Abstract: The application of food-specific IgG antibody detection in allergy dermatoses was explored. 181 patients with allergy dermatoses were diagnosed from January to September 2014 and 20 healthy subjects were selected. Fourteen kinds of food-specific IgG antibodies were detected by ELISA method among all the subjects. The positive rates of IgG antibody of the patient group and the healthy group were respectively 65.2% and 5.0%. The positive rates of IgG antibody of egg, milk, shrimp and crab took a large proportion in three groups of patients with three kinds of allergy dermatoses of urticaria, eczema and allergic dermatitis, the proportion of which was respectively 70.2%, 77.8% and 71.7%. Among urticaria and allergic dermatitis patients with positive antibody, the positive rate of children was significantly higher than that of adults (p<0.05) while there was no significant difference between children and adults among eczema patients with positive antibody (p>0.05). Allergy dermatoses are closely related to food-specific IgG antibodies, and the allergy dermatoses patients have a high incidence rate of food intolerance; detecting IgG antibody in the serum of patients is of great significance for the diagnosis and treatment of allergy dermatoses.

Keywords: food-specific IgG antibody, allergy dermatoses, food intolerance

1 Introduction

The allergy dermatosis is an inflammatory dermatosis caused by allergic reaction, featuring complicated causes, uncertain pathogenesis and high recurrence rate [1-3]. Allergy dermatoses are mainly caused by an allergen, which can lead to inflammatory reaction through ingesting and touching [4,5]. Research in the literature [7,8] shows that as much as 40% of people have tolerance against some food to some degree, and food intolerance can cause allergy dermatoses. Meanwhile, clinical data [9] show that IgG antibody detection is closely related to adverse reactions to food and allergy dermatoses involved in IgG is not being paid more attention. This report presents tests conducted to detect food-specific IgG antibodies in the serum of allergy dermatosis patients to explore its role in allergy dermatoses.

2 Data and methods

2.1 General data

181 allergy dermatosis patients who were diagnosed in dermatological department of our hospital from January 2014 to September 2014 were selected including 98 males and 83 females with age ranging from 2 months to 73 years old. The patients include 75 patients with urticaria, 27 patients with eczema and 79 patients with allergic dermatitis. There were also 20 healthy subjects of 11 males and 9 females who participated and had no significant difference on age and gender with patient group.

Ethical approval: The research related to human use has been complied with all the relevant national regulations, institutional policies and in accordance the tenets of the Helsinki Declaration, and has been approved by the authors’ institutional review board or equivalent committee.
Informed consent: Informed consent has been obtained from all individuals included in this study.

2.2 Methods

2ml routine venous blood was collected (There is no need to empty stomach). The food intolerance detection kits of BIOMERICA Company of America and the ELISA method were used to detect the specific antibody IgG in 14 kinds of foods including shrimp, crab, cod, beef, pork, chicken, egg, milk, wheat, corn, rice, tomato, soybean and mushroom. The results were based on the different IgG antibody concentration of the different foods and were classed according to the following categories: weak, mild, moderate, and severe. For details see Table 1.

2.3 Statistical method

All the data was input in the form of EXCEL and the statistical software SPSS18.0 was used to analyze the data. The count data were tested by Chi-square test and when p<0.05, the difference has statistical significance.

3 Result

3.1 Positive rates of IgG antibody of healthy subjects and allergy dermatosis patients

Among 20 healthy subjects, 1 subject was tested positive IgG antibody and the positive rate was 5.0%; among 181 patients with allergy dermatoses, 118 patients were tested positive IgG antibody and the total positive rate was 65.2%. There was a significant difference in the positive rate of IgG antibody of healthy group, yet not for the patient group ($X^2=71.8, p < 0.05$); See Figure 1. There was no significant difference in the positive rate of IgG antibody among urticaria, eczema and allergic dermatitis groups ($p > 0.05$). For details see Table 2.

3.2 Positive rate of food-specific IgG of 14 kinds of foods

For the serum of all the subjects, ELISA method was used to detect the food-specific IgG antibody of 14 kinds of foods; for details see Table 3. It was found that among 14 kinds of foods, the positive rate of IgG antibody of egg, milk, shrimp and crab took a large proportion in the three groups of allergy dermatosis patients. In the group of urticaria, eczema and allergic dermatitis, the proportion of the IgG antibody positive of egg, milk, shrimp and crab in that of the 14 kinds of foods was respectively 70.2%, 77.8% and 71.7%. Among the three groups of allergy dermatoses, the positive rates of egg and milk were above 10%. For the sequence of positive rates of IgG antibody of egg, milk, shrimp and crab in the three groups of allergy dermatosis patients see Table 4.

