Consensus on DEfinition of Food Allergy SEverity (DEFASE) an integrated mixed methods systematic review

Stefania Arasi*1, Ulugbek Nurmatovb,1, Audrey Dunn-Galvinc, Shahd Daherd, Graham Roberts e, f, Paul J. Turner g, h, Sayantani B. Shinder i, Ruchi Guptaj,2, Philippe Eigenmannk, Anna Nowak-Wegrzynl, m, Mario A. Sánchez Borges n, Ignacio J. Ansotegui o, Montserrat Fernandez-Rivasp, Stavros Petroud, Luciana Kase Tannoq, r, Marta Vazquez-Ortiz j, Brian P. Vickery u, Gary Wing-Kin Wong v, Motohiro Ebisawaw, and Alessandro Fiocchi a

ABSTRACT

Background and aims: The term “Food Allergy” refers to a complex global health problem with a wide spectrum of severity. However, a uniform definition of severe food allergy is currently missing. This systematic review is the preliminary step towards a state-of-the-art synopsis of the current evidence relating to the severity of IgE-mediated food allergy; it will inform attempts to develop a consensus to define food allergy severity by clinicians and other stakeholders.

Methods: We undertook a mixed-methods systematic review, which involved searching 11 international biomedical databases for published studies from inception to 31 December 2019. Studies were independently screened against pre-defined eligibility criteria and critically appraised by established instruments. The substantial heterogeneity of included studies precluded meta-analyses and, therefore, narrative synthesis of quantitative and qualitative data was performed.

Results: We found 23 studies providing eligible primary data on symptom-specific severity of food allergic reactions, and 31 previously published symptom-severity scoring systems referred to food allergic reactions. There were seven studies which assessed quality-of-life measures in patients (and family members) with different food allergy severity and two studies that investigated the economic burden of food allergy severity. Overall, the quality and the global rating of all included studies were judged as being moderate.

Conclusions: There is heterogeneity among severity scoring systems used and even outcomes considered in the context of severity of food allergy. No score has been validated. Our results will
be used to inform the development of an international consensus to define the severity of food allergy.

**Systematic review registration:** A protocol was prospectively registered with the International Prospective Register of Systematic Reviews (PROSPERO) database with the registration number CRD42020183103 (https://www.crd.york.ac.uk/prospero/#recordDetails).

**Keywords:** Definition, Food allergy, Severity, Systematic review, Mixed-methods study

**BACKGROUND AND RATIONALE**

Over the last few decades, with increasing prevalence, food allergy (FA) has emerged as a global health problem affecting up to 10% of the population.\(^1\)\(^-\)\(^4\) Epidemiological studies demonstrate an increase not only in prevalence, particularly in children,\(^5\),\(^6\) but also in severity with remarkable morbidity and in some cases, mortality.\(^7\),\(^8\) A diagnosis of FA can result in a significant adverse impact on health-related quality of life for both allergic individuals and their families, with an increase in emotional, social, and financial burdens.\(^9\) However, as for other diseases, including allergic pathologies, there are different phenotypes of FA with variability in allergen-specific clinical symptoms and eliciting doses.\(^10\) Patients with milder forms, such as those with only oral symptoms, are certainly worthy of diagnostic attention, but may not require all the therapeutic and management resources that are necessary for the patient at higher risk of life-threatening food-induced anaphylaxis. Diagnosing FA remains highly complex, and although numerous biomarkers are under exploration,\(^11\),\(^12\) the most commonly used tests remain food allergen-specific immunoglobulin E (sIgE) and skin prick testing (SPT) which do not correlate with severity.\(^13\) Presently, without a reliable classification system, we risk treating all FA patients in the same way, in effect, a one-size-fits-all approach that is unhelpful to patients, their families, and their providers. Severity classifications are available for both allergic rhinitis and asthma; however, for FA, no such specific scoring system for classifying severity currently exists.\(^14\),\(^15\)

Without a standardized classification system in place, terminology and definitions that are currently in use are not comparable across studies and among different stakeholders.\(^16\) Standardizing the classification of FA severity will benefit not just patients and providers, but also patient advocacy groups, disease registries, research, food and drug industries, government agencies and regulators, as well as legislative bodies. There is presently a great need for an international consensus-based system to define FA severity.

World Allergy Organization (WAO) is undertaking the development of an international definition and classification of severity associated with food allergy (“DEFinition of Food Allergy SEverity”, DEFASE). The preliminary step in the formulation of a uniform definition and classification of FA severity includes a state-of-the-art synopsis of the current evidence. This systematic review focuses exclusively on IgE-mediated food allergy (ie, acute allergic reactions manifesting as a broad spectrum of signs/symptoms ranging from urticaria to vomiting and wheezing, up to fatal or near-fatal anaphylaxis).\(^17\) To our knowledge, this is the first systematic review of the literature on current severity classifications used for FA.

**METHODS**

This systematic review (SR) was conducted by a panel of allergy specialists, psychologists, other health-care professionals, economists, academicians, researchers, patient representatives, and methodologists. The members of DEFASE team come from Europe, North and South America, Asia, and Australasia.

**Plan of investigation**

The methods are briefly described herein. A detailed SR protocol “Consensus on the DEFinition of Food Allergy SEverity (DEFASE): protocol for an
integrated mixed methods systematic review” was registered with International Prospective Register of Systematic Reviews (PROSPERO) CRD42020183103 and accepted for publication in World Allergy Organization Journal.

Population

Studies on patients of any age with a physician-confirmed diagnosis of IgE-mediated food allergies to eggs, milk, peanuts, tree nuts, and/or any other food were eligible for inclusion.

Outcomes

Our outcomes of interest were: (a) symptom-related severity scores for food allergic reactions; (b) non-symptom related severity scores for food allergy, ie, health-related quality of life and economic evaluations; (c) methodological approaches used to derive definitions of food allergy severity; (d) the features used to define them (ie, variables, either clinical [eg, type and numbers of reactions to the culprit food; comorbidities; cofactors; disease-related quality of life impairment], or immunological characteristics [eg, pattern of sensitization to allergenic molecules, IgE-specific activities]); (e) the characteristics associated with severity category.

We considered and categorised food allergy severity as either symptom-related or non-symptom-related severity scores. We have considered all scores designed for or applied to food induced allergic reactions. However, we included primary data on symptom severity of food allergic reactions only from papers reporting on physician confirmed diagnosis of IgE mediated food allergy based on a positive history and IgE sensitization (skin prick test, SPT) and/or serum levels of specific IgE (sIgE) with/without oral food challenge (OFCs). We also evaluated validated scoring systems for food allergy quality of life (FAQL) and how FAQL and/or food allergy independent measure (FAIM) is impacted by "severity" of food allergic reactions in patients with physician - confirmed food allergy.

This mixed-methods SR was designed to capture and include all types of partial and full economic evaluations of food allergy severity. The economic evaluations could either be partial economic evaluations (cost analyses or cost-cost offset analyses) or they could be full economic evaluations that identify, measure, and value costs and outcomes of the severity of food allergy with an appropriate comparator(s). The different types of full economic evaluations include cost-effectiveness analyses, cost-utility analyses, cost-benefit analyses, cost-consequence analyses, and cost-minimisation analyses. The results were analysed to determine the number of studies that support the severity of food allergy on cost-effectiveness grounds, and where available an overall recommendation was made based on the results of partial economic evaluations (eg, cost analysis).

Study types

Papers whose primary or secondary aim is to define or identify severity classifications of IgE-mediated food allergies in humans were considered eligible for inclusion in our SR. The following study types were eligible for inclusion:

- All analytical studies: ie, cohort, case-control, and cross-sectional studies; case series involving 40 or more participants; and economic evaluation of FA severity.
- All interventional studies: ie, randomized controlled trials, RCT; quasi- RCTs; controlled clinical trials, CCT; interrupted time series, ITS; and controlled before after studies.

In addition, we also included reviews, SRs, guidelines, position and consensus papers, editorials, and rostrums.

