of non-carriers were found to have TSST-1 antibody; other superantigen antibodies were not measured, however. In another study, 33% of Scandinavian subjects less than 10 years old but 88% over 10 years had detectable antibody to TSST-1. In a small cohort of 22 control children from a study of Kawasaki disease, the proportion of antibodies detected to TSST-1, SEA, and SEB was 45%, 27%, and 27%, respectively. These proportions are significantly less than we observed to SEB, SEA, and TSST-1 in children of similar ages. This may be caused by the increased sensitivity of our assay, as an amplification step using avidin peroxidase was included in our assay but not the previous study, and avidin peroxidase has been reported to enhance the sensitivity of sandwich ELISAs. As the controls in this study were undergoing laboratory investigations, it is also possible that they do not represent completely healthy children, although all the investigations requested were negative. Although this is a limitation of the study, it would be difficult to obtain this number of samples from completely healthy non-atopic children. Furthermore, the levels of antibody in these controls would likely have been equivalent or higher than those of “healthy controls”, and thus the comparison of “healthy controls” with atopic children is likely to have given similar results to the present controls.

The prevalence of colonisation of normal subjects with various SAG producing strains within our study community are not known, although their prevalence has been examined elsewhere. Jacobson *et al* examined nasal carriage of superantigen producing *S aureus* and simultaneous prevalence of detectable IgG to TSST-1 in 168 children in the USA. They observed TSST-1 to be the most prevalent SAG identified, being present in 29% of *S aureus* strains with a carriage rate of 9.5%. SEA, B, C, and D were identified in only 6.5, 3.0, 1.2, and 1.2% of *S aureus* isolates, respectively. A lower carriage rate of 5% of superantigen producing *S aureus* was observed by Hoeger *et al* in a group of normal children from...
Antibody response to Staphylococcus aureus superantigens in atopic dermatitis

We postulate that the diminished cell mediated immunity to S. aureus and SEB previously observed in children with atopic dermatitis is due to increased antibody levels against superantigens. Infants and young children with atopic dermatitis had higher titres of anti-SEB antibody we observed were caused by increased superantigen exposure. As colonisation with S. aureus in atopic dermatitis is largely confined to cutaneous sites, it is possible that cell mediated rather than antibody responses are important for S. aureus eradication in this situation. We postulate that SAG do play a role in the inflammation of atopic dermatitis. Decreased production of IFN-γ in response to S. aureus and SAG in atopic dermatitis could reduce eradication of superantigen producing strains of S. aureus. Persistence of SAG on the skin would therefore contribute to T cell activation, production of proinflammatory cytokines and thus to cutaneous inflammation. We have not found evidence to suggest that an inability to make IgG antibody to SAG is a factor in the development or ongoing inflammatory response of atopic dermatitis.

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Wilson C, Lewis D, Penix L. The absence of evidence that superantigens are protective in this regard? In this useful paper Campbell and Kemp observed that all children were exposed at some point to staphylococcal superantigens (as one might expect with the large environmental reservoir and ubiquity of these organisms). By 16 years of age all their cases had antibodies; however, these did not appear to protect them against toxic shock syndrome, or against staphylococcal colonisation. This resembles the situation in adults, in whom the presence of antibodies to staphylococcal cell wall components and toxins and secondary responses do not protect against invasive or toxin-mediated staphylococcal disease.

What are the possible links between staphylococcal allergy? Staphylococci frequently infect children with very high levels of IgE, as in Job’s syndrome. Their superantigens and protein A (a B cell mitogen) may stimulate IgE production directly, and it is possible that their superantigens switch off or anergise T cells normally involved in preventing allergic responses. Alternatively, by activating regulatory γδ T lymphocytes in skin, superantigens may influence local inflammation, a situation observed in contact dermatitis. It is also possible that there is an association between susceptibility to staphylococcal colonisation and allergic disease. The observations made by Campbell and Kemp help begin to dissect the complex relation between staphylococcal superantigens and allergic disease: they demonstrated frequent, early antibody responses to staphylococcal superantigens in healthy children. It seems that staphylococci and their superantigens do not cause allergy in most children, but they are strongly associated with atopic dermatitis in some.

This paper leaves us with two particularly important questions. What determines if and when an individual becomes persistently colonised by staphylococci, if not antibodies—such factors are central to the control of atopic dermatitis. Second, is there a causal relation between staphylococcal and atopic dermatitis. It is also possible that there is an association between susceptibility to staphylococcal colonisation and allergic disease. The observations made by Campbell and Kemp help begin to dissect the complex relation between staphylococcal superantigens and allergic disease: they demonstrated frequent, early antibody responses to staphylococcal superantigens in healthy children. It seems that staphylococci and their superantigens do not cause allergy in most children, but they are strongly associated with atopic dermatitis in some.

Commentary

Children are never alone: colonised by vast numbers of prokaryotes in the gut, upper airways, and skin, they grow and develop in a sea of microbial activity. Most commensal organisms are not pathogenic, but a number can cause serious illness. Staphylococcus aureus or coagulase positive staphylococcus is one of our oldest commensals. Most newborns have the organism around their healing umbilical stump. This proportion falls to some 30% in older children with nasal or fingernail carriage. Colonisation increases following skin trauma, burns or the use of occlusive dressings close to the skin. At some time or another we will all have an invasive infection with staphylococci, usually minor, perhaps with the characteristic hallmark of a localised abscess.

Staphylococci also give rise to toxin mediated diseases including food poisoning, toxic shock syndrome, and scalded skin syndrome. Within a genome of some 3000 kbp, staphylococci may harbour a potential armamentarium of over 10 toxins. Many staphylococcal toxin mediated disorders are mediated by molecules that are particularly effective T lymphocyte activating agents or superantigens. Nanomolar quantities of these can cause T cell proliferation.

The clinical significance of these molecules, powerfully demonstrated in the test tube or animal model, has been difficult to assess. Although devastating at one end of a spectrum with fatal cases of toxic shock syndrome, at the other end most staphylococcal food poisoning probably never presents to medical staff.

General paediatricians encounter staphylococcal superantigens daily in atopic dermatitis. Superantigens are excellent candidates for promoting the inflammatory T cell infiltrates typical of moderate or severe atop dermatitis. These molecules produce atopic dermatitis-like lesions if applied to normal skin and in patients recovering from toxic shock syndrome. Removal of staphylococci and their toxins improves most atopic dermatitis. Surely, therefore, antibodies to the superantigens are protective in this regard? In this useful paper Campbell and Kemp observed that all children were exposed at some point to staphylococcal superantigens (as one might expect with the large environmental reservoir and ubiquity of these organisms). By 16 years of age all their cases had antibodies; however, these did not appear to protect them against toxic shock syndrome, or against staphylococcal colonisation. This resembles the situation in adults, in whom the presence of antibodies to staphylococcal cell wall components and toxins and secondary responses do not protect against invasive or toxin-mediated staphylococcal disease.