

# Critical Issues in Food Allergy: A National Academies Consensus Report

Scott H. Sicherer, MD,<sup>a</sup> Katrina Allen, MD, PhD,<sup>b,c</sup> Gideon Lack, MD,<sup>d,e</sup> Steve L. Taylor, PhD,<sup>f</sup> Sharon M. Donovan, PhD, RD,<sup>g</sup> Maria Oria, PhD<sup>h</sup>

The National Academies of Sciences, Engineering, and Medicine convened an expert, ad hoc committee to examine critical issues related to food allergy. The authors of the resulting report, “Finding a Path to Safety in Food Allergy: Assessment of the Global Burden, Causes, Prevention, Management, and Public Policy,” evaluated the scientific evidence on the prevalence, diagnosis, prevention, and management of food allergy and made recommendations to bring about a safe environment for those affected. The committee recommended approaches to monitor prevalence, explore risk factors, improve diagnosis, and provide evidence-based health care. Regarding diagnostics, emphasis was placed on utilizing allergy tests judiciously in the context of the medical history because positive test results are not, in isolation, diagnostic. Evidence-based prevention strategies were advised (for example, a strategy to prevent peanut allergy through early dietary introduction). The report encourages improved education of stakeholders for recognizing and managing as well as preventing allergic reactions, including an emphasis on using intramuscular epinephrine promptly to treat anaphylaxis. The report recommends improved food allergen labeling and evaluation of the need for epinephrine autoinjectors with a dosage appropriate for infants. The committee recommended policies and guidelines to prevent and treat food allergic reactions in a various settings and suggested research priorities to address key questions about diagnostics, mechanisms, risk determinants, and management. Identifying safe and effective therapies is the ultimate goal. This article summarizes the key findings from the report and emphasizes recommendations for actions that are applicable to pediatricians and to the American Academy of Pediatrics.

Food allergy has become an increasingly recognized global health concern. Defined as an adverse health effect arising from a specific immune response that occurs reproducibly on exposure to a given food,<sup>1</sup> the disease impacts health and quality of life for sufferers and their caregivers.<sup>2</sup> A new report entitled “Finding a Path to Safety in Food Allergy: Assessment of the Global Burden, Causes, Prevention, Management, and Public Policy” (referred to hereafter as “the

report”) was recently released by The National Academies of Sciences, Engineering and Medicine (NAS) (available at [www.nationalacademies.org/FoodAllergies](http://www.nationalacademies.org/FoodAllergies)).<sup>3</sup> Here we describe the Report, focusing on key findings and recommendations applicable to pediatricians and the American Academy of Pediatrics (AAP).

The National Academy of Sciences was established in 1863 by an Act of Congress signed by President Abraham Lincoln as a nongovernmental

## abstract

FREE

<sup>a</sup>Division of Pediatric Allergy and Immunology, Department of Pediatrics, Jaffe Food Allergy Institute, Icahn School of Medicine at Mount Sinai, New York, New York; <sup>b</sup>Centre for Food and Allergy Research, Murdoch Children's Research Institute, The University of Melbourne, Victoria, Australia; <sup>c</sup>Institute of Inflammation and Repair, University of Manchester, Manchester, United Kingdom; <sup>d</sup>Division of Asthma, Allergy and Lung Biology, Department of Paediatric Allergy, King's College London, London, United Kingdom; <sup>e</sup>Guy's and St Thomas' NHS Foundation Trust, London, United Kingdom; <sup>f</sup>Food Allergy Research and Resource Program, Department of Food Science and Technology, University of Nebraska, Lincoln, Nebraska; <sup>g</sup>Department of Food Science and Human Nutrition, University of Illinois, Urbana, Illinois; and <sup>h</sup>National Academies of Sciences, Engineering and Medicine, Washington, DC

Dr Sicherer authored the initial draft of this manuscript; Drs Allen, Lack, Taylor, Donovan, and Oria contributed additional perspectives and edited subsequent drafts of this manuscript; and all authors approved the final manuscript as submitted and agree to be accountable for all aspects of the work.

The authors are responsible for the content of this article, which does not necessarily represent the views of the National Academies of Sciences, Engineering, and Medicine, their committees, or convening bodies.

**DOI:** <https://doi.org/10.1542/peds.2017-0194>

Accepted for publication May 1, 2017

Address correspondence to Scott H. Sicherer, MD, Mount Sinai Hospital, Box 1198, 1 Gustave L Levy Place, New York, NY 10029. E-mail: [scott.sicherer@mssm.edu](mailto:scott.sicherer@mssm.edu)

PEDIATRICS (ISSN Numbers: Print, 0031-4005; Online, 1098-4275).

