How dangerous is food allergy in childhood? The incidence of severe and fatal allergic reactions across the UK and Ireland

C F Macdougall, A J Cant, A F Colver

METHODS
The British Paediatric Surveillance Unit (BPSU) of the Royal College of Paediatrics and Child Health studies the epidemiology of uncommon childhood disorders in the UK and Ireland. All consultant paediatricians receive a card every month for reporting any of the 10 to 12 conditions currently under surveillance. The overall response rate was 93.4% in 1999.

Severe food allergic reactions are characterised by angioedema, hypotensive shock, and wheeze. The phrase “severe food allergic reaction” is preferable to “anaphylaxis”, which sometimes describes a severe reaction and sometimes an allergic mechanism. There is professional11 and public12 concern that the number of severe and fatal childhood food allergic reactions is increasing dramatically, leading to increased prescription of epinephrine (adrenaline) autoinjectors,1 anxiety for schools,2 costly actions by food manufacturers,2 and significant parental concern.

Food allergy in childhood is common with a prevalence of between 0.3% and 8%.1 Fatal and severe food allergic reactions are well documented in children.11–13 Although their incidence is not known, some claim the incidence in adults and children is increasing,11–13 while others suggest there is no evidence of this in children.11 We conducted a prospective survey of fatal and severe allergic reactions to food in children from March 1998 to February 2000; together with a retrospective search for fatal allergic reactions from 1990 to February 1998.

Aims: To discover the incidence of fatal and severe allergic reactions to food in a large population of children.

Methods: A retrospective search for fatalities in children 0–15 years from 1990 to February 1998, primarily of death certification at offices of national statistics. A prospective survey of fatal and severe reactions from March 1998 to February 2000, primarily through the British Paediatric Surveillance Unit. Main outcome measures were deaths and severe reactions. A case was deemed severe if one or more of the following criteria was met: cardiorespiratory arrest; need for inotropic support; fluid bolus >20 ml/kg; more than one dose of epinephrine; more than one dose of nebulised bronchodilator. A case was deemed near fatal if intubation was necessary.

Results: The UK under 16 population is 13 million. Over the past 10 years, eight children died (incidence of 0.006 deaths per 100 000 children 0–15 years per year). Milk caused four of the deaths. No child under 13 died from peanut allergy. Two children died despite receiving early epinephrine before admission to hospital; one child with a mild food allergic reaction died from epinephrine overdose. Over the past two years, there were six near fatal reactions (none caused by peanut) and 49 severe ones (10 caused by peanut), yielding incidences of 0.02 and 0.19 per 100 000 children 0–15 years per year respectively. Coexisting asthma is more strongly associated with a severe reaction than the severity of previous reactions.

Conclusions: If 5% of the child population have food allergy, the risk that a food allergic child will die from a food allergic reaction is about 1 in 800 000 per year. The food allergic child with asthma may be at higher risk. Prescribing an epinephrine autoinjector requires a careful balance of advantages and disadvantages.
uncertain or other processes such as an exercise induced reaction were more likely to have been responsible. The severity of previous reactions was classified as mild (rash, only needed advice and antihistamine), moderate (distressed breathing, dizziness, and anxiety), or severe (needing urgent treatment and assessment in hospital).

**Ascertainment of fatal reactions: 1990 to February 2000**

Notifications of deaths were sought from the offices of national statistics as described above. However, we could not be certain that these offices recorded all deaths, and it is important to examine the extent to which other sources yield the same or additional cases. The following sources were also used:

- BPSU 1998–2000
- Database of allergy deaths as a result of all types of allergy in all age groups held by Dr R Pumphrey, an immunologist in Manchester
- Anaphylaxis Campaign—a voluntary organisation with nationwide contacts
- Personal letters to 10 experts in paediatric allergy in the UK
- Asthma and Allergy Information Research, a web based information service for the public, whose director contacted the study
- *Daily Telegraph* and *The Times* stored on CD-ROM.

Paediatricians have shown great interest in the study, and possible cases they recall over the past 10 years have been mentioned to us informally on six occasions. We have followed up such details and all six were already known to us.

**Ascertainment of severe reactions: March 1998 to February 2000**

After cases were notified to the BPSU, additional information was sought from the notifying consultant by means of a questionnaire about severity, management, associations, previous reactions, and outcomes.

**RESULTS**

There were three deaths in the prospective period March 1998 to February 2000 and five in the retrospective period 1990 to February 1998 (table 1). Four cases arose in the searches by offices of national statistics and appeared in at least one of the other sources. There were also two cases from Dr Pumphrey, one from BPSU, and one from Asthma and Allergy Information Research, for which there was no other source. The under 16 population in 1999 was 13 028 933, based on data from the offices of national statistics. Thus, the period 1990–2000 yields an incidence of 0.006 deaths per 100 000 children per year. Table 2 shows the provoking allergens. Table 3 shows the association of the notified cases with a history of asthma. The more severe the reaction, the greater the chance the patient had coexisting asthma. Although the cases with non-severe reactions represent an unknown proportion of comparable reactions not admitted to hospital, we included them because coexisting asthma was so much less common. The fatal cases appeared to have especially troublesome asthma but we cannot be any more precise because historical accounts of antiasthma medication do not satisfactorily define severity. Table 4 shows the association with the severity of previous reactions. In 53% there had been a previous reaction and in 33% of these it was reported to be reported in error. Complete data were therefore available on 231 cases; 58 of these met the severity criteria.

