Adverse reactions to food: Navigating the maze in primary health care

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Abstract
Primary health care providers are often confronted with patients describing adverse reactions to food. With a multiplex of differential diagnoses, it is essential to have a structured approach to these complaints. Just as allergies can have a negative impact on quality of life, inappropriate food allergy labelling can cause unnecessary distress and hardship for patients. Understanding the diagnostic approaches to adverse food reactions, alongside an appreciation of multidisciplinary collaboration, can assist the clinician in making appropriate decisions in the management of these patients.

Keywords: food allergy; food hypersensitivity; food intolerance; allergy-focused history; allergy diagnosis.

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Introduction
An adverse food reaction is defined as an abnormal response linked to the ingestion of food and can be broadly classified into immune-modulated reactions and non-immune-modulated reactions (food intolerances) as demonstrated in Figure 1.1 It is a common complaint, affecting 15%–20% of the global population, with multisystem involvement, ranging from gut to mucocutaneous, respiratory, cardiovascular and central nervous system symptoms.2 In addition, patients and caregivers may suffer from anxiety and reduced quality of life related to attending or being excluded from social events, bullying at school, the affordability of special or ‘free-from’ products and the extra time spent in-store to read food product labels. Fear of recurrence after events such as anaphylaxis may also lead to conditions such as post-traumatic stress reactions.3

Whilst the majority of adverse reactions to food can be attributed to non-allergic processes,4 patients may label their reactions as food allergy (FA). This self-reported diagnosis may lead to an overestimation as much as four times higher than the true prevalence,5 which may lead to self-imposed food avoidance6 and a risk of nutritional deficiencies.7

Food allergy is an adverse food reaction associated with a reproducible hypersensitive immune response.8 Despite beliefs that FAs are rare in Africa, evidence suggests that it is a growing problem9 and that it remains underdiagnosed.8 Studies have found that the true prevalence of immunoglobulin E (IgE)-mediated FA in children in South Africa varies from 0.5% in rural populations to 2.5% in urban populations,8 whilst the prevalence of IgE sensitisation to common foods is estimated at 11.6%.10 At present, non-IgE-mediated FAs (non-IgE FA) are less well studied than other food allergies.11 There is currently no prevalence data available for non-IgE FA in South Africa.

Figure 1: Classification of adverse food reactions.
Source: Adapted from Cox AL, Sicherer SH. Classification of adverse food reactions. J Food Allergy. 2020;2(1):3–6. https://doi.org/10.2500/jfa.2020.2.200022 IgE, immunoglobulin E.

It is the dichotomy of the increasing incidence of FA against the backdrop of self-reported overestimation that creates a diagnostic conundrum for primary health care providers. Considering that there are no validated tests for non-IgE-mediated allergies yet, clinicians may find themselves stuck in a maze of uncertainty.
The clinician should be able to discern a true FA from food intolerance and therefore provide relevant counselling or make an appropriate referral. Truly allergic individuals are at risk of developing life-threatening anaphylaxis and should therefore receive education on avoidance measures as well as the need for an allergy action plan and adrenaline autoinjector. Conversely, those without concrete evidence of harmful effects because of food ingestion should be discouraged from practising unnecessary avoidance.

It is therefore essential to have a structured approach to the diagnosis of an FA.

Diagnostic approach to food allergies

The diagnosis of FA may be seen as a process of triangulation involving sequential steps. One such model recommended by the European Academy of Allergy and Clinical Immunology supports a five-step approach that can be followed to confirm or exclude an FA:7

- Food allergy-focused history
- Selecting appropriate investigations to determine the presence of sensitisation
- Short-term elimination of suspected foods for diagnostic purposes
- Performing oral food challenges
- Evaluating for non-IgE-mediated FA.