Copyright © 2017 by the American Academy of Pediatrics

**To cite:** Sicherer SH, Allen K, Lack G, et al. Critical Issues in Food Allergy: A National Academies Consensus Report. *Pediatrics*. 2017;140(2):e20170194

institution to advise the nation and, in concert with the National Academy of Engineering and the National Academy of Medicine (formerly the Institute of Medicine), now functions as the NAS to provide independent, objective advice to the nation. For the study on food allergy, support was received by 3 federal and 8 nonfederal sponsors\* and was the product of a study by a committee of 15 international experts with diverse expertise who were vetted for conflicts of interest and who considered a vast array of issues in the field. The evidence base included selective literature reviews and evaluation of published guidelines and practice parameters, and it favored meta-analyses and systematic reviews as applicable to a variety of topics. Recognizing that the etiology and management of food allergy involves not only the patient, an ecological-developmental model of food allergy was considered to address numerous interacting facets including the environment, family, home, child care, the health care system, industry, cultural practices, and government policy and regulation.

The report underscores the central role of the immune response in defining food allergy, thus distinguishing it from other adverse

\*As indicated in the full report, the National Academies of Sciences, Engineering and Medicine activity was supported by federal sponsors: the Food and Drug Administration (contract HHSP233201400020B/HHSP23337025), the Food and Nutrition Service of the US Department of Agriculture (grant # FS\_NAS\_IOM\_FY2015\_01), and the National Institute of Allergy and Infectious Diseases; and nonfederal sponsors: the Asthma and Allergy Foundation of America, the Egg Nutrition Center, Food Allergy Research & Education, the International Life Sciences Institute North America, the International Tree Nut Council Nutrition Research & Education Foundation, the National Dairy Council, the National Peanut Board, and the Seafood Industry Research Fund. The full report indicates that any opinions, findings, conclusions, or recommendations expressed do not necessarily reflect the views of any organization or agency that provided support for the project.

responses to food. Food allergy results from immunoglobulin E (IgE)-mediated and/or non-IgE-mediated mechanisms, the former being more common (potentially resulting in anaphylaxis), and is the focus of the report. Often, alternative causes of adverse reactions to foods may be misinterpreted as a food allergy. Lactose intolerance is one example; the inability to digest the sugar lactose results in bloating and diarrhea, but is not an allergy. Adverse reactions to foods may be caused by metabolic, pharmacologic, or toxic factors that must be distinguished from allergy. In the United States, it is common for parents to avoid a food on the basis of the perception of food allergy when in fact most of the time diagnostic testing will reveal that there is none and the food could be added back to the diet.<sup>1,4,5</sup> Here we review and comment on highlights of the report that are pertinent to pediatricians, pediatric care of food allergies, and the AAP as a professional organization named as a potential actor to accomplish some of the committee's recommendations.

## PREVALENCE

A previous US review concluded that food allergy affects >1% to 2% and <10% of the population.<sup>6</sup> A US study published in *Pediatrics* that was based on parental report of allergy concluded that up to 8% of children have food allergy.<sup>7</sup> Several studies, including data presented by the Centers for Disease Control and Prevention (CDC), suggest an increase in prevalence in the United States.<sup>8,9</sup> However, the report points out that prevalence estimates vary among studies depending on numerous factors, including the definition of allergy; the foods being considered; whether diagnosis is by self-report, testing, or a combination of factors; selection of participants; age of participants;

geographical regions; and many other factors. Ideal studies include medically supervised feeding to confirm allergy or tolerance (an oral food challenge [OFC]). Studies of self-reported allergy typically overestimate prevalence.<sup>10</sup> Few studies include gold standard, medically supervised feeding tests, with no US studies including them since the 1980s.<sup>5</sup> Although many foods have been noted to cause allergic reactions, it is clear that cow's milk, hen's egg, peanuts, tree nuts, and seafood are responsible for most of the serious allergic reactions.<sup>1,11</sup> Despite an exhaustive literature search, the authors of the report could not present definitive prevalence data for the United States. The committee recommended approaches for performing studies to obtain more accurate prevalence estimates, including using modalities such as the National Health and Nutrition Examination Survey with incorporation of additional testing of subsamples with OFCs.

## DIAGNOSIS AND PROGNOSIS

The committee recommended that "physicians use evidence-based, standardized procedures as the basis for food allergy diagnosis and avoid nonstandardized and unproven procedures....When food allergy is suspected, the patient should be evaluated by a physician who has the training and experience to select and interpret appropriate diagnostic tests." The AAP's Clinical Report on allergy testing emphasizes these fundamentals<sup>12</sup> and, like the National Academies report, suggests consideration of referral to or consultation with a physician specialist (for example, an allergist-immunologist).

The basis of these comments stems in part from the observation that self-reported allergy typically overestimates true allergy, and no simple tests provide a diagnosis.