There were 55 severe non-fatal reactions (21 female, 34 male) in the prospective period, six requiring intubation (near fatal). This yields an incidence of severe non-fatal events of 0.2 per 100 000 children per year. Table 2 shows the provoking allergens. Table 3 shows the association of the notified cases with a history of asthma. The more severe the reaction, the greater the chance the patient had coexisting asthma. Although the cases with non-severe reactions represent an unknown proportion of comparable reactions not admitted to hospital, we included them because coexisting asthma was so much less common. The fatal cases appeared to have especially troublesome asthma but we cannot be any more precise because historical accounts of antiasthma medication do not satisfactorily define severity. Table 4 shows the association with the severity of previous reactions. In 53% there had been a previous reaction and in 33% of these it was reported to be

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**Table 1** Deaths 1990–2000

<table>
<thead>
<tr>
<th>Year</th>
<th>Case</th>
<th>Age</th>
<th>Allergen</th>
</tr>
</thead>
<tbody>
<tr>
<td>1991</td>
<td>1</td>
<td>13 years</td>
<td>Milk</td>
</tr>
<tr>
<td>1992</td>
<td>2</td>
<td>15 years</td>
<td>Peanut</td>
</tr>
<tr>
<td>1994</td>
<td>3</td>
<td>3 months</td>
<td>Egg white</td>
</tr>
<tr>
<td>1994</td>
<td>4</td>
<td>9 years</td>
<td>Milk (in ice cream)</td>
</tr>
<tr>
<td>1995</td>
<td>5</td>
<td>13 years</td>
<td>Peanut</td>
</tr>
<tr>
<td>1998</td>
<td>6</td>
<td>13 years</td>
<td>Milk</td>
</tr>
<tr>
<td>1999</td>
<td>7</td>
<td>15 years</td>
<td>Milk</td>
</tr>
<tr>
<td>1999</td>
<td>8</td>
<td>5 years</td>
<td>Mixed food</td>
</tr>
</tbody>
</table>

**Table 2** Provoking allergens in the severe non-fatal food allergic reactions, 1998–2000

<table>
<thead>
<tr>
<th>Allergen</th>
<th>Severe</th>
<th>Near fatal</th>
</tr>
</thead>
<tbody>
<tr>
<td>W</td>
<td>3</td>
<td>2</td>
</tr>
<tr>
<td>Mixed food</td>
<td>9</td>
<td>2</td>
</tr>
<tr>
<td>Cows’ milk—including ice cream, yoghurt, formula milk</td>
<td>5</td>
<td>1</td>
</tr>
<tr>
<td>Egg</td>
<td>3</td>
<td>1</td>
</tr>
<tr>
<td>Peanuts</td>
<td>10</td>
<td>2</td>
</tr>
<tr>
<td>Yeast extract (Marmite)</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>Soya</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Sesame seed</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Potato</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Wheat</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Lentil</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Mixed nuts</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>Walnut</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Brazil nut</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>Cashew nuts</td>
<td>7</td>
<td></td>
</tr>
<tr>
<td>Macadamia nut</td>
<td>1</td>
<td></td>
</tr>
</tbody>
</table>

**Table 3** Association of asthma with severity of food allergic reaction, 1998–2000

<table>
<thead>
<tr>
<th>Reported reaction</th>
<th>Cases without asthma</th>
<th>Cases with asthma (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Non-severe</td>
<td>109</td>
<td>64 (37)</td>
</tr>
<tr>
<td>Severe</td>
<td>21</td>
<td>28 (57)</td>
</tr>
<tr>
<td>Near fatal</td>
<td>1</td>
<td>5 (83)</td>
</tr>
<tr>
<td>Fatal</td>
<td>0</td>
<td>3 (100)</td>
</tr>
</tbody>
</table>

χ² for trend 14.4, df=1, p=0.0002.

**Table 4** Association of severity of previous reaction with severity of food allergic reaction, 1998–2000

<table>
<thead>
<tr>
<th>Previous reaction</th>
<th>Non-severe</th>
<th>Uncertain</th>
<th>None</th>
<th>Mild/moderate</th>
<th>Hospitalised</th>
</tr>
</thead>
<tbody>
<tr>
<td>Non-severe</td>
<td>11</td>
<td>77</td>
<td>74</td>
<td>11</td>
<td></td>
</tr>
<tr>
<td>Severe</td>
<td>4</td>
<td>15</td>
<td>22</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>Near fatal</td>
<td>1</td>
<td>1</td>
<td>3</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Fatal</td>
<td>0</td>
<td>0</td>
<td>2</td>
<td>1</td>
<td></td>
</tr>
</tbody>
</table>

χ² for trend comparing mild/moderate with hospitalised: p>0.05.
to the same allergen, although the allergen judged responsible after a meal of mixed foods may be influenced by the previous history.

