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Psychometric properties of disease-specific health-related quality of life (HRQoL) instruments for food allergy and food intolerance: protocol for a COSMIN-based systematic review

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ABSTRACT

Introduction Food allergies and food intolerances can bring burdens on patients and their caregivers and reduce health-related quality of life (HRQoL). An increasing number of disease-specific HRQoL instruments for food allergies and food intolerances has been developed, and some of them have been adapted for different cultures and languages. This report describes a protocol for a systematic review of the psychometric properties of these instruments. The aims of this systematic review are to: (1) formulate recommendations for the usage of existing validated disease-specific HRQoL instruments for patients with food allergies and/or food intolerances and their caregivers; and (2) identify knowledge gaps to inform future research relating to these instruments.

Methods and analysis This protocol adheres to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) Protocol 2015 checklist. The future review will follow the Consensus-based Standards for the Selection of Health Measurement Instruments (COSMIN) guideline for systematic reviews of patient-reported outcome measures (PROMs) and PRISMA 2020 statement guideline. Six databases (PubMed, EMBASE, Web of Science, Scopus, CINAHL and ProQuest -Health & Medical Collection) will be searched to retrieve studies focusing on the development and psychometric properties of disease-specific HRQoL instruments for patients with food allergies and/or food intolerances and their caregivers between 1 December 2021 and 31 December 2021. Two researchers will be responsible for literature screening, data extraction and literature evaluation, independently. Disagreements will be addressed by discussion or the involvement of a third researcher. The methodological quality of the included studies and the quality of the identified instruments will be assessed based on the COSMIN guideline for systematic reviews of PROMs.

Ethics and dissemination Ethical approval is not applicable for this study. We will disseminate the findings through publication in peer-reviewed journals and/or academic conferences.

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INTRODUCTION

Adverse reactions to food encompass food allergies and food intolerances. Food allergies and food intolerances are common, especially in children, with 6.25%–28.0% of children having had this experience. Food allergies and food intolerances have become severe public health problems worldwide. A food allergy is an abnormal immunological reaction to specific food(s) that results in the development of symptoms. Food allergies include three types of immunological reactions: IgE-mediated, non-IgE-mediated and Mixed IgE- and non-IgE-mediated. Food intolerances are non-immune mediated adverse reactions to food; lactose intolerance...
is one such example. Food allergies and food intolerances impact multiple organs and systems, and are associated with a range of symptoms (e.g., urticaria, eczema, colic, vomiting, reflux, diarrhoea or constipation, blood in stool and growth retardation). Meanwhile, food allergies may lead to life-threatening anaphylaxis.

Adverse food intake reactions bring great challenges to healthcare, education, food and catering industries in many countries. Critically, the physiological, psychological, social and financial burdens relating to food allergies and food intolerances also undermine health-related quality of life (HRQoL) of patients and their family caregivers. Instruments measuring HRQoL can help quantify the impacts of food allergies and food intolerances, and may support better prevention and management of this problem. For example, healthcare professionals could use these instruments to assess the quality of life of patients and their caregivers.

There are two types of HRQoL instruments: generic and disease-specific. Disease-specific HRQoL instruments are more likely to have a higher level of sensitivity compared with generic HRQoL instruments. A number of disease-specific HRQoL instruments for food allergies has been developed. Some of them have been validated and adapted into multiple versions for different cultures and languages. Since 2014, there have also been some studies reporting on the overall development of instruments for food intolerances. Two literature reviews relating to HRQoL instruments for food allergies were published in 2009 and 2014, respectively. The 2009 review summarised and described generic and disease-specific instruments for food allergies in children and adults. However, this review was limited by its use of a narrative review approach, rather than a systematic review following corresponding guidelines, which can lead to the omission of some important literature. The 2014 review systematically summarised and evaluated all disease-specific HRQoL instruments for IgE-mediated food allergies. However, this systematic review failed to include other types of food allergies (non-IgE-mediated, and Mixed IgE- and non-IgE-mediated), as well as food intolerances. These omitted types of adverse reactions to food also have significant influences on quality of life of patients and caregivers, and as such, should be included in a broader systematic review of the literature. Furthermore, the 2014 review did not follow specific guidelines for evaluating the methodological quality of the included studies and the quality of the included instruments. In 2018, the Consensus-based Standards for the Selection of Health Measurement Instruments (COSMIN) guideline was developed for systematic reviews of Patient-Reported Outcome Measures (PROMs). This guideline could improve the quality of the systematic review of PROMs, offering researchers a critical and comprehensive evaluation of the available instruments.