In one meta-analysis, the rate of self-reported food allergy among children was 12%, compared with 3% when confirmatory testing was performed.<sup>10</sup> A serious misconception about food allergy diagnostics relates to equating a “positive test result” by a serum food-specific IgE (sIgE) blood test or skin prick test (SPT) to having an allergy to the tested food. These tests detect IgE antibodies to the food but are not typically intrinsically diagnostic. For example, in 111 OFCs performed in 44 children avoiding foods because of positive test results, 93% were tolerant of the avoided food.<sup>13</sup> It is clear that these tests are misunderstood by physicians. For example, in a survey of 407 primary care physicians, 38% indicated incorrectly that these sIgE blood tests or SPTs were sufficient for diagnosis.<sup>14</sup> Although overdiagnosis is a concern and may be driven by overtesting and misinterpretation of results, under- or misdiagnosis must also be addressed. Assuming incorrectly that an allergen has been identified without confirmatory testing could lead to reexposure to the true culprit, resulting in serious reactions if the unidentified trigger is consumed again. The report summarized information from previously published guidelines and systematic reviews to highlight appropriate tests and their use and to discuss tests that are not recommended.<sup>1,15,16</sup>

A previous guideline<sup>1</sup> suggests that the medical history is key in diagnosis, and food allergy should be considered when allergic symptoms occur proximate (within minutes to hours) to ingestion of a specific food, especially when symptoms occur on more than 1 occasion. For example, suspicion would be high if a child developed urticaria within minutes after the ingestion of a peanut, especially if an acute reaction such as this occurred with more than 1 exposure. However, suspicion might

be low if the child routinely ingested peanuts and the urticaria persisted for days, suggesting a viral rash. Food allergy diagnostic testing may also be warranted for infants and children with moderate to severe atopic dermatitis because there is a higher rate of food allergy in this population, and the food allergy may be contributing to the rash. Disorders with subacute or chronic symptoms that are indicative of a food-related origin, such as food protein–induced enterocolitis syndrome (severe vomiting, lethargy, and sometimes hypotension and acidosis starting 2 hours after ingestion) and allergic colitis (mucous-containing, bloody stools) also warrant investigation for food-allergic triggers, but these disorders occur without IgE antibodies.<sup>1</sup> Food allergy should also be considered in children with eosinophilic esophagitis. Importantly, food allergy is not a typical trigger of chronic asthma or chronic rhinitis in childhood.

Medical history can help to identify the likelihood of a food allergy diagnosis (pretest probability) and can suggest whether the pathophysiology is IgE- or non-IgE-mediated, which is pertinent for test selection and helps to identify potential triggers. Importantly, details of the history may disclose alternative reasons for symptoms other than a food allergy. Additional diagnostic modalities include diagnostic elimination diets, SPT, sIgE blood test, and OFC. Neither SPT nor sIgE blood test results alone are considered sufficient for diagnosis. However, in the presence of a strong suggestive history of food allergy, these can confirm the diagnosis without further need for an OFC.<sup>1,12</sup> Increasingly large skin test wheal sizes or sIgE levels correlate with higher risks of allergy, but sensitivity and specificity are often inadequate to confirm a diagnosis.<sup>16</sup> Tests that measure sIgE to specific proteins in foods, a modality called “component

testing,” may improve specificity of tests. For example, Ara h 2 is a protein in peanuts associated with clinical reactions.<sup>15,17</sup> The OFC, particularly when performed in a double-blind manner with a placebo, is considered a gold standard.<sup>18</sup> Unmasked OFCs are typically performed for clinical purposes, and double-blind procedures are used more often for research. The report concluded that a number of diagnostic modalities were not recommended for routine use, including food allergy patch testing (atopy patch test), measurement of total IgE, and the basophil activation test. Other tests were not recommended and were considered “unproven and non-standardized” for diagnosing food allergy. These tests include allergen-specific IgA, IgG, or IgG<sub>4</sub>; provocation neutralization; immune complexes; human leukocyte antigen screening; lymphocyte stimulation; facial thermography; gastric juice analysis; endoscopic allergen provocation; hair analysis; applied kinesiology; cytotoxic assays; electrodermal testing; mediator release assays; bioresonance; and iridology.

Although some studies suggest laboratory results might help to provide insights on severity and prognosis,<sup>19–23</sup> there are no simple reliable means currently to identify these aspects of food allergy. The report emphasizes that on the basis of test limitations, particularly that sensitization alone is not diagnostic, physicians should not order “panels” of food tests without a rationale.

## RISK AND PREVENTION

Food allergy is undoubtedly caused by a combination of genetic and environmental factors.<sup>24</sup> The report presents a number of risk factors and prevention strategies that have been evaluated in published studies.

Regarding solid food introduction, the authors of the report concluded that

there is strong evidence regarding the early introduction of peanuts as being protective against peanut allergy in infants at high risk, defined by early-onset eczema or coexistent egg allergy. This is generally in line with recent consensus publications and guidelines<sup>25,26</sup> that have been endorsed by the AAP and whose authors have promoted the introduction of peanut protein in infant-safe forms for high-risk infants as early as 4 to 6 months of age after appropriate testing. Regarding foods other than peanuts, the report indicates that more studies are needed to assess whether early introduction affects food allergy. However, the report concludes that “limited evidence... suggests that delaying the introduction of egg, cow milk and wheat to decrease risk of those food allergies has no benefits” and discusses a potential benefit of introducing these foods in the first year of life when the infant is developmentally ready (at ~6 months but not before 4 months of age).