The following study types were excluded: studies of non-IgE mediated food allergy; studies that used only self-reported diagnosis of food allergy; primary data from studies on allergen immunotherapy; non-research letters and editorials; case reports; case-series with less than 40 participants; and in-progress phenotyping studies (abstracts) as they are unlikely to provide sufficient detail on the definitions of food allergy severity score; animal studies; and studies that examined food allergy as a predictor of a separate outcome (eg, asthma development).
Research methods for identification of studies: electronic databases

We systematically searched 11 international databases: AMED (1985–2019); CAB (1910–2019); CINAHL (1937–2019); Cochrane Library (1992–2019); Econlit (1886–2019); EMBASE (1980–2019); Global Health (1987–2019); Google Scholar (2000–2019); ISI Web of Science (which contains the Science Citation Index) (1970–2019); MEDLINE (1966–2019); TRIP (2003–2019).

Search strategy for electronic databases

A search strategy was developed in Medline format and adopted for other databases. MEDLINE and EMBASE databases were searched using the controlled vocabulary search terms (MeSH and EMTREE, respectively) combined using Boolean terminology with a wide-range of free-text terms. The results were limited to humans (see Online Supplementary material, section “Search strategy”). There were no publication year or publication status restrictions; however, the searches were restricted to only English language. Searches were undertaken from inception up to 31 December 2019.

Additional search methods

All references of published studies were hand searched. The bibliographies of all eligible studies were scrutinised to identify possible additional studies. In addition, we contacted the primary study authors to clarify discordant data [Table S1]. We also reviewed the reference lists of relevant studies.

Study selection

Duplicate publications were removed. Titles and abstracts of identified studies were checked against the inclusion/exclusion criteria independently by two reviewers.

Full-text papers were retrieved if their titles and/ or abstracts appeared to meet the eligibility criteria or if the decision could not be made based on the titles and/or abstracts alone. Assessment of the full texts of each retrieved paper was undertaken independently by two reviewers using the same criteria. Disagreements about inclusion were resolved through discussion at the meetings.

Assessment of methodological quality

The methodological quality of included observational studies was independently assessed by two reviewers (UN, SA) by using the Effective Public Health Practice Project (EPHPP). We focused on the following domains to assess the quality of included studies: selection bias; study design; confounders; blinding; data collection method; withdrawals and dropouts; and final global rating. Each component-specific parameter (ie, suitability of the study design for the research question; risk of selection bias; exposure measurement; outcome assessment; and generalizability of findings) was given a judgment: “strong”; “moderate”; and “weak”. At the end of critical appraisal, we also provided the overall grading for each study.

Data extraction

Data were independently extracted onto a customized data extraction sheet by two reviewers (SA, UN), and any discrepancies were resolved by discussion or, if agreement was not reached, the third reviewer arbitrated.

Meta-analysis

Meta-analysis was inappropriate given the substantial heterogeneity of the populations, exposures, outcomes and study designs.

RESULTS

An overview: characteristics of included studies

Our searches identified 12,148 potentially relevant papers and 10 further papers identified by experts; 2365 duplicate papers were removed; a further 9705 papers were excluded for not meeting our inclusion criteria. Furthermore, 88 papers were at full text level, and in total 52 studies satisfied our inclusion criteria and were thus included in our systematic review (see Fig. 1, PRISMA flow diagram). Manuscripts excluded at full-text screening phase and reasons for exclusion are explained in Table S2.

We found 23 studies providing eligible primary data on symptom-specific severity of food allergic reactions [Table S3].
Fourteen studies reported aggregated symptom-specific primary data referred to allergic reactions triggered by any allergenic source (ie, not only by food allergens). We tried to contact the respective authors several times; however, only in three studies, primary data referred specifically to food allergic reactions were provided by the contacted authors [Table S1] and, therefore, those three studies have been included in the category of primary data.

In terms of study design, the 23 eligible studies were: 12 cohort; two case-control; four cross-sectional; and five case series studies.

Additionally, our SR identified 31 previously published symptom-severity scoring systems referred to food allergic reactions [Table 1]. Twenty-three were primary studies providing new symptom-severity scoring systems to assess food-induced allergic reactions [Tables 1 and 2]. Of note, we were able to pool primary data eligible for our SR only from 11 of them. The remaining 20 studies provided new symptom-severity scores but not eligible for our SR.

Fig. 1 PRISMA flow diagram
| Study | (First author, year, country) | No. | Gender (male:female) | Target age group | Setting | Level 1 | Level 2 | Level 3 | Level 4 | Level 5 | Development / Notes | Applying Primary Studies (included in our systematic review) |
|-------|--------------------------------|-----|----------------------|------------------|---------|--------|--------|--------|--------|--------|-------------------|-------------------------------------------------|
| PRIMARY STUDIES | | | | | | | | | | | | |
| Amin, 2012, USA | 3 | food ped org aller gy | y | skin and mucosal $ | GI $ or resp $ or multisystemic involvement (≥ 3 systems) | urticaria or hypoxia/neurological compromise/ resp $ or vascular $ (flushing, headache, hypotension) or multisystemic involvement (≥ 4 systems) | Amin, 2012 |
| Astier, 2006, France & USA | 5 | peanut ped org aller gy | n | transient GI, rhinoconjunctivitis, <10 h, eczema onset | 2 organs involved | 3 organs involved or 1 of: asthma or needing Tx + laryngeal oedema or hypotension | cardiacl & resp $ requiring ICU admission | Modified from iwan 2001 |
| Ben-Shoshan, 2010, Canada* | 3 | all ped org Self-reported FA | n | At least 1 of: pruritus, urticaria, flushing or rhinoconjunctivitis | 2 organs involved | At least 1 of: angioedema or GI/S + respiratory distress, gastrointestinal complaints, or breathing difficulties (other than wheeze) | At least 1 of: wheeze, cyanosis or circulatory collapse | Hourihane 1997; Brown 2004; Sicherer 1999 |
| Boyano-Martinez, 2009, Spain | 3 | milk (egg, soy, dairy) ped org aller gy | n | skin $ (angioedema excluded), rhinitis, or conjunctivitis | angioedema or GI/S | lower resp tract $ (loss of voice, dysphonia, wheezing, stridor, and/or cyanosis) or systemic $ (LOC, weakness, or dizziness) | Boyano-Martinez, 2009; Boyano-Martinez, 2012 |
| Bragantza, 2006, Australia | 2 | all ped org EO | n | skin and/or GI $ | resp $ (asphyxia/dysphonia/laryngeal oedema/cyanosis) and/or CVS and/or neurological $ (hypotension/syncope/slurred/GCS<15) | Modified from ASCIA Guidelines for EpiPen prescription 2004* |
| Brown, 2004, Australia | 3 | all all org EO | n | skin $ only (Generalized erythema, urticaria, periorbital oedema, or angioedema) | non-severe resp/CVS GI $ (hypoxia, stridor, wheeze, nausea, vomiting, dizziness (presyncope), diaphoresis, chest or throat tightness, or abdominal pain) | cyanosis or hypoxia (SO₂<92%) or hypotension (systolic blood pressure < 90 mm Hg in an adult) or collapse or altered consciousness or incontinence | Brown, 2004; Brown, 2013; Moro-Moro, 2011; Tejedor-Alonso, 2013; Ye, 2015 |
| Burks, 2012 (COFAR) | S food | all organ | aller V | n | Grade 1 (Mild) | Grade 2 (Moderate) | Grade 3 (Severe) | Grade 4 (Life-Threatening) | Grade 5 (Death) |
|---------------------|-------|----------|--------|---|----------------|------------------|-----------------|--------------------------|---------------|
|                     |       |          |        |   | Transient or mild discomforts (< 48 hours), no or minimal medical intervention/Tx required. These $ may include pruritus, swelling or rash, abdominal discomfort or other transient symptoms. | Symptoms that produce mild to moderate limitation in activity, some assistance may be needed; no or minimal intervention/Tx is required. Hospitalization is possible. These $ may include persistent hives, wheezing, dyspnea, abdominal discomfort/ increased vomiting or other $ | Marked limitation in activity, some assistance usually required; medical intervention/Tx required, hospitalization is possible. $ may include bronchospasm with dyspnea, severe abdominal pain, throat tightness with hoarseness, transient hypotension among others. Parenteral medication(s) are usually indicated. | Extreme limitation in activity, significant assistance required; significant Tx, intervention is required; hospitalization is probable. Symptoms may include persistent hypotension and/or hypoxia with resultant decreased level of consciousness associated with collapse and/or incontinence or other life threatening $ | Death |