Table 1: EATERS mnemonic.

| Focus         | IgE-mediated allergies                                                                 | Non-IgE-mediated allergies                      |
|---------------|----------------------------------------------------------------------------------------|-------------------------------------------------|
| Exposure      | Proximity to allergens is not deemed adequate to suggest exposure, the suspect food must be ingested. Other occasional methods for exposure could include handling raw food, being kissed by someone who has just eaten or inhalation of aerosolised pan-fried foods. | Ingestion of allergens via infant formula, food consumption or exposure through allergens excreted in breastmilk.8,9 |
| Allergen      | The most common allergens include milk, egg, nuts, soy, wheat and fish. The allergen is often determined by host immune responses. | Any food can cause a reaction. Cow’s milk, soy and egg are often implicated. More common foods include corn, wheat, rice, oat, barley, sweet potato, white potato, chicken, vegetables, fruit, peanuts, nuts, fish and shellfish.8,9 |
| Timing        | Most reactions occur immediately on exposure. Symptoms are often delayed and typically occur within 72 hours after ingestion. | More commonly seen in young infants and children. May occur during breastfeeding, formula feeding or during the introduction of solid foods. |
| Environment   | Weaning is the most common scenario as new foods are introduced to the diet. Food allergy also tends to occur when eating away from home – nurseries, restaurants, parties and holidays. | |
| Reproducible  | Strictly reproducible and occur at all subsequent exposures to the food. Strictly reproducible, and symptoms may occur with small volumes of the implicated food. | |
| Symptoms      | Skin: hives, swelling, redness, itching.GIT: vomiting, abdominal pain, diarrhoea. Respiratory tract: cough, wheeze, stridor, hoarseness, hypoxaemia. Cardiovascular: shock, hypotension, syncope, collapse | Skin: pruritus, erythema, atopic eczema. GIT: acute or chronic diarrhoea, gastrolesophageal reflux, loose or frequent stools, constipation, food aversion or refusal, abdominal pain, severe or recurrent vomiting, intermittent bloody stools or mucous in stools, dysphagia, oesophageal food impaction, erythema around the anus, malabsorption. Others: failure to thrive, changed behaviour because of discomfort and pain, persistent clear rhinorhoea, lethargy, possible cyanosis, repetitive emesis may be associated with progressive lethargy, shock, dehydration and acidosis, hyponatraemia and hypotension. |

Source: Adapted from Eleywun-Lajeunesse M, Weir T, Brown L, et al. Fifteen-minute consultation: The EATERS method for the diagnosis of food allergies. Arch Dis Child Educ Pract. 2019;104(6):286–291. https://doi.org/10.1136/archdischild-2018-316397

Step 1: Food allergy-focused history taking

An allergy-focused clinical history remains the cornerstone of any investigative process. Ignoring this step in favour of other uninformed investigations will likely create further diagnostic confusion. The EATERS (exposure, allergen, timing, environment, reproducible, symptoms) mnemonic8 provides a method for clinicians to interpret each case and makes sense of test results (Table I).

Some caveats may not fit in with the classical history of FA. For instance, food-dependent, exercise-induced anaphylaxis occurs when food and exercise are tolerated independently, but reactions occur when physical exertion follows ingestion of a food allergen.15 Secondly, a condition called alpha-gal syndrome presents a true IgE-mediated allergy to red meat, but the timing of symptoms is delayed by 2–6 hours,16 resulting in the phrase ‘midnight anaphylaxis’.17 These examples provide the potential limitations of an allergy-focused history and further demonstrate the fact that diagnosis in allergic conditions is a process rather than a single investigation.

Step 2: Investigations to confirm sensitisation

Immunoglobulin E (IgE) sensitisation can be determined via skin prick testing (SPT) or specific IgE levels. It is important to remember that sensitisation may be used as a proxy for confirming FA in the context of a positive history.7 Food-specific SPT and IgE have a sensitivity of 70%–100%, but a specificity of 40%–70%. Therefore,
Step 4: Oral food challenges

An oral food challenge is the gold standard for the diagnosis of FA. However, it is mostly reserved for scenarios where further clarification is required (e.g., in the setting of a suggestive history but equivocal SPT or IgE results), as it is time-consuming and carries a risk of inducing a severe allergic reaction. It should therefore be done under the supervision of a trained clinician.