The committee also concluded that there is limited evidence to support or discourage eliminating allergenic foods from the diet of pregnant or lactating women and that evidence for a protective effect of breastfeeding is also limited.

There were no changes in recommending breastfeeding as the preferred feeding for all infants. However, the authors of the report gave recommendations and drew conclusions that differ from findings and recommendations on allergy prevention that were published in a 2008 AAP Clinical Report.<sup>27</sup> The authors of the AAP Clinical Report suggested that specific types of infant formulas may be useful for allergy prevention in high-risk infants if the infants could not exclusively be breastfed. In contrast, the NAS committee concluded that studies on the effects of partially or extensively hydrolyzed infant formulas for preventing food allergies are

inconsistent or have methodologic flaws and that evidence is therefore limited. They concluded that high-quality randomized clinical studies would be needed before these formulas could be recommended for food allergy prevention.

As a global recommendation, the committee advised that public health authorities and clinical practice guidelines include consistent, clear, and evidence-based advice for families and health care providers about the potential benefits of introducing allergenic foods in the first year of life to infants when the infant is developmentally ready (at ~6 months of age but not before 4 months of age, particularly for those at high risk of allergy). As noted above, the AAP has been involved in updating such guidelines<sup>26</sup> and is in the process of creating an updated Clinical Report on prevention of atopic disease.

Regarding risk factors, the committee identified limited but consistent evidence that skin barrier defects (for example, eczema or filaggrin gene mutations) play a role in sensitization. Although individuals with 1 type of atopy (eg, eczema) may be at risk for another manifestation (eg, food allergy), a new theory suggests that skin barrier defects and inflammation may provide a sensitizing route of exposure to proteins in the environment. Unfortunately, firm conclusions could most often not be reached on a number of issues because of limited data. The committee considered, but could not identify because of limited evidence, a link between food allergy and nutritional factors such as vitamin D, maternal omega-3 fatty acid intake, folate supplementation, and other nutrients. The committee could not identify randomized clinical trials or other convincing evidence to address other potential hypotheses for an increase in food allergy. Only a few studies have been published on the relationship between changes in the microbiota and food allergen

sensitization, so evidence supporting the relationship is limited. Trials on probiotics and prebiotics also have methodologic limitations, and the committee concluded that there is not yet evidence to support a decreased risk of food allergy from using these interventions. Also with regard to microbial exposure and the “hygiene hypothesis,” firm conclusions could not be reached regarding a relationship between food allergy and cesarean delivery, use of antibiotics, or exposure to pets or animals. Although there is indirect evidence that genetics contributes to the development of food allergies, with those with a family history at moderately increased risk, there are no conclusive findings regarding specific loci. Similarly, epigenetic mechanisms are likely involved because the apparent rise in food allergy has occurred in a short period of time and therefore is presumed to be secondary to environmental causes. These hypotheses are underpinned by a recent finding that Asian infants born in Asia who migrate to Australia in the first 5 years of life are protected from nut allergy, whereas Australian-born Asian infants are at a 2 to 3 times higher risk of nut allergy than white infants born in Australia when evaluated for allergy at age 5 years.<sup>28</sup>

## EMERGENCY MANAGEMENT

The authors of the report reviewed the management of food allergic reactions and anaphylaxis and focuses on prompt use of epinephrine to treat anaphylaxis. Researchers suggest that epinephrine is safe but often underutilized<sup>29–31</sup> and that there is poor recognition about both how and when to use epinephrine autoinjectors. The authors of the report identified teenagers as a high-risk group because they may take risks that result in ingestion of an allergen and may postpone proper treatment. In addition, those with

**TABLE 1** Recommendations Regarding Management Directed to Pediatricians and/or the AAP

Recommendations From the Report Applicable to Pediatricians at the Patient-Care Level or Addressable by the AAP	Features, AAP Resources, Current Actions, Potential Future Work
The AAP should regularly update guidelines on diagnosis, prevention, and management of food allergy on the basis of strong scientific evidence as emerging scientific data become available.	The AAP has a number of Clinical Reports on food allergy diagnosis, management of food anaphylaxis, and food allergy management in schools. <sup>12,27,32,33,37</sup> It also participated in development of guidelines. <sup>26</sup> Participation in updating and developing resources is needed.
Health care providers should receive training on approaches to counseling patients and their caregivers. Counseling training is envisioned to be provided in part by professional organizations through various means, including the Internet.	In addition to the guidelines mentioned above, the AAP has educational modules on selected topics including programming at the annual National Conference and Exhibition. More information and materials to fulfill these recommendations are still needed.
Health care providers should counsel patients and their caregivers on food allergies by following the most recent guidelines emphasizing the need to take age-appropriate responsibility. Counseling is particularly important for those at high risk and with severe food allergy, such as adolescents and those with both food allergy and asthma.	The areas of counseling include emergency management, avoidance in a variety of settings, nutritional counseling, and psychosocial counseling.
Health care providers and others should use intramuscular epinephrine as a first line of emergency management for episodes of food allergy anaphylaxis.	The AAP recently released 2 Clinical Reports on these topics, including information regarding the use of epinephrine for first aid management in the community and for providing written plans for recognition and treatment of food allergy and anaphylaxis. <sup>32,33</sup>