| Clark, 2004, USA | 1 food | all organ | ED y | N/R | N/R | N/R | N/R | 2 organ systems (skin, respiratory, CVS or GI) or hypotension alone (<ystolic blood pressure < 100 mm Hg) | Clark, 2004 |
|-------------------|-------|----------|------|-----|-----|-----|-----|---------------------------------|----------------|
| Ewan, 2001, UK | 5 peanut and nut | all organ | aller V | n | Localised skin $ (urticaria/angioedema) | Generalised skin $ (urticaria/angioedema) | Skin + GI $/rhinocconjunctivitis | Mild laryngeal $/oedema (voice change/lightening of throat) or mild asthma | Severe dyspnea or any hypertensive $ (light-headedness/collapse/loss of consciousness) | Ewan, 2001 |
| Fleisher, 2005, USA | 3 tree nuts | all organ | aller V | n | Skin $ (any rash, pruritus, worsening edema, angioedema, eye or facial swelling) and/or oral $ (oral, throat pain, palatal hives or erythema, or itching of the palate, tongue, or lips) only | Upper respiratory (rhinorrhea, nasal congestion, nasal pruritus, sneezing) and/or GI (nausea, vomiting, diarrhea, abdominal pain) only or any 3 systems | Resp lower (cough, wheeze, shortness of breath, stridor, hoarseness) and/or CVS $ (hypotension, light-headedness, syncope, or collapse or any 4 systems) | }
| Arasi et al. World Allergy Organization Journal (2021) 14:100503 |
|------------------------------------------------------------------|
| Gupta, 2011, USA | 2 | food | ped | org | allerg | n | parent: report of angioedema of the lips, eyes, or face, other angioedema, coughing, other oropharyngeal symptoms, eczema, flushing, hives, pruritus, and vomiting |
| | | | | | | | parent: report of anaphylaxis, low blood pressure, trouble breathing, or wheezing. A reaction including vomiting, angioedema, and coughing in combination was also categorized as severe. |
| Gupta, 2018, USA | 2 | food | ped | org | allerg | n | patient report of only 1 of the following specific “stringent” symptoms: hives, swelling, lip and/or tongue swelling, difficulty swallowing, throat tightening, trouble breathing, wheezing, vomiting, chest pain, rapid heart rate, fainting, low blood pressure, WITH OR WITHOUT any of the following symptoms: itching, rash, hoarse voice, achy mouth, mouth or throat tingling, nasal congestion, repetitive cough; belly pain, cramps, diarrhea, nausea |
| | | | | | | | report of specific “stringent” symptoms within ≥2 organ systems (skin, respiratory, CV or GE) Skin: hives, Swelling, Lip and/or tongue swelling, difficulty swallowing, throat tightening Respiratory: Chest tightening, trouble breathing, wheezing CV: Vomiting GE: Low blood pressure, fainting, rapid heart rate, chest pain |
| Hourihane, 1997, UK | 3 | peanut | ped | org | validated questionnaire (allegry) | n | urticaria / pruritus |
| | | | | | | | angioedema or laryngeal swelling or breathing difficulty (except wheeze) |
| | | | | | | | wheeze / cyanosis / collapse/faint |
| Hourihane, 2005, UK | 3 | peanut | all | amount | allergy | n | >3/4 nut eaten/ <100 mg peanut protein |
| | | | | | | | 1/4 – 3/4 nut eaten/ 3-99 mg peanut protein |
| | | | | | | | Fragment swallowed/ <3 mg mg peanut protein |
| | | | | | | | Mucosal touch, none eaten |
| | | | | | | | Inhalation / skin contact only |
| | | | | | | | Modified from Hourihane 1999 Furthermore authors proposed to combine dosage and symptom grades to give an overall score for reactions to peanut in the community or during low-dose DBPCFC. |
| | | | | | | | Hourihane, 1997 |
| | | | | | | | Hourihane, 2005 |

Score 2–5, mild; 6–13, moderate; 14–25, severe.

DBPCFC, double-blind, placebo-controlled food challenge.
| Author(s) | Year, Country | Case Type | Description | Symptoms | Outcome |
|-----------|---------------|-----------|-------------|----------|---------|
| Itazawa, 2020, Japan | Volume 14, No. 3, March 2021 | 5 | food | ped | aller | GV | Grade 1: At least 1 of: • localised skin • subjective oral • generalised skin • mild GI • mild nasal • decrease in activity level |
| Kimchi, 2015, Canada | 3 | all | all | org | ED/ambulance | n | Grade 3: At least 1 of: • marked nasal S or • x2 vomiting or diarrhoea • persistent abdominal pain • marked nasal congestion and/or rhinorrhea • repeated sneezing • persistent cough • laryngeal pruritus • tachycardia |
| Mardougias, 2002, USA | 1 | food | ped | Tx | GP | n | Grade 4: At least 1 of: • laryngeal tightness • hoarseness • barking cough • dysphagia • wheezing • dyspnoea • cyanosis • arrhythmia • hypotension • fear of death |
| Mehli, 2005, Germany | 4 | all | ped | org | GP-reported FA | n | Grade I: • mild systemic reaction: reddening induration itching, etc. |
| Mueller, 1959, USA | 5 | ven | org | ICU | n | Slight general reaction Generalized urticaria, itching, malaise and anxiety |

**Modified slightly from Sampson 2003. The symptom grade is determined according to the organ S of the highest grade.**

**Modified from Brown, 2004.**

**At least 1 of: • cardiorespiratory arrest; • need for isotropic support; • fluid bolus ≥ 20 mL/kg; • ≥ 1 dose of epinephrine by any route; • ≥ 1 dose of nebulised bronchodilator. Cases where intubation was necessary were deemed “near-fatal.”**

**The investigator defined criteria.**

**Only in combination with other grade IV symptoms.**

**Severity score originally designed for allergic reactions to insect stings but applied also to food induced allergic reactions.**
### Primeau, 2000, Canada
3 peanut all organ allergy n

| Symptom         | Score 1: | Score 2: | Score 3: | Score 4:          |
|-----------------|----------|----------|----------|-------------------|
| hypotension     | faint/dizzy | unconscious | prolonged unconsciousness | cardiac arrest |
| bronchospasmy   | wheezy    | severe    | poor response to Tx |              |
| angioedema      | lips/face | generalized | severe | |
| pharyngeal edema| hoarse    | difficulty in swallowing | severe difficulty in breathing | |
| urticaria       | mild      | generalized | generalized + severe | |
| pruritus        | mild      | severe    |         | |
| conjunctivitis  | mild      | severe    | ulcerated | |
| vomiting        | once      | severe    | prolonged over hours | |
| diarrhoea       | once      | severe    | prolonged over hours | |
| uterine pain    | period   | severe    | cramps   | uterus bleeding   |

### Punchey, 1996, UK
3 all all algorythm allergy n

Reactions were classified as fatal, major or minor. Major reactions were taken as those where the sum of the scores for the principal 5 (hypotension, bronchospasmy and angioedema) was ≥ 9, or was 7 with ≥ 3 additional points from the other 5.
An empirical formula was used to combine the 5 scores to allow ranking of the severity of the worst reaction for each patient: severity index = 4 x (score for hypotension + bronchospasmy + pharyngeal oedema) + 2 x (score for angioedema + urticaria) + (sum of all other scores).

### Sinclair, 1999, USA
3 peanut all self-reported FA n

| Symptom         | Score 1: | Score 2: | Score 3: | Score 4:          |
|-----------------|----------|----------|----------|-------------------|
| significant skin or GI S or mild S in other systems | wheezing alone or significant involvement of 2 organs/systems | At least 1 of: hypotension, wheezing, rhives, diarrhoea/vomiting, ≥ 3 systems involved |

### van der Zee, 2011, The Netherlands
5 peanut ped algorythm ED n

All S were classified into 5 categories: skin (score 1), upper airway (eyes, nose, and throat); (score 3), lower airway (lungs (score 3), GI (score 2), and CV (score 3) or neurological S (score 3).
To compute a severity index for the reaction at home, the scores for each category were summed. This index ranges from 0 to 12 and was classified into tertiles as follows: tertile 1:0 to 2; tertile 2:0 to 6; and tertile 3:7 to 12.