Step 5: Evaluation for non-IgE food allergy

Overview of non-IgE-mediated food allergy

Non-IgE FAs are characterised by gastrointestinal reactions, including food protein-induced allergic proctocolitis (FPIAP), food protein-induced enterocolitis syndrome (FPIES) and food protein-induced enteropathy (FPE) (see Box 1 for disease specific characteristics); cutaneous reactions (such as atopic dermatitis, contact dermatitis and dermatitis herpetiformis) and respiratory reactions (such as Heiner syndrome) are often reported. Eosinophilic gastrointestinal disorders (EGIDs), including eosinophilic esophagitis (EoE), allergic eosinophilic gastroenteritis (AEG) and eosinophilic colitis (EC), are also classified as non-IgE FAs. Eosinophilic gastrointestinal disorders often present with non-specific symptoms such as repeated regurgitation, vomiting, and watery, mucoid or haemorrhagic diarrhoea, combined with other features such as poor growth and crying crises (colic). Food allergy itself can also cause gastrointestinal reflux disease.

Non-IgE-mediated food allergic gastrointestinal disorders, including gastroesophageal reflux, abdominal pain, constipation and frequent vomiting, can adversely affect a child’s feeding ability and skills, depending on the associated symptoms, disease severity, age at which the child is affected, social circumstances and the degree of dietary restriction recommended. Early diagnosis and management of these feeding difficulties are of utmost importance, as they can result in nutritional deficiencies or growth faltering. Growth faltering is a commonly described symptom, with low weight and in particular low height found in approximately 10% of children with non-IgE FA. Cow’s milk is a predominant FA in children, of which non-IgE gastrointestinal FA accounts for up to 50% of reactions. Similarly, cow’s milk is responsible for the majority of reactions in FPIES, FPE and FPIAP.

Diagnostic approach to non-IgE-mediated food allergy

With the lack of laboratory tests to support the diagnosis of non-IgE FAs, the diagnosis remains challenging, as it relies on clinical evaluation and sound history taking (refer to Table I). The onset of non-IgE FA symptoms is delayed and may have a chronic presentation, making their association with the food allergen less evident. The diagnosis of non-IgE FAs is based on the occurrence of symptoms after the culprit allergen is consumed, followed by their resolution once the suspected food has been eliminated and reappearing when it is re-introduced.

Multidisciplinary collaboration: The role of the registered dietitian

A targeted, individualised approach to each allergic patient remains essential for the effective management of FA. As part of a thorough nutritional assessment, an experienced dietitian will conduct an allergy-focused clinical history and a comprehensive dietary assessment, including the evaluation of the food and symptom diary, as well as monitor growth. Dietary management includes advice regarding allergen avoidance, food label ingredients and terminology, hidden allergens in foods, suitable replacement foods and ensuring dietary adequacy. Ideally, no food should be
removed from an individual’s diet without the involvement of a dietitian. A dietitian can help with the identification of possible allergen–symptom associations and offer the necessary support to improve the allergic individual’s quality of life.

**Summary**

Adverse reactions to food are commonly seen in primary health care providers. In addition to the obvious physical effects, it also has negative psychological and social consequences. Given the variety of causes and presentations, clinicians may easily be confused, resulting in haphazard testing and counselling.

Central to the accurate diagnosis of an FA is a sound allergy-focused clinical history. Specific information regarding the presenting symptoms should include the following: the age of symptom onset; the timing of onset following food contact; the duration, severity and frequency of symptoms; and the quantity of food required to induce a reaction, as is the reproducibility of symptoms on re-exposure. Other pertinent information includes personal history of atopic disease (other FA, asthma, eczema or allergic rhinitis); family history of atopic disease in parents or siblings; cultural and religious factors that affect food preferences; comprehensive dietary assessment, including the age of weaning and food preferences; and whether breastfed, mixed or formulated. If the patient is breastfed, enquire about maternal dietary intake; details of any previous treatment, including medication such as antihistamines; and details on food elimination and reintroduction which may guide further investigation and management. A ‘14-day food and symptom diary’ is a valuable reintroduction which may guide further investigation and intake; details of any previous treatment, including medication such as antihistamines; and details on food elimination and reintroduction which may guide further investigation and management. A ‘14-day food and symptom diary’ is a valuable diagnostic aide.

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J.S. and I.v.Z. contributed equally to the design and the writing of the article.

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