food allergy and comorbid asthma are considered at high risk because they may be prone to more severe reactions. Epinephrine autoinjectors are currently only available in doses of 0.15 and 0.30 mg, which are not ideal for infants. The committee recommended that the Food and Drug Administration (FDA) evaluate the need for an epinephrine autoinjector with a dosage appropriate for use in infants and, if indicated, that industry should develop an autoinjector with 0.075 mg of epinephrine specifically designed for use in infants. The authors of an AAP Clinical Report discuss the dosing of epinephrine autoinjectors for first aid management of anaphylaxis in the community and also raise concerns regarding the best dosing for infants.<sup>32,33</sup>

### DAILY MANAGEMENT

Pediatricians should counsel families on allergen avoidance for a variety of settings, including home, school,

and travel. At home, care is needed during food preparation to avoid cross-contact of allergen with safe foods. Information about obtaining safely packaged foods and managing food allergy outside the home is reviewed below. Two additional areas of emphasis especially pertinent to pediatric patients are (1) the need for careful nutritional monitoring and intervention for children avoiding foods because of allergy<sup>34</sup>; and (2) attention to psychosocial aspects of managing food allergy, including an increased risk of bullying.<sup>35,36</sup> The committee recommended that the CDC work with other public health authorities to initiate a public health campaign for the general public, those with food allergy, and relevant stakeholders to increase awareness and empathy as well as to dispel misconceptions about food allergy and its management. This global approach could reduce bullying and risk-taking behavior by raising awareness. The committee made a number

of additional recommendations applicable to pediatricians and to the AAP as outlined in Table 1.

### Packaged Foods, Travel, Restaurants

Patients and families living with food allergies often depend on others to obtain safe foods. Obtaining safely packaged food is one consideration. The current status of labeling is often confusing for allergic consumers. In the United States, labeling laws require plain English terminology to identify “major allergens” as ingredients, including cow’s milk, hen’s egg, wheat, soy, peanuts, tree nuts, fish, and crustacean shellfish. For categorical foods, specific types must be listed (for example, codfish instead of “fish” or walnut instead of “nuts”). However, the foods considered “nuts” include ones that are not necessarily true nuts. The committee identified, for example, that lychee is erroneously considered a nut by the FDA (when botanically it is actually a fruit). Other allergens, such as sesame, are not included in US labeling laws and yet different countries include different foods in their labeling laws, including sesame and others. Advisory labeling, also called precautionary labeling (eg, “may contain” or “in a facility with”), is voluntary and not regulated. There has been a proliferation of advisory labeling, leading to consumer concerns about the veracity of the warnings. The committee provided a number of recommendations directed toward the FDA to improve labeling and create a system in which advisory labeling is meaningful. For example, evaluation of a risk-based labeling system like the Voluntary Incidental Trace Allergen Labelling program, which is used in Australia, could be considered.

Regarding restaurants and food service as well as foods prepared in groceries, the committee recommended a number of training approaches and regulatory actions to improve safety and the flow of information

for consumers. In the meantime, it is important for pediatricians to advise patients to inform restaurants and food service establishments of their allergy and to discuss details of the food's safety with staff. With regard to airline travel, where concerns include the safety of food provided and exposure to food proteins in the closed environment, recommendations are directed toward food safety and also anaphylaxis recognition and management. The recommendations include ensuring emergency epinephrine capabilities are in place for children. The AAP has been committed to these issues, and recently resolutions were submitted at the Annual Leadership Forum regarding availability of epinephrine for children on airplanes.

### **Schools and Early Child Care and Education Settings**

The committee recommended that various stakeholders, including advocates such as the AAP, participate in a task force to address emergency management and prevention strategies for venues such as schools and early care and education settings. The authors of the report review some of the approaches, including state and federal policies, which are important for managing food allergy in early care and education settings and schools. The AAP has been committed to these issues as well. The AAP produced a Clinical Report on food allergy management in schools<sup>37</sup> and had representation on the CDC Voluntary Guidelines for Food Allergy in Schools that was referred to in the report and was recommended for implementation. The CDC guidelines report is available at: [http://www.cdc.gov/healthyschools/foodallergies/pdf/13\\_243135\\_a\\_food\\_allergy\\_web\\_508.pdf](http://www.cdc.gov/healthyschools/foodallergies/pdf/13_243135_a_food_allergy_web_508.pdf).

### **RESEARCH NEEDS**

The committee identified numerous areas in need of additional research. Some of the priority areas include

the prevalence and cost of food allergies; diagnosis and prognosis; risk determinants and prevention; management in health care settings, food establishments, schools and traveling; and curative therapies. Some of the areas that may be of interest to general pediatric research include the best modalities to educate families and physicians about management, examining barriers to proper testing, identifying educational approaches and tools to improve physician and patient education, utilizing birth cohorts and other opportunities for prevalence estimates, determining the effectiveness of evidence-based guidelines and educational programs on food allergy management, improving the understanding of nutritional needs, establishing the best means to identify and manage psychosocial concerns, monitoring the number of food allergic reactions that occur in settings such as schools, and defining best practices regarding management.