Developed by the authors

Developed by the authors

Developed by the authors
| Study                        | GRADE I                                                                 | GRADE II                                                                 | GRADE III                                                                 | GRADE IV                                                                 | GRADE V                                                                 | Notes                                                                 |
|------------------------------|------------------------------------------------------------------------|------------------------------------------------------------------------|------------------------------------------------------------------------|------------------------------------------------------------------------|------------------------------------------------------------------------|-----------------------------------------------------------------------|
| Cox, 2010** (WAO)            | • skin (urticaria and/or erythema-warmth and/or pruritus, other than localized at the injection site) or | • lower resp: mild asthma (cough, wheeze, shortness of breath) with good response to Tx and/or | • lower airway (severe asthma not responding or worsening in spite of Tx) and/or | • resp failure and/or | • score designed primarily for adverse reactions to subcutaneous immunotherap y (SCIT) but also for any allergic reaction. The symptom grade is determined according to the organ $SO$ of the highest grade. |
| Muraro, 2007 (EAACI)         | • upper resp or *conjunctival $                               | • GI $ (abdominal cramps and/or vomiting/diarrhea) or | • upper airway (laryngeal edema with stridor) or | • CVS collapse/ hypotension and/or | |                                                                 |
|                              | • nausea                                                            | • uterine cramps                                                      | • nausea                                                            | • LOC (vasovagal excluded)                                             |                              |                                                                 |
| Muraro, 2018 (EAACI)         | • MILD                                                                | • MODERATE                                                            | • SEVERE                                                             | Modified from Sampson 2003                                             |                                                                                     |                                                                                      |
|                              | • skin $ (Sudden itching of eyes and nose, generalized pruritus, flushing, urticaria, angioedema) | • GI (crampy abdominal pain, diarrhoea, recurrent vomiting)         | • cyanosis or SpO$_2$ <92%, resp arrest                              |                                                                                     |                                                                                     |                                                                                      |
|                              | • mild GI $ (oral pruritus, oral tingling, mild lip swelling, nausea or emesis, mild abdominal pain) | • resp $ (hoarseness, barking cough, difficulty swallowing, stridor, dyspnea, moderate wheeze) | • hypotension and/or collapse, dysrhythmia, severe bradycardia, CVS arrest, confusion, LOC |                                                                                     |                                                                                     |                                                                                      |
|                              | • mild upper respiratory $ (nasal congestion and/or sneezing, rhinorrhea, throat pruritus, throat tightness) mild wheezing | • tight headedness feeling of impending doom                         | • severe, potentially life-threatening $ involving:                   |                                                                                     |                                                                                     |                                                                                      |
|                              | • tachycardia                                                        | | • CV$S$, neurologica, bronchial and/or laryngeal $                   |                                                                                     |                                                                                     |                                                                                      |
|                              | • change in activity level = anxiety                                 | |                                                                                     |                                                                                     |                                                                                     |                                                                                      |
|                              |                                                                       | |                                                                                     |                                                                                     |                                                                                     |                                                                                      |

Simplified classification of severity according to the organ system involved proposed across different allergic triggers to address the needs of different stakeholder groups.
| Author, Year, Location | i.all | iall | all | all | n | GRADE I: | GRADE II A: | GRADE IIIA: | GRADE IIIB: | GRADE IIIC: | GRADE IV: | GRADE IIC: | Proposal for a uniform grading system of allergic reactions across the full spectrum of allergic triggers | Author, Year |
|------------------------|------|------|-----|-----|----|---------|-----------|------------|------------|------------|-----------|------------|--------------------------------------------------|------------|
| Niggemann, 2016, Germany | 6    | all  | all | all | all | 1 | • local S (e.g. redness, swelling, pruritus) | • skin (e.g. urticaria, angioedema, flush) OR • GI tract (e.g. abdominal pain, vomiting, diarrhea) | • resp (cough, wheezing, stridor) OR • CVS (tachycardia, lowered BP) | • severe resp (e.g. objective dyspnoea, assoesory muscles) and/or • severe CVS (shock) | Proposal for a uniform grading system of allergic reactions across the full spectrum of allergic triggers | Vrhovk, 2019 |
| Ring, 1999, Germany | 4    | all  | all | all | all | 1 | • local S (e.g. pruritus, flush, urticaria, angioedema) | • skin (e.g. urticaria, angioedema, flush) | • resp (cough, wheezing, stridor) OR • CVS (tachycardia, lowered BP) | • severe resp (e.g. objective dyspnoea, assoesory muscles) and/or • severe CVS (shock) | Proposal for a uniform grading system of allergic reactions across the full spectrum of allergic triggers | Vrhovk, 2019 |
| Sampson, 2003, USA | 5    | food | all | all | all | 1 | • local S (e.g. pruritus, flushing, urticaria, angioedema) | • skin (e.g. urticaria, angioedema) | • GI S (nausea, vomiting, diarrhea) | • severe resp (e.g. objective dyspnoea, assoesory muscles) and/or • severe CVS (shock) | Proposal for a uniform grading system of allergic reactions across the full spectrum of allergic triggers | Vrhovk, 2019 |
| Sampson, 2012 PRACTALL | 3    | food | all | all | all | 1 | • at least one of: | • nasal S (nasal congestion and/or sneezing) | • change in activity level | • respiratory arrest | Modified from Ring and Meßmer, 1977 | Proposal for a uniform grading system of allergic reactions across the full spectrum of allergic triggers | Vrhovk, 2019 |

*Note: The table includes examples of grading systems for allergic reactions, demonstrating the use of specific symptoms and their severity levels. Each grading system is designed to provide a uniform approach to the grading of allergic reactions, facilitating a more consistent and standardized assessment across different allergic responses.*

**Authors:**
- Arasi et al.
- Niggemann
- Ring
- Sampson, USA
- Sampson, 2012 PRACTALL

**Years:**
- 2016
- 1999
- 2003
- 2012

**Topics:**
- Localized skin reactions
- Generalized skin reactions
- GI tract reactions
- Respiratory reactions
- Cardiovascular reactions

**Proposals:**
- Proposal for a uniform grading system of allergic reactions across the full spectrum of allergic triggers.
primary data for our SR since they: a) included allergic reactions triggered by a different\textsuperscript{58} or any allergenic source (ie, not only food);\textsuperscript{46-48} b) or were based on self-reported diagnosis of food allergy;\textsuperscript{1,42,45,49} c) or included food allergy diagnosis only based on IgE-sensitization without history of ingestion of the suspected culprit food;\textsuperscript{44} d) or referred to oral immunotherapy trials.\textsuperscript{43} Eight out of the 31 that included symptom-severity scores were provided by secondary research papers.\textsuperscript{15,51-57} The following four secondary studies were from international collaboration, specifically, from: World Health Organization (WHO),\textsuperscript{51} European Academy of Allergy and Clinical Immunology (EAACI),\textsuperscript{15,52} and PRACTALL;\textsuperscript{56} two studies from Germany,\textsuperscript{53,54} two studies were from the United States,\textsuperscript{55} and one study from Sweden.\textsuperscript{57}

There were seven studies which assessed quality of life (QoL) measures in patients (and family members) with different food allergy severity.\textsuperscript{59-65} All were primary studies, with five out of the seven employing a cross-sectional,\textsuperscript{59,61,63,65} and two a longitudinal,\textsuperscript{60,64} design [Table S4].

Two studies investigated the economic burden of food allergy severity\textsuperscript{66,67} [Table S5].

The studies were undertaken in Australia (n = 3); Canada (n = 4); France (n = 1); Germany (n = 3); Japan (n = 1); Korea (n = 1); Spain (n = 4); Sweden (n = 2); The Netherlands (n = 4); United Kingdom (n = 5); United States (n = 14); and international collaboration (n = 10).

Critical appraisal of studies

Quality assessment of the 23 included primary studies on symptom-severity assessment suggested that out of 12 cohort studies 10 were judged as strong.\textsuperscript{19,29,34-40,68} Out of two case-control studies one was judged as moderate and two as weak. Three cross-sectional studies\textsuperscript{22,23,30} were judged as moderate, and one study\textsuperscript{31} was judged as weak. Among the five case-series, two were judged as moderate,\textsuperscript{24,26} and three studies were judged as weak\textsuperscript{25,28,32} [Table S6A].

Each of the seven included studies on QoL utilized a cross-sectional design. In terms of critical appraisal, four of them have been judged as moderate\textsuperscript{60,61,64,65} and three as weak\textsuperscript{59,62,63} [Table S6B].