![Figure 1: Comparison of IgG antibody positive rate between healthy subjects and patients](image)

| Groups                  | n  | Positive Cases | Positive Rate (%) |
|-------------------------|----|----------------|-------------------|
| Healthy Subjects        | 20 | 1              | 5.0               |
| Urticaria               | 75 | 47             | 62.7              |
| Eczema                  | 27 | 18             | 66.7              |
| Allergic dermatitis     | 79 | 53             | 67.1              |
3.3 Positive Rates of 3 groups of allergy dermatoses in different food tolerance degrees

According to the standard identification of food intolerance, the mild, moderate and severe positive rates of IgG antibody positive patients of urticaria are respectively 29.8%, 40.4% and 29.8%; those of IgG antibody positive patients of eczema are respectively 33.3%, 44.4% and 22.2%; those of IgG antibody positive patients of allergic dermatitis are respectively 58.5%, 32.1% and 9.4%; for details see Table 5.

3.4 Food-specific IgG antibody positive distribution of children and adults

Among 118 patients, 31 children and 87 adults were tested antibody positive patients. The proportion of children and adults among antibody positive patients with urticaria was respectively 19.1% and 80.9%. The proportion of children and adults among antibody positive patients with eczema was respectively 55.6% and 44.4%. The proportion of children and adults among antibody positive patients with allergic dermatitis was respectively 22.6% and 77.4%; among antibody positive patients with urticaria, the antibody positive rate of children was significantly below that of adults ($X^2=58.26$, p=0.031). Among antibody positive patients with allergic dermatitis, the antibody positive...
The rate of children was also significantly below that of adults ($X^2=61.02$, $p=0.029$). Among antibody positive patients with eczema, there was no significant difference between children and adults ($X^2=25.57$, $p=0.11$). For details see Table 6.

Note: *There was a significant difference in comparison with adults group ($p < 0.05$); ∆There was no significant difference in comparison with adults group ($p > 0.05$)

### 4 Discussion

Allergy dermatosis is a kind of dermatosis caused by allergen by various ways [1-3]. Studies [4,5] have shown that food intolerance can lead to cutaneous inflammation reaction and that IgG antibody is closely related to food intolerance. In this paper, ELISA method was used to detect the different food-specific IgG antibodies in the serum of patients with urticaria, eczema and allergic dermatitis which are the three kinds of common seen allergy dermatoses in clinics, aiming at providing references for the diagnosis of clinical allergy dermatoses.

In this detection result, one of the 20 healthy subjects was tested with positive IgG antibody and the positive rate was 5.0%. This rate was lower than that in the reference report [10], which was probably caused by the lack of subjects. Among 181 patients with allergy dermatoses, 118 patients were tested with positive IgG antibody and the total positive rate is 65.2%. There was significant difference compared with healthy group ($p < 0.05$); meanwhile, a comparison was conducted on the different positive rates of IgG antibody of urticaria, eczema and allergic dermatitis and there was no significant difference among these groups of positive rates ($p > 0.05$).

Through detecting the food-specific IgG antibody of 14 kinds of foods, it was found that among 14 kinds of foods, the positive rate of IgG antibody of egg, milk, shrimp and crab took large proportion in the three groups of allergy dermatosis patients. In the group of urticaria, eczema and allergic dermatitis, the proportion of the IgG antibody positive of egg, milk, shrimp and crab in that of the 14 kinds of foods was respectively 70.2%, 77.8% and 71.7%; among the three groups of allergy dermatoses, the positive rates of egg and milk were above 10% and the moderate intolerance and severe intolerance were mainly found in milk, egg and shrimp, which was similar to that in the literature report [11-13]. Egg was the most intolerant food for urticaria and eczema while milk was the most intolerant food for allergic dermatitis. Among IgG antibody positive patients with urticaria, the mild, moderate and severe positive rates were respectively 29.8%, 40.4% and 29.8%. Among IgG antibody positive patients with eczema, the mild, moderate and severe positive rates were respectively 33.3%, 44.4% and 22.2%. Among IgG antibody positive patients with allergic dermatitis, the mild, moderate and severe positive rates were respectively 58.5%, 32.1% and 9.4%. The data show that there was mild and moderate food intolerance in the allergic dermatitis group and there was no distribution difference of food intolerance in the urticaria and eczema group.

Among 118 patients, 31 children and 87 adults were tested as antibody positive patients. The proportion of children and adults among antibody positive patients with urticaria was respectively 19.1% and 80.9%. The proportion of children and adults among antibody positive patients with eczema was respectively 55.6% and 44.4%. The proportion of children and adults among antibody positive patients with allergic dermatitis was respectively 22.6% and 77.4%. Among antibody positive patients with urticaria, the antibody positive rate of children was significantly below that of adults ($p < 0.05$). Among antibody positive patients with allergic dermatitis, the antibody positive rate of children was also significantly below that of adults ($p < 0.05$). Among antibody positive patients with eczema, there was no significant difference between children and adults ($p > 0.05$). The data show that the allergy dermatosis of adults was different from that of children. Adult patients with urticaria and allergic dermatitis were more than child patients. Child patients with eczema were more than adult patients, which were different from that in the literature report and probably caused by the lack of eczema subjects.

To sum up, allergy dermatoses are closely related to food-specific IgG antibody, and the allergy dermatoses patients have a high incidence rate of food intolerance. Detecting IgG antibody in the serum of patients is of great significance for the diagnosis and treatment of allergy dermatoses.

**Conflict of interest statement:** Authors state no conflict of interest
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