### **SIX MAJOR ACTIONS REQUIRED TO ACHIEVE FOOD ALLERGY SAFETY**

The committee conceptualized a roadmap to safety involving 6 major actions that require input from various stakeholders, including pediatricians and professional organizations such as the AAP, to achieve short- and long-term goals. The first action would involve collecting better information about prevalence (which is crucial for identifying the scope of the problem), prioritizing food allergy in context of other health diseases, and exploring risk factors. The second action involves improving the quality of diagnosis and providing evidence-based health care. The third action concerns evidence-based prevention strategies. As noted previously, data are lacking on a variety of potential interventional strategies, but exciting new approaches, particularly regarding prevention of peanut allergy, are being promulgated. The fourth action is to improve education and training

of all stakeholders for recognizing and managing as well as preventing allergic reactions. This work includes updating, creating, and implementing various guidelines and educational programs. The fifth action is to develop policies and related practices to help prevent and treat severe reactions. The sixth and most critical action regards research priorities to address key questions about diagnostics, mechanisms, risk determinants, and management. Identifying safe and effective therapies is the ultimate goal.

### **CONCLUSIONS**

Food allergy appears to be increasing in prevalence and particularly affects children. The National Academies report identifies a large number of immediately obtainable practice goals to improve safety for children with food allergies. The committee also identified numerous research goals. Although the recommendations of the report are directed toward numerous stakeholders (including consumers, patients, allergy researchers, health care providers, school leaders, manufacturers, government agencies, and others), pediatricians and the AAP can take a prominent seat at the table to ensure that the report's smorgasbord of findings and recommendations are digested and implemented to reduce risks and improve safety for children with food allergies.

### **ABBREVIATIONS**

AAP: American Academy of Pediatrics  
CDC: Centers for Disease Control and Prevention  
FDA: Food and Drug Administration  
IgE: immunoglobulin E  
NAS: National Academies of Sciences, Engineering, and Medicine  
OFC: oral food challenge  
sIgE: serum food-specific IgE  
SPT: skin prick test

**FINANCIAL DISCLOSURE:** Dr Sicherer received grant support for research from the National Institute of Allergy and Infectious Diseases (NIAID) and Food Allergy Research and Education and royalties from UpToDate. Dr Donovan received grant support for pediatric nutrition research from the National Institutes of Health as well as food and pharmaceutical companies. Dr Taylor received grant support for research from the National Institute of Food and Agriculture and from the Food Allergy Research and Resource Program consortium of over 90 food manufacturers and/or suppliers as well as royalties from Neogen Corp. Drs Allen, Lack, and Oria have indicated they have no financial relationships relevant to this article to disclose.

**FUNDING:** No external funding.

**POTENTIAL CONFLICT OF INTEREST:** Drs Sicherer, Allen, Lack, Taylor, and Donovan served as members of the committee for the report discussed herein. Dr Oria was the study director for the report discussed herein. Dr Lack has received research support from the NIAID and the UK Food Standards Agency, is on the DBV Technologies scientific advisory board, has received a contribution to NIAID contract and/or grant from Food Allergy Research and Education, has received a contribution to King's College London Division of Asthma Allergy and Lung Biology from MRC & Asthma UK Centre, has received the Biomedical Research Centre award to Guy's and St Thomas' NHS Foundation from the UK Department of Health through the National Institute for Health Research, has received support for a pediatric allergy clinical trial's unit from the National Peanut Board, has received a discounted Bamba peanut snack from Osem, and has stock and/or stock options in DBV Technologies.