The absence of full economic evaluation studies in this SR precluded the use of Consolidated Health Economic Evaluation Reporting Standards (CHEERS) checklist for critical appraisal of only two cost-evaluation studies.

Primary data on symptom-specific food allergy severity

Food allergy diagnosis

We included primary data on symptom severity of food allergic reactions only from papers reporting on physician confirmed diagnosis of IgE mediated food allergy based on a positive history and IgE sensitization (SPT and/or serum levels of...
| Uncommon Criteria | N. organ Systems | Neurological | Cardiovascular | Respiratory Tract | Gastrointestinal Tract | Skin |
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Table 2. Overview of all included symptom-severity scores in included studies for each listed symptom ordered by organ

| Legend          | Description             |
|-----------------|-------------------------|
| Light green     | mild allergic reaction  |
| Light yellow    | moderate allergic reaction |
| Dark red        | severe allergic reaction |
| Dark red        | unspecified              |

**List of Abbreviations:** PEF, peak expiratory flow; $S$, symptom(s)

If a severity score includes more than 3 grades of severity (mild, moderate, severe), the box reports the respective grade. For symptoms included in different grades, the box is coloured as for the most severe grade and grade numbers are specified.

1 Primary studies providing symptom-severity score but data referred to allergic reactions triggered by different allergenic sources (Kimchi, 2015; Mehl, 2005; Pumphrey, 1996; van der Zee, 2011).
2 Primary studies providing symptom-severity score but data based on self-reported diagnosis of food allergy (Ben-Shoshan 2010, Gupta, 2011, Gupta, 2018, Sicherer, 1999).
3 Primary studies providing symptom-severity score but data based on food allergy diagnosed only based on IgE-sensitization without history of ingestion of the suspected culprit food (Fleisher, 2005).
4 Severity score originally designed for allergic reactions to insect sting but afterwards applied also to food-induced allergic reactions [Mueller, 1959]
# An empirical formula was used to combine the symptom scores for each symptom to rank the severity of reaction (Hourihane, 2005; Pumphrey, 1996; van der Zee, 2011).
6 Score symptom combining dosage and symptom grades (Hourihane, 2005).
* Combined symptoms.
* <10 hives.
1 Symptom classified as grade 1 if not requiring treatment or transient, otherwise grade 2.
4 Breathing difficulty other than wheeze.
6 Report of specific “stringent” symptoms within ≥2 organ systems (skin, respiratory, CVS or GI) is mandatory [Skin: Hives, Swelling, Lip and/or tongue swelling, difficulty swallowing, throat tightening; Respiratory: Chest tightness, trouble breathing, wheezing; GI: Vomiting; CVS: Low blood pressure, fainting, rapid heart rate, chest pain]
9 Uterine symptoms: Pumphrey, 1996: period pains (grade 1, mild); severe cramps (grade 2, moderate); uterine bleeding (grade 3, severe); Cox, 2010: uterine cramps (grade 2, mild)
1 Incontinence: Brown, 2004 (grade 3, severe); Mueller 1959 (grade 4, severe, shock reaction).
4 Reduction PEF<40%; good response to inhaled bronchodilator.
6 Maccougall, 2002: criteria for severe grade: at least 1 of: cardiorespiratory arrest; need for inotropic support; fluid bolus ≥ 20 ml/kg; ≥ 1 dose of epinephrine by any route; ≥ 1 dose of nebulised bronchodilator.
3 Loss of bowel control: Murara 2007 (grade 3, severe); Sampson 2003 (grade 5, severe); Vetander 2011 (grade 3, severe).
6 Chest tightness: Mueller 1959 (grade moderate, general reaction); Vetander, 2011 (grade 1, mild).
specific IgE) with/without OFCs. In 10 out of 23 included primary studies assessing symptom severity, food allergy was confirmed by OFC. Among these 10 studies, 1 used a validated questionnaire with a sensitivity of 100% and a specificity of 87% for detecting peanut allergy compared with the gold standard of double-blind placebo-controlled food challenge (DBPCFC). 41

Setting
The assessment of food allergy severity was carried out in different emergency departments (EDs) of hospitals in nine studies. 24–26,28,34,35,39,40,68 Food allergy severity was assessed by allergist specialist consultations in 15 studies. 4,19–23,29–33,35–38

Recurrence of adverse reactions
Recurrence of adverse reactions (ARs) were reported in 10 studies. 22,23,28–31,35,36,38,40 However, these studies reported data in different formats, and we could not pool data statistically.

Epinephrine use was reported in 11 studies; 22,27–32,33,36,38,40 ED admission was recorded in five studies; 22,23,27,31,40 and admission to hospital was provided in two studies. 28,30 Admission to intensive care unit (ICU) was recorded in six studies; 22–24,28,31,38 ranging from 0 up to 1.1%.

Symptom-severity scoring systems for food allergy
Our SR identified 31 previously published instruments focusing on severity of food allergic reactions. 1,4,15,19,26,22,24,26,27,29–33,42–58 [Table 1].

Twenty-three were primary studies providing new symptom-severity scoring systems to assess food-induced allergic reactions. 1,4,19,20,22,24,26,27,29–33,42–50,58 [Tables 1 and 2]. We included eight additional symptom-severity scores from secondary literature (eg, editorials, rostrum, consensus reports, theoretical reviews, position papers). 15,51–57

Setting
The instruments have been designed and developed in several settings [Table 1]: allergy specialist centres (including clinical trials) 4,19,20,22,29,30,32,43,44,48,50 or emergency rooms 24,26,27 or intensive care unit 58 or general practitioner setting 33,47 or self (/parental) reported survey. 1,42,45,49 Three instruments have been designed for the context of OFC. 30,32,56

Targeted age group
Thirteen instruments have been originally created targeting pediatric allergic patients. All of them were primary studies. 1,19,20,22,24,31–33,43–45,47,50 One old primary study did not report age of participants. 58 The other 17 scores, including all of those reported in secondary literature, are applicable to all age groups. 4,15,26,27,29,30,42,43,46,48,49,51–57 [Table 1].

Targeted allergenic source
We included 17 symptom-severity scoring systems primarily designed to assess allergic reactions elicited by food 1,4,19,20,22,29–33,42–45,49,50,68 [Table 1] as follows: seven for any food; 1,19,27,32,33,43,45 six for peanut only; 4,20,30,31,49,50 one for peanut and nuts; 29 one for peanut, tree-nuts, fish, shell-fish, and sesame; 42 one for tree-nuts; 44 one for milk 22 but also applied to egg. 23 One score has been created to assess severity of food induced reactions occurring during oral immunotherapy. 43 We identified 11 instruments created to evaluate the severity of allergic reactions triggered by any allergenic source, including food. 15,24,26,46–48,51–54,57 One of them was designed by authors primarily for adverse reactions to subcutaneous immunotherapy (SCIT) but with the indication to be applied for any allergic reaction. 51 In particular, secondary literature, highlighted the need to identify a uniform instrument to be applied to any allergenic sources at any patient age by different stakeholders. 15,51–54,57 Our SR included also one symptom scoring system that was originally designed to assess the overall severity of an allergic reaction elicited from other allergic condition (ie, hymenoptera venom allergy) but that has been applied afterwards to classify food-induced allergic reactions. 58

Organ-specific symptoms
All included scoring systems had organ-specific outcomes covering the whole spectrum of clinical symptoms and signs in the context of IgE-
mediated allergic reactions. Several of them used the term “anaphylaxis” to describe the entire spectrum of severity, although non-anaphylactic milder symptoms neither fulfil the main current definitions of anaphylaxis nor the new ICD-11 criteria.

Of note, all scoring systems divided symptoms according to their anatomical involvement, ie, skin, respiratory, gastro-intestinal (GI), cardio-vascular (CVS), or neurological subjective symptoms/objective signs. Table 2 provides a detailed overview of the 30 included symptom-severity scores for each listed symptom ordered by organ. At least two reading levels of Table 2 are possible. A macroscopic evaluation suggests that there is overall concordance in assigning a progressive severity grade proceeding from the left (overall coloured in green) to the right side (in red colour), ie, spanning from skin involvement up to the lower respiratory tract, cardio- and neurological involvement passing by the gastrointestinal and upper respiratory tract (in yellow). However, a closer evaluation highlights the presence of some heterogeneities.