**DISCUSSION**

This first large child population based study of fatal and severe reactions to food covered a 0–15 years childhood population of 13 million and yielded a low incidence of 0.006 fatal events per 100 000 children per year over the period 1990–2000. Such deaths are much less common than sudden unexpected non-violent deaths from any cause in childhood which are about 3 per 100 000 per year. If 5% of the child population have food allergy, the risk that a food allergic child will die from a food allergic reaction is 1 in 800 000 per year.

The accuracy of death certificates pertaining to food allergy has been criticised. Furthermore, a postmortem diagnosis of allergy may be difficult, and a diagnosis of asthma (a “natural cause”) may be easier for a family to cope with than a diagnosis of allergy (an “unnatural cause” requiring an inquest). Nevertheless, we think cases are likely to have been identified because of the multiple sources we have used and the fact that the tragedy of a child’s death always leads to careful inquiry.

The finding that peanut was the cause of two of eight fatal reactions and no near fatal reactions differs from studies that report peanut to be the most common cause of fatal reactions to food in children in the United States. Additionally, there were no deaths from peanut allergy in children under 13. Milk caused the greatest number of fatal reactions (four of eight); this is in line with reports of the frequency and severity of reactions to milk.

In the prospective period, the reason for notification was that the child was admitted to hospital. The referring physician was not asked to assess severity. The severity criteria, deliberately set so that all children meeting the criteria would have had to be admitted to hospital, were based on clear treatment decisions that could be recorded. The criteria were applied by us, based on the answers in the questionnaire completed by the notifying physician. Any inaccuracy in the number of severe reactions so classified will be an overestimate because initial under treatment demands further treatment, whereas initial over treatment would not be noticed.

The incidence of non-fatal severe events was 0.2 per 100 000 children per year. The allergens responsible were similar to those reported in other series, the commonest being peanut (18%), albeit less common than previously reported.

Those suffering the most severe reactions tended to have had severe previous reactions, but it is notable that in two of the three fatal reactions and five of the six near fatal reactions, the previous event had not required urgent hospital treatment. This challenges the view that a previous mild reaction makes a fatal reaction less likely. All three children had eaten food shortly before the reaction. In one, the allergen to which the child was known to be allergic was not ingested; in the second, although milk protein may have been ingested, there was a concurrent and intense exposure to grass pollen; and in the third there was concurrent direct contact with a stuffed animal and the only previous reaction had been to an unknown allergen. These cases illustrate a group of asthmatic deaths where food may be initially implicated but subsequently shown to be unlikely.

On the other hand, could an apparently idiopathic attack of asthma be a manifestation of an unrecognised food allergic reaction? Our definition of a case excluded children whose only symptoms were asthmatic (no rash or oedema) and the allergen unknown because there is no means for attributing such reactions to food or for knowing if a causal link exists.

**Children who received a single dose of epinephrine**

Our criteria for a severe reaction excluded children who received only one dose of epinephrine and fulfilled none of the other criteria. There were 78 such excluded children but they cannot be all such cases, as some may not be admitted to hospital as recommended for observation for late or biphasic reactions. In 27 of the 78 (34%), no respiratory symptoms were noted before epinephrine administration; in 14 of these cases only mild cutaneous symptoms were seen. This suggests that some epinephrine is being administered unnecessarily. However, in the other instances, there were worrying symptoms that required urgent referral to hospital. Our study cannot determine whether epinephrine prevented potentially fatal occurrences in some of these.

Two of the three children in the prospective period who died received epinephrine before hospital. All the children who died had asthma and the precipitation of a severe asthmatic attack may have led to death. Administration of salbutamol by inhalation, together with intramuscular epinephrine and rapid transfer to hospital, might then better emergency treatment than epinephrine alone.

Treatment with epinephrine carries some risk of cardiac arrhythmia, even when given in recommended doses, and can cause death if given inadvertently in overdose. One fatality reported to us, which we have not classified as a food allergic death, was in a child with a mild food allergic reaction who died from overdose of intravenous epinephrine (coroner’s verdict).

**Conclusion**

The finding of so few deaths in such a large population should reassure parents and doctors that the risk of death is small. The findings are especially relevant to children under 10, in whom the risk of severe or fatal reaction is even smaller than in the 10–15 year old group, especially as there is increasing evidence that some young children grow out of their allergy by age 10. The child with food allergy and troublesome asthma may be at particular risk, and their asthma should be kept under optimal control. While a previous mild reaction may not be as reassuring as has been thought, absence of asthma may be. Early administration of epinephrine may not prevent death and concomitant treatment for the asthmatic component of an allergic reaction may be very important.

We emphasise that our findings relate to children, not adults; and to food, not other allergens.

**ACKNOWLEDGEMENTS**

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AC had the original idea for the study, and with CM, coordinated the study. All three authors were involved in study design, analysis, and writing of the paper.

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