## REFERENCES

1. Boyce JA, Assa'ad A, Burks AW, et al; NIAID-Sponsored Expert Panel. Guidelines for the diagnosis and management of food allergy in the United States: summary of the NIAID-sponsored expert panel report. *J Allergy Clin Immunol*. 2010;126(6):1105–1118
2. Lieberman JA, Sicherer SH. Quality of life in food allergy. *Curr Opin Allergy Clin Immunol*. 2011;11(3):236–242
3. National Academies of Sciences, Engineering, and Medicine. *Finding a Path to Safety in Food Allergy: Assessment of Global Burden, Causes, Prevention, Management, and Public Policy*. Washington, DC: The National Academies Press; 2017
4. Burks AW, Jones SM, Boyce JA, et al. NIAID-sponsored 2010 guidelines for managing food allergy: applications in the pediatric population. *Pediatrics*. 2011;128(5):955–965
5. Bock SA. Prospective appraisal of complaints of adverse reactions to foods in children during the first 3 years of life. *Pediatrics*. 1987;79(5):683–688
6. Chafen JJ, Newberry SJ, Riedl MA, et al. Diagnosing and managing common food allergies: a systematic review. *JAMA*. 2010;303(18):1848–1856
7. Gupta RS, Springston EE, Warrier MR, et al. The prevalence, severity, and distribution of childhood food allergy in the United States. *Pediatrics*. 2011;128(1). Available at: [www.pediatrics.org/cgi/content/full/128/1/e9](http://www.pediatrics.org/cgi/content/full/128/1/e9)
8. Branum AM, Lukacs SL. Food allergy among children in the United States. *Pediatrics*. 2009;124(6):1549–1555
9. Sicherer SH, Muñoz-Furlong A, Godbold JH, Sampson HA. US prevalence of self-reported peanut, tree nut, and sesame allergy: 11-year follow-up. *J Allergy Clin Immunol*. 2010;125(6):1322–1326
10. Rona RJ, Keil T, Summers C, et al. The prevalence of food allergy: a meta-analysis. *J Allergy Clin Immunol*. 2007;120(3):638–646
11. Sicherer SH. Epidemiology of food allergy. *J Allergy Clin Immunol*. 2011;127(3):594–602
12. Sicherer SH, Wood RA; American Academy of Pediatrics Section On Allergy And Immunology. Allergy testing in childhood: using allergen-specific IgE tests. *Pediatrics*. 2012;129(1):193–197
13. Fleischer DM, Bock SA, Spears GC, et al. Oral food challenges in children with a diagnosis of food allergy. *J Pediatr*. 2011;158(4):578–583.e1
14. Gupta RS, Springston EE, Kim JS, et al. Food allergy knowledge, attitudes, and beliefs of primary care physicians. *Pediatrics*. 2010;125(1):126–132
15. Sampson HA, Aceves S, Bock SA, et al; Joint Task Force on Practice Parameters; Practice Parameter Workgroup. Food allergy: a practice parameter update-2014. *J Allergy Clin Immunol*. 2014;134(5):1016–1025.e43
16. Muraro A, Werfel T, Hoffmann-Sommergruber K, et al; EAACI Food Allergy and Anaphylaxis Guidelines Group. EAACI food allergy and anaphylaxis guidelines: diagnosis and management of food allergy. *Allergy*. 2014;69(8):1008–1025
17. Klemans RJ, van Os-Medendorp H, Blankestijn M, Bruijnzeel-Koomen CA, Knol EF, Knulst AC. Diagnostic accuracy of specific IgE to components in diagnosing peanut allergy: a systematic review. *Clin Exp Allergy*. 2015;45(4):720–730
18. Sampson HA, Gerth van Wijk R, Bindslev-Jensen C, et al. Standardizing double-blind, placebo-controlled oral food challenges: American Academy of Allergy, Asthma & Immunology-European Academy of Allergy and Clinical Immunology PRACTALL consensus report. *J Allergy Clin Immunol*. 2012;130(6):1260–1274
19. Neuman-Sunshine DL, Eckman JA, Keet CA, et al. The natural history of persistent peanut allergy. *Ann Allergy Asthma Immunol*. 2012;108(5):326–331.e3
20. Wood RA, Sicherer SH, Vickery BP, et al. The natural history of milk allergy in an observational cohort. *J Allergy Clin Immunol*. 2013;131(3):805–812
21. Sicherer SH, Wood RA, Vickery BP, et al. The natural history of egg allergy in an observational cohort. *J Allergy Clin Immunol*. 2014;133(2):492–499
22. Savage J, Sicherer S, Wood R. The natural history of food allergy. *J Allergy Clin Immunol Pract*. 2016;4(2):196–203, quiz 204
23. Santos AF, Douiri A, Bécares N, et al. Basophil activation test discriminates between allergy and tolerance in peanut-sensitized children. *J Allergy Clin Immunol*. 2014;134(3):645–652
24. Allen KJ, Koplin JJ. Prospects for prevention of food allergy. *J Allergy Clin Immunol Pract*. 2016;4(2):215–220
25. Fleischer DM, Sicherer S, Greenhawt M, et al; American Academy of