The majority of scoring systems used a detailed predefined list of symptoms, each of them presented as a dichotomous variable (ie, “present/non present”) or, in some cases, as a detailed grading of specific symptoms, (eg, urticaria, into mild/local or severe/generalized). A few used a more general "catch-all symptoms" approach for specific organ/system to embrace all possible symptoms for that specific organ/system (eg, all symptoms related to the “GI tract”).

All scoring systems utilized an ordinal scale ranging over 2-6 incomparable steps. The majority of them defined the overall allergic reaction severity based on the organ symptom(s) of the highest grade (ie, most severe symptoms). Some considered the number of organ-systems involved or the fulfilment of “2-or-more” symptoms/organ. Three scoring tools used a mathematical formula/summation of symptoms to obtain symptom severity regardless of number of observed symptoms. Two studies considered explicitly the need of medical treatment as a criterion for assessing the symptom-severity. One scoring system correlated the severity of any allergic symptom to the amount of exposed food allergen.

Studies investigating predictors for symptom-severity of food allergy

This SR assessed if the included studies identified any predictors for symptom-severity of food allergy. We found that 13 included primary studies reported on the assessment of host-related and food allergen-related factors, including demographic, clinical and/or laboratory variable(s), as predictors for severe allergic reaction to food.

Host-related factors

Three studies reported on the assessment of gender as a predictor for severity of allergic reactions; all of them reported no significant results.

Six papers evaluated age as a potential parameter associated with increased risk of symptom-severity; only two studies found that adolescence and adulthood are risk factors.

Asthma has been analyzed and reported as a predictor for severe ARs by five studies describing this analysis, one reported that patients with a clinical history of asthma were more likely to suffer severe ARs (χ² = 17.9, P.00013) and, of note, wheeze as the most common severe symptom of AR (~40% of pts). In another study, the frequency of severe ARs compared with moderate, mild, or no ARs was 10-fold higher in asthmatic children but did not reach statistical significance (OR, 10.19; 95% CI, 1.13–91.54; P .022).

One study evaluated the concomitant use of drugs ie, ACE inhibitor and β-blocker) but no correlation with symptom severity was found.

Three studies evaluated the role of recurrence of ARs as a predictor. The symptom-severity of the previous AR(s) or the first AR did not significantly predict the symptom-severity of the next, in the two studies reporting on this outcome. Similarly, one study evaluating if a previous reaction to peanut in the clinical history predicted symptom severity in peanut allergic patients found no correlation.
Food allergen-related predictors

Two of the included studies assessed the type of food as potential risk factor.\textsuperscript{19,39} One found that wheat was the only predictor of severe anaphylaxis (OR 2.425, 95% CI 1.054–5.581, p < 0.037).\textsuperscript{39} The second study found the highest (but statistically non-significant) risk of severe ARs for peanuts (OR = 1.76, 95%CI: 0.9–3.45) and shellfish (OR = 1.54, 95%CI: 0.49–5.64) and the lowest for sesame, soy, and wheat.\textsuperscript{19}

One study reported total IgE level as a protective factor.\textsuperscript{23} Total IgE levels were significantly lower in patients with moderate/severe ARs (adjusted odds ratio for every 1-unit increase in the decimal logarithm, 0.16; 95% CI, 0.05–0.54; \( P = .001 \)).

On the other side, five out of the eight studies\textsuperscript{20–23,30,31,36,38} reporting on serum level of specific IgE (sIgE) as a predictor for symptom severity found it as a risk factor.\textsuperscript{21–23,30,38} The remaining three studies found no significant results for sIgE as predictor.\textsuperscript{20,31,36} Of note, the five studies assessed specific IgE level to different culprit foods: sIgE to cow’s milk and to casein;\textsuperscript{22} sIgE to egg white;\textsuperscript{23} and sIgE to whole peanut proteins,\textsuperscript{21,30} and rAra h 1, rAra h 2.\textsuperscript{21} In particular, one study reported that sIgE to peanut and challenge score correlated significantly in the whole group but this correlation was stronger in adults than in children, despite the median values of peanut sIgE being similar; in adults Spearman’s \( r \)-value increased to 0.766 (\( P = 0.001 \), compared with children (\( r = 0.49 \), \( P = .018 \)).\textsuperscript{30}

Another study reported that age, sIgE and SPT to almond at challenge when combined demonstrated good predictive value for grade 2/3 allergic reactions by AUC (area under the curve, 0.83).\textsuperscript{38}

A further study found that patients monosensitized to rAra h 2 had a significantly lower severity score than those polysensitized to the same source (i.e. rAra h 2 and rAra h 1 and/or rAra h 3) (\( P < .02 \)).\textsuperscript{20} Two studies reported on SPT itself with no significant results.\textsuperscript{20,31}

Quality of life studies

Our SR identified 7 papers that met our inclusion criteria, namely that the studies used a validated scoring system to measure FAQL, and reported how this scoring is impacted by "severity" (symptoms/anaphylaxis). All papers included participants with confirmed FA by specialist/allergist [Table S4]. We note here that the majority of recent papers not included in the review investigated the impact of Allergy Immunotherapy on FAQL.

Setting and population samples

The majority of studies recruited participants through allergy specialty clinics,\textsuperscript{59–61,64} and two studies also recruited through general medical clinics, community support groups and media advertisements.\textsuperscript{62,63,65} All studies took place in The Netherlands, Ireland, and United States. The measures used were distributed through hospital allergy clinics either on site or online through the clinic to patients diagnosed with food allergy (or parents of patients diagnosed with food allergy).

Measures

All studies used a validated age appropriate version of the Food Allergy Quality of Life Questionnaires (FAQLQ) which are recommended as gold standard by EAACI. The FAQLQ includes questions on demographics, symptoms, reaction history, diagnosis, prescription, and use of an auto-injector. The FAQLQ also incorporates the Food Allergy Independent Measure (FAIM) which assesses the perception of severity/chance of adverse outcome, if an allergen is accidentally ingested. FAIM also operates as an anchor instrument for the FAQLQ. The instruments used were designed for data collection in general and treatment settings, cross-sectionally and longitudinally, and have reported a minimal clinical important difference (MCID) score of 0.45/0.5.

The version of the FAQLQ chosen reflected the population(s) targeted in the study. The Parent Form was used in two studies,\textsuperscript{59,60} the Child Form (CF) and Teen Form (TF) were used in three studies\textsuperscript{59,61,64} and the Adult Form in three studies.\textsuperscript{51,62,64} One study used only the FAIM section of FAQLQ,\textsuperscript{53} with all other studies using the FAIM in addition to the FAQLQ, and \textsuperscript{1} used the Parental Burden (PB) version of FAQLQ.

In addition to FAQLQ, generic measures were used to measure outcomes in three studies, namely Parental Empowerment Scale,\textsuperscript{65} CHQ-
CF87 and Rand-36, Food Insecurity Scale (FIS), and use of food assistance programs (SNAP, food banks).

Severity

At minimum to satisfy the inclusion criteria, all studies included questions on reaction history (eg, a list of symptoms reflecting all levels of severity), diagnosis (eg, how and by whom a patient had been diagnosed) and whether an epinephrine auto-injector (EAI) had been prescribed and reported how FAQL is impacted by "severity" (eg, symptoms/reactions). Severe food allergy was defined as having a prescription for an EAI, or self-reported previous episodes of anaphylaxis (ie, the symptoms “difficulty breathing”, “inability to stand”, collapse and/or loss of consciousness).

Outcomes targeted

The majority of the studies were carried out for psychometric purposes, specifically to assess the longitudinal validity and responsiveness of the FAQLQ-AF, FAQLQ-TF, FAQLQ-CF and the FAQLQ-PF and cross-cultural validity of the adult form (AF) and parent form (PF) respectively, and one study compared FAQL measured with generic and disease-specific questionnaires. The impact of a food challenge on FAQL was evaluated in three studies. Relationships between allergen severity, type, or comorbidities and FAQL was the focus of 2 in the context of parental empowerment, and uncertainty or inability to meet family food requirements (FIS).