Therefore, the overall aim of this systematic review is to critically describe, appraise, and summarise the existing disease-specific HRQoL instruments for patients with food allergies and/or food intolerances and their family caregivers, based on the COSMIN guideline for systematic reviews of PROMs. The specific objectives of the proposed systematic review are to: (1) identify and describe all existing validated disease-specific HRQoL instruments for patients with food allergies and/or food intolerances and their caregivers; (2) evaluate the methodological quality of studies on measurement properties of the instruments and (3) assess and compare the psychometric properties and other key characteristics of these instruments.

This systematic review will answer the following questions: (1) What are existing disease-specific HRQoL instruments for patients with food allergies and/or food intolerances and their caregivers? (2) What are the characteristics of these instruments? (3) What is the methodological quality of studies on measurement properties of these instruments? (4) What are the measurement properties, interpretability and feasibility of these instruments? (5) What are the similarities and differences among these instruments? and (6) What are the knowledge and research gaps in this area?

**METHODS**

This is a protocol for a systematic review following Preferred Reporting Items for Systematic Reviews and Meta-Analyses Protocols (PRISMA-P) 2015 checklist. Based on COSMIN guideline for systematic reviews of PROMs, we adapted the ‘Objectives’ section of the PRISMA-P 2015 checklist by replacing ‘Participants, Interventions, Comparators and Outcomes’ with ‘Construct, Population(s), Type of Instrument(s) and Measurement properties’. We registered the protocol in the International Prospective Register of Systematic Reviews (PROSPERO), and the title of the protocol has been registered on the Joanna Briggs Institute website. We will conduct this systematic review based on the COSMIN guideline for systematic reviews of PROMs and report it following the updated PRISMA 2020 checklist.

**Inclusion and exclusion criteria of studies**

**Inclusion criteria**

Studies will be included if they: (1) report disease-specific HRQoL instrument(s) designated for patients with food allergies and/or food intolerances, and/or their caregivers; (2) describe the processes of development and/or evaluation of one or more measurement properties for eligible instrument(s) and (3) have full-text availability. The authors of the articles will be approached if a full-text version is not available online. However, if the authors’ contact information is not available or the authors do not respond to the inquiry, these studies will be excluded but their information will be recorded in online supplemental file of the formal systematic review.

Exclusion criteria
Studies will be excluded if they: (1) are not primary studies (e.g., discussion papers, letters and editorials) or case studies or (2) are reports that used the instruments only for outcome measurement.

Search strategy
Between 1 December 2021 and 31 December 2021, we will search PubMed, EMBASE, Web of Science, Scopus, CINAHL and ProQuest (Health & Medical Collection) using comprehensive and sensitive search strategies that combine Medical Subject Heading and free text words. All databases will be searched from the date of inception to the date of searching. The major search concepts will be quality of life (Construct), food allergies/food intolerances (Population), PROMs (Type of instrument(s)) and measurement properties. Three comprehensive and sensitive keyword search strategies developed by other researchers for reviews of the concepts will be used in this literature search. They are: (1) the search filter of ‘quality of life’ for medical and health bibliographic databases developed by Dutch medical information specialists; (2) the search filter for finding PROMs developed by the University of Oxford; (3) the sensitive PubMed search filter for measurement properties developed by Terwee et al, and corresponding search filters adapted for other databases. These literature search filters will improve the comprehensiveness, effectiveness and quality of the literature search in this study. Furthermore, a health science librarian had been consulted for developing the search strategies. Online supplemental table S1 and online supplemental tables S9–S13 show the search strategies we developed for the databases searched. The search will not be limited to a specific language; that is, we will include eligible publications in any language, and a translation service will be used if needed. Database searches will be carried out again to provide a final update of the searches after the systematic review is accepted by a journal. The systematic review will be updated if new eligible studies are identified.