- Allergy, Asthma & Immunology; American Academy of Pediatrics, American College of Allergy; Asthma & Immunology, Australasian Society of Clinical Immunology and Allergy; Canadian Society of Allergy and Clinical Immunology; European Academy of Allergy and Clinical Immunology; Israel Association of Allergy and Clinical Immunology; Japanese Society for Allergology; Society for Pediatric Dermatology; World Allergy Organization. Consensus communication on early peanut introduction and the prevention of peanut allergy in high-risk infants. *J Allergy Clin Immunol*. 2015;136(2):258–261
26. Togias A, Cooper SF, Acebal ML, et al. Addendum guidelines for the prevention of peanut allergy in the United States: report of the National Institute of Allergy and Infectious Diseases-sponsored expert panel. *J Allergy Clin Immunol*. 2017; 139(1):29–44
  27. Greer FR, Sicherer SH, Burks AW; American Academy of Pediatrics Committee on Nutrition; American Academy of Pediatrics Section on Allergy and Immunology. Effects of early nutritional interventions on the development of atopic disease in infants and children: the role of maternal dietary restriction, breastfeeding, timing of introduction of complementary foods, and hydrolyzed formulas. *Pediatrics*. 2008;121(1):183–191
  28. Panjari M, Koplin JJ, Dharmage SC, et al. Nut allergy prevalence and differences between Asian-born children and Australian-born children of Asian descent: a state-wide survey of children at primary school entry in Victoria, Australia. *Clin Exp Allergy*. 2016;46(4):602–609
  29. Campbell RL, Bellolio MF, Knutson BD, et al. Epinephrine in anaphylaxis: higher risk of cardiovascular complications and overdose after administration of intravenous bolus epinephrine compared with intramuscular epinephrine. *J Allergy Clin Immunol Pract*. 2015;3(1):76–80
  30. Muraro A, Roberts G, Worm M, et al; EAACI Food Allergy and Anaphylaxis Guidelines Group. Anaphylaxis: guidelines from the European Academy of Allergy and Clinical Immunology. *Allergy*. 2014;69(8):1026–1045
  31. Lieberman P, Nicklas RA, Randolph C, et al. Anaphylaxis—a practice parameter update 2015. *Ann Allergy Asthma Immunol*. 2015;115(5):341–384
  32. Sicherer SH, Simons, FE; Section on Allergy and Immunology. Epinephrine for first-aid management of anaphylaxis. *Pediatrics*. 2017;139(3):e20164006
  33. Wang JS, Sicherer SH; Section on Allergy and Immunology. Guidance on completing a written allergy and anaphylaxis emergency plan. *Pediatrics*. 2017;139(3):e20164005
  34. Nowak-Węgrzyn A, Groetch M. Nutritional aspects and diets in food allergy. *Chem Immunol Allergy*. 2015;101:209–220
  35. Annunziato RA, Rubes M, Ambrose MA, Mullarkey C, Shemesh E, Sicherer SH. Longitudinal evaluation of food allergy-related bullying. *J Allergy Clin Immunol Pract*. 2014;2(5):639–641
  36. Herbert L, Shemesh E, Bender B. Clinical management of psychosocial concerns related to food allergy. *J Allergy Clin Immunol Pract*. 2016;4(2):205–213, quiz 214
  37. Sicherer SH, Mahr T; American Academy of Pediatrics Section on Allergy and Immunology. Management of food allergy in the school setting. *Pediatrics*. 2010;126(6):1232–1239

**Critical Issues in Food Allergy: A National Academies Consensus Report**  
Scott H. Sicherer, Katrina Allen, Gideon Lack, Steve L. Taylor, Sharon M. Donovan  
and Maria Oria

*Pediatrics* originally published online July 24, 2017;

<b>Updated Information &amp; Services</b>	including high resolution figures, can be found at: <a href="http://pediatrics.aappublications.org/content/early/2017/07/19/peds.2017-0194">http://pediatrics.aappublications.org/content/early/2017/07/19/peds.2017-0194</a>
<b>References</b>	This article cites 36 articles, 10 of which you can access for free at: <a href="http://pediatrics.aappublications.org/content/early/2017/07/19/peds.2017-0194#BIBL">http://pediatrics.aappublications.org/content/early/2017/07/19/peds.2017-0194#BIBL</a>
<b>Subspecialty Collections</b>	This article, along with others on similar topics, appears in the following collection(s): <b>Allergy/Immunology</b> <a href="http://www.aappublications.org/cgi/collection/allergy:immunology_sub">http://www.aappublications.org/cgi/collection/allergy:immunology_sub</a>
<b>Permissions &amp; Licensing</b>	Information about reproducing this article in parts (figures, tables) or in its entirety can be found online at: <a href="http://www.aappublications.org/site/misc/Permissions.xhtml">http://www.aappublications.org/site/misc/Permissions.xhtml</a>
<b>Reprints</b>	Information about ordering reprints can be found online: <a href="http://www.aappublications.org/site/misc/reprints.xhtml">http://www.aappublications.org/site/misc/reprints.xhtml</a>

American Academy of Pediatrics

DEDICATED TO THE HEALTH OF ALL CHILDREN™



# PEDIATRICS®

OFFICIAL JOURNAL OF THE AMERICAN ACADEMY OF PEDIATRICS

**Critical Issues in Food Allergy: A National Academies Consensus Report**  
Scott H. Sicherer, Katrina Allen, Gideon Lack, Steve L. Taylor, Sharon M. Donovan  
and Maria Oria

*Pediatrics* originally published online July 24, 2017;

The online version of this article, along with updated information and services, is  
located on the World Wide Web at:

<http://pediatrics.aappublications.org/content/early/2017/07/19/peds.2017-0194>

Pediatrics is the official journal of the American Academy of Pediatrics. A monthly publication, it has been published continuously since 1948. Pediatrics is owned, published, and trademarked by the American Academy of Pediatrics, 141 Northwest Point Boulevard, Elk Grove Village, Illinois, 60007. Copyright © 2017 by the American Academy of Pediatrics. All rights reserved. Print ISSN: 1073-0397.

American Academy of Pediatrics

DEDICATED TO THE HEALTH OF ALL CHILDREN™