Findings/results

FAQLQ was found to be responsive to change in a food-allergic patient population with disease-specific clinical outcomes with good cross-cultural validity. All studies identified positive associations between FAQLQ impact on was found according to severity, positive challenge result, number of allergens avoided, and number of symptoms. The FAQLQ of American food-allergic adults was found to be more impaired than Dutch food-allergic adults and Irish food-allergic children. Caregivers classified as FIS reported an increased perceived risk of accidental ingestion, severe reaction, and death, and it was also associated with utilization of food assistance programs and food banks. Mothers reported greater empowerment and worse FAQL compared with fathers, regardless of allergen severity, type, or comorbidities, but was not significantly associated with FAQL for mothers or fathers. Highest FAQLQ-PF impact was for items involving fear of allergen exposure outside the home.

Economic burden

A SR was conducted to identify and summarise evidence regarding economic analyses of food allergy severity. Of the final articles selected for full screening, two met inclusion criteria. Articles that did not identify grades of food allergy severity (mild, moderate and severe) were excluded.

The first study was based on 402 cases of severe anaphylaxis reported by the Allergy Vigilance Network, in years 2004, 2005, and 2006. The setting was hospital and general practices in France. International classification of Diseases codes for anaphylaxis used in the study, included T780 (shock due to adverse food reaction), T782 (anaphylactic shock, not specified), T805 (shock due to serum/vaccine/immunization), T886 (shock due to adverse drug reactions), and T882 (anaesthetic shock). Direct and indirect costs were estimated from a national perspective. Direct costs consisted of the costs of medications, consultations, use of emergency units, diagnosis, and hospitalisations, as well as nonmedical costs such as transport, and diet. Indirect costs were based on the costs of absenteeism with a mean of three days (two days at the time of event, and one day after an event). Indirect cost data was calculated on the basis of Belgian costs.

Direct medical costs

Results indicated that the average direct cost was €1580 per patient, ranging from €74.88 to €4445.47 (as the currency year was not indicated, it was assumed to be one year prior to the year of the publication). Costs were equivalent in purchasing power to €1,889 per patient, with a range of €89.51 to €5,314.08, in year 2020. Table S8 includes the direct medical costs for severe anaphylaxis management obtained from Flabbee.
et al, 2008, and adjusted to the currency year of 2020. The hospitalization had the greatest cost, ranging from €239.08 for Emergency ambulance brigade called, to severe cases of hospitalization in intensive care unit with an average cost of €2,528.25/day.

Indirect costs

Indirect costs were estimated to be €315 per patient; equivalent to €376.55 in year 2020.

Total costs

The total average cost per patient was €1895, equivalent to the cost of €2,265.27 in 2020.

The second study, investigated health service costs for food allergic individuals in Europe (Greece, Iceland, Poland, Spain, Czech Republic, France, Italy, The Netherlands, and United Kingdom), and the relationship between severity and the cost of illness. The time frame was from January 2007 until July 2009. The Geary-Khamis dollar (I$) was used to estimate unit costs of services at 2016 prices. The setting was in general practitioners’ patient lists, city council registration databases, local authority/hospital databases, and primary schools. Participants were recruited through the EuroPrevall study in a case-control study design, and completed an economic questionnaire. Participants with possible food allergy were identified by clinical history, and those with sIgE were defined as having probable allergy. Results indicated that the average health care cost for adults with possible food allergy was I$2016 (equivalent to €1 933.61 in year 2020) and I$1089 (€1 044.49 in year 2020) for controls. For children aged 7–11 years, the costs are I$2197 (€2 107.2 in year 2020) for those with possible food allergy and for controls it was I$863 (€827.73 in year 2020). The mean average yearly cost of possible and probable food allergy was I$1778 (€1705.33 in currency year 2020) in 9 participating centres. The study indicated that the cost of treating individuals with moderate allergy symptoms was 68% higher than for those with mild symptoms. Health care costs for those with severe food allergy were estimated to be double the amount for those with mild food allergy. No significant differences in health care costs were observed for children when compared with adults.

DISCUSSION

Summary of main findings

To our best knowledge, this mixed methods SR represents the most comprehensive investigation ever undertaken of literature on current classification of food allergy severity.

We have tried to cover all the perspectives of food allergy severity from different stakeholders, including patients and parents/families, patient advocacy groups, disease registries, health professionals, researchers, academicians, food and drug industries, government agencies and regulators, as well as legislative bodies, as they all perceive the concept of severity differently. All included studies were observational studies that investigated symptom-specific and non-symptom specific severity of food allergy in children and adults.

- A consensus on definition of food allergy severity is missing. This systematic review is the preliminary step towards a state-of-the-art synopsis of the current evidence relating to the severity of IgE-mediated food allergy. It will inform attempts to develop a consensus to define food allergy severity by clinicians and other stakeholders. Each participating stakeholders perspectives on food allergy severity has been covered.
- All included studies were observational studies that investigated symptom-specific and non-symptom specific severity of food allergy in children and adults.
- The overall body of epidemiological evidence in relation to the food allergy severity classification is moderate.
- There is heterogeneity among severity scoring systems used and even outcomes considered in the context of severity of food allergy. No score has been validated.
- To assess comprehensively predictors of severity of food allergy is urgently required in order to develop and use worldwide the best prediction model for severe food allergy.
- Research into FAQL has helped to raise awareness of patient issues and provided a means of individualized assessment for a patient or parent.
- There is lack of full economic evaluation studies on the severity of food allergy.
- Shared decision making is needed to address all issues regarding the definition of food allergy severity from each stakeholder’s perspectives.

Table 3. Summary of the DEFASE systematic review
We found that many severity scoring systems for food allergic reactions have been generated; however, they are heterogeneous and none of them has been validated in practice. They differ for number of steps and are only partially interchangeable. No standardized or validated method exists to compare multiple heterogeneous scoring systems. The inconsistency and non-validity of these scoring systems highlights the urgent need to develop a harmonised, consensus-based definition for the severity of food allergy in children and adults useful for all stakeholders involved.

The severity spectrum of an allergic reaction can range from subjective local mild symptoms to fatal anaphylactic shock. Type of allergen, dosage, individual threshold, route of exposure, age, comorbidity, and involvement of cofactors may influence the severity of a food allergic reaction. In turn, these variables make severity difficult to capture. Furthermore, onset and severity of each symptom can lead to progression and interaction of symptoms. Allergic reactions can occur in different incomparable settings: they range from accidental exposure in an unknown environment to controlled titrated oral food challenges in a highly specialized clinical setting.

The variability between instruments included in this SR was overall wide: some instruments are purposed solely for single allergens (e.g., peanut), others developed exclusively for specific populations (e.g., children) and some to specific situations (e.g., OFC).

At present, the global severity of a food allergic reaction is generally either based on the highest/most severe reported symptoms or calculated by different algorithms. Some instruments used “catch-all” definitions in contrast to others based on a predefined “symptom list”, which contains more information for research purposes, and avoids the pitfall of overlooking milder symptoms.

Research into FAQL has helped to raise awareness of patient issues and provided a means of individualized assessment for a patient or parent. If a validated consensus-driven severity scoring system for FA could be developed, it could be used to harmonize outcome assessment in clinical trials and also facilitate understanding of important determinants of FAQL. This could widen the parameters of benefits and harms of new treatments and allow for the development and improvement in process and outcome quality indicators. A standardised protocol that incorporates FAQL, severity, and agreed definitions of outcomes would allow for the comparison of efficacy of food allergy treatments between centres, trials or countries. The use of such measures can help to differentiate between levels of severity on multi-dimensional patient outcomes. To this
end, it is vital that improved FAQL is seen as a primary outcome, and is measured at multiple intervals during trial duration and beyond. In addition, few studies have used the minimal clinical important difference (MCID) when assessing change in FAQL. The MCID can help determine the effect of a given therapy on a patient and add meaning to statistical inferences made in research; therefore, their use is critical for the conduct and interpretability of clinical trials.

Decision science modelling is another method that could help us understand variations in preferences for treatment, which could affect the health and economic impact of allergen immunotherapy (AIT). Assessment of patient/caregiver attribute preference and how this translates to health economic outcomes will provide a basis to understand if strategies used in food allergy can deliver value-based care, which can be applied to the development of future food allergy research. Utility valuations should be derived from responses to FAQLQ instruments, providing more accurate measurement of this construct.

Strengths and limitations of this work

To our knowledge, this is the first SR of the literature on current severity classifications used for FA. The strengths of this SR are the comprehensiveness of the searches, including all available sources of 11 international electronic databases without any geographical restrictions and with higher methodological rigour. In addition, we were able to contact an international panel of experts. We carefully grouped and categorised food allergy severity as either symptom-related or non-symptom related under several categories to find the effect of each compounds on the severity of allergic reactions to culprit food.

The methods used to verify symptom specific and non-symptom specific measures of food allergy severity were carefully assessed and graded for methodological rigour.

The main limitations of this systematic review stem from the substantial heterogeneity of studies with moderate methodological quality, to the fact that the included studies were only observational (cohort, case-control, or cross-sectional) with no interventional studies.

CONCLUSIONS

The overall body of epidemiological evidence in relation to the food allergy severity classification is moderate. We found only observational studies, data are generally of moderate quality. This systematic review confirms the variability and diversity of severity scoring systems for FA. Overall, in terms of symptom severity systems there is a general suggestion that cardiovascular and lower respiratory tract reactions are severe and the cutaneous, and gastrointestinal ones are mild to moderate. Quality of life and economic evaluation of severity of food allergy should be incorporated into the food allergy severity definition alongside the symptom score assessment. Unfortunately, at present there is no validated and broadly accepted categorization of severity of food allergy that can be used by all stakeholders (patients, family members, guardians, healthcare professionals, researchers, food industry, policy makers and other public health authorities). A validated severity scoring system for FA could be used both for standardised patient monitoring and also to define the eligibility of allocating patients for clinical studies. This review also will represent a preliminary work for generating a consensus-based definition of severity of food allergy in children and adults, developing and implementing the algorithm by a multidisciplinary panel of experts.

There is an excess of severity scoring systems for allergic reactions including to food in children and adults. The usability of these severity scoring systems remains unclear because of methodological shortcomings, incomplete presentation, lack of internal and external validation, and testing for reliability and validity of the severity scoring systems in a range of settings and populations. The standardised, harmonised, and consensus-based uniform definition of severity scoring systems that will be used by all stakeholders is urgently needed. Rather than developing yet another severity scoring system for allergic reactions, future research should focus on external validation of scoring systems, tailoring of these models to different allergens, populations, and settings. In addition, as a gold standard, a standardised, harmonised, consensus-based severity scoring system for food allergy needs to be tested for reliability and validity in a range of settings and
populations. To reach this crucial goal, expert consultation, e-Delphi study and impact studies will be the main platforms in the risk assessment and risk management of patients with food allergy [Table 3].

**IMPORTANCE TO STAKEHOLDERS AND IMPLEMENTATION**

The concept of food allergy severity is important not only for healthcare professionals in evaluating patients, but also for patients’ family members, food and drug industries, research (clinical trials, epidemiologic, genetic, immunological, and mechanistic studies), government agencies and regulators, as well as for policy makers. The terminology and definitions currently applied to food allergy severity are not standardized, and often misleading. Furthermore, different stakeholders perceive the concept of severity differently. Therefore, a common approach is needed for an international consensus-based system to define food allergy severity and our mixed-methods systematic review comprehensively assesses and addresses this crucial issue in the management of food allergy.

**FUTURE RESEARCH**

This SR will be a background work to reach an international consensus on the definition of food allergy severity in children and adults. As described in the roadmap (see Fig. 2), the next step will be to conduct expert consultation, through an e-Delphi study and by taking into consideration the perspectives from the different stakeholders involved. After developing the consensus document, there is a need for well-designed clinical impact studies by using large clinical databases that test the reliability and validity of severity scoring systems for food allergy. These clinical impact studies may then need to be followed up via well conducted large RCTs to evaluate the correct usage of consensus-based definitions of food allergy severity, effectiveness and cost-effectiveness of interventions aiming to reduce food allergy severity burden and risk of developing its complications in the future.

**Abbreviations**

ACE, Angiotensin-Converting Enzyme; AIT, Allergen Immunotherapy; AMED, Allied and Alternative Medicine; AR, Allergic Reaction; AUC, Area Under Curve; CAB, the Commonwealth Agricultural Bureaux; CF, Child Form; CINAHL, Cumulative Index to Nursing and Allied Health Literature; CASP, Critical Appraisal Skills Programme; CBA, controlled before after studies; CCT, controlled clinical trials; CI, confidential interval; CoFAR, Consortium of Food Allergy Research; CVS, Cardio-vascular System; DBPCFC, Double-blind placebo-controlled food challenge; DEFASE, Definition of Food Allergy Severity; EAACI, European Academy of Allergy and Clinical Immunology; EAI, epinephrine auto-injector; EconLit, Economics Literature; ED, Emergency Department; EMBASE, Excerpta Medica database; EMTREE, EMBASE Subject Headings; EPHPP, the Effective Public Health Practice Project; FA, IgE-mediated food allergy; FAQL, food allergy quality of life; FAQLO:AF, Food allergy quality of life questionnaire-adult form; FAQLO:CF, Food allergy quality of life questionnaire-child form; FAQLO:PF, Food allergy quality of life questionnaire-parent form; FAQLO:TF, Food allergy quality of life questionnaire-teen form; FAIM, food allergy independent measure; FARE, Food Allergy Research & Education; FIS, Food Insecurity Scale; GI, Gastro-intestinal; GRADE, Grading of Recommendations Assessment, Development and Evaluation; HRQL. Health related quality of life; ICU, Intensive Care Unit; iFAAM, Integrated Approaches to Food Allergen and Allergy Risk Management; ISI, the Institute for Scientific Information; ITS, interrupted time series; MCID, Minimal clinical important difference; MEDLINE, Medical Literature Analysis and Retrieval System Online; MeSH, Medical SubHeadings; OFC, Oral Food Challenge; OR, Odds Ratio; PF, Parent Form; PRACTALL, PRACTical ALLergy; PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analyses; PROSPERO, Prospective Register of Systematic Reviews; QoL, Quality of Life; RCT, randomized controlled studies; SCIT, Subcutaneous Immunotherapy; SNAP, the Supplemental Nutrition Assistance Program; SPT, Skin Prick Test; SR, systematic review; TF, Teen Form; TRIP, Turning Research Into Practice; USA, the United States of America; WAO, World Allergy Organization

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Appendix A. Supplementary data
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Author details
*Allergy Unit - Area of Translational Research in Pediatric Specialities, Bambino Gesù Children’s Hospital, IRCCS, Rome, Italy. †Division of Population Medicine, School of Medicine, Cardiff University, Wales, UK. ‡Applied Psychology and Paediatrics and Child Health, University College Cork, Cork, Ireland. §Nuffield Department of Primary Care Health Sciences, University of Oxford, Radcliffe Observatory Quarter, England, UK. ¶NIHR Southampton Biomedical Research Centre, University Hospital Southampton NHS Foundation Trust, Clinical and Experimental Sciences and Human Development in Health, Faculty of Medicine, University of Southampton, Southampton, UK. ¶¶The David Hide Asthma and Allergy Research Centre, St Mary’s Hospital, Isle of Wight, UK. ¶¶¶National Heart & Lung Institute, Imperial College London, London, UK. ¶¶¶¶Discipline of Paediatrics and Child Health, School of Medicine, University of Sydney, Sydney, Australia. ¶¶¶¶¶Division of Pulmonary, Allergy, and Critical Care Medicine, Department of Medicine, Stanford University, Sean N. Parker Center for Allergy and Asthma Research at Stanford University, Stanford University, Stanford, CA, USA. ¶¶¶¶¶¶Center for Food Allergy and Asthma Research (CFAAR), Northwestern University Feinberg School of Medicine, Department of Pediatrics & Medicine, Ann & Robert H. Lurie Children’s Hospital of Chicago, USA. ¶¶¶¶¶¶¶Pediatric Allergy Unit, Department of Women-Children-Teenagers Pediatries, University Hospitals of Geneva, Geneva, Switzerland. ¶¶¶¶¶¶¶¶Allergy and Immunology, Department of Pediatrics, New York University School of Medicine, Langone Health, New York, NY, USA. ¶¶¶¶¶¶¶¶¶Department of Pediatrics, Gastroenterology and Nutrition, Collegium
Medicum, University of Warmia and Mazury, Olsztyn, Poland. 8Department of Paediatrics, Imperial College London, United Kingdom. 9Department of Paediatrics, The Chinese University of Hong Kong, Prince of Wales Hospital, Shatin, Hong Kong, China. 10Clinical Research Center for Allergy and Rheumatology, National Hospital Organization, Sagamihara National Hospital, Kanagawa, Japan.

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