Study screening
Endnote and Covidence will be used to manage the references screening. First, we will use EndNote to recognise and remove duplicates, and then conduct manual screening. Following this initial screening, titles, abstracts and full-text articles will be reviewed and screened independently by two researchers with the support of Covidence. Disagreements between the two researchers will be addressed through discussion. Consultation of a third researcher will be adopted where necessary. Reference lists from all eligible papers will also be screened using the aforementioned inclusion and exclusion criteria. The processes of study screening are shown in figure 1.

Data extraction
Two researchers will conduct the data extraction independently. A third researcher will review the extracted data and address the discrepancies between the two researchers if identified. We will extract data on (1) basic characteristics of the included instruments (online supplemental table S2, including: the name of the instrument, developer(s)/year developed, construct(s), targeted population, mode of administration, recall period, (sub) scale(s)/(number of items), response options, range of scores/scoring, original language and available translations); (2) characteristics of the included study populations (online supplemental table S3, including sample size, age, gender, disease, disease duration and severity, setting, country, language); (3) results of measurement properties of the included instruments (Result columns in online supplemental table S5, including content validity, structural validity, internal consistency, cross-cultural validity, measurement invariance, reliability, measurement error, criterion validity, construct validity, responsiveness and (4) interpretability (online supplemental table S7) and feasibility (online supplemental table S8) of the included instruments.

Quality appraisal and data synthesis
Two researchers will conduct the quality assessment for included studies and instruments independently. A third researcher will be consulted if consensus could not be reached. The COSMIN guideline will be used to assess each subscale of a multidimensional PROM separately. Therefore, the measurement properties for subscale scores and the entire PROM will be rated separately in this study.

In the first step, COSMIN standards for design requirements and preferred statistical methods will guide the evaluation of the methodological quality of the included studies on the development and measurement properties of the instruments. The following COSMIN resources will be used in this phase: the COSMIN Risk of Bias checklist for PROMs, the COSMIN methodology for systematic reviews of PROMs—User manual, and the COSMIN methodology for assessing the content validity of PROMs—User manual. In this step, an Excel sheet file named ‘Scoring form COSMIN boxes’ will be used to manage the evaluation records (refer to online supplemental additional file 3; this file is also available at https://www.cosmin.nl/tools/guideline-conducting-systematic-review-outcome-measures/); this file is provided by the COSMIN guideline. The final consensus on the results of the methodological quality will be presented in online supplemental table S4 and Meth qual column in online supplemental table S5.

In the second step, we will evaluate the results associated with measurement properties of identified instruments according to the COSMIN quality criteria for good measurement properties. The corresponding results will be reported in the rating columns in online supplemental table S5. However, the rating results of content validity will be separately presented in online supplemental table S5-1 given that criteria and rating systems...
for evaluation of content validity of PROMs differ from other measurement properties in COSMIN guideline. The COSMIN guideline also provides a separate table (online supplemental table S6) to synthesise evidence and results associated with measurement properties. In the third step, we will statistically pool or qualitatively summarise the results on measurement properties from different studies provided and show the summarised or pooled results in the column of Summary or pooled results of online supplemental table S6. A meta-analysis approach (weighted means and 95% CIs) will be used when possible. We will evaluate the pooled or summarised results per measurement property for each PROM according to the COSMIN quality criteria for good measurement properties; the corresponding results will be shown in the Overall rating columns in online supplemental table S6.

Finally, we will use the modified GRADE (Grading of Recommendations Assessment, Development and Evaluation) approach developed by the COSMIN guideline for systematic reviews of PROMs to grade the quality of the evidence. The quality of the evidence will be defined as the level of confidence based on the level of trustworthiness of the pooled or summarised result (shown in Quality of evidence columns in online supplemental table S6). The COSMIN guideline classifies the quality of the evidence into four levels: ‘high’, ‘moderate’, ‘low’ or ‘very low’. These findings will enable us to formulate recommendations on the usage of existing disease-specific HRQoL instruments for patients with food allergies and/or food intolerances and their caregivers. Our findings will also identify knowledge gaps in this area and inform future research.

**Patient and public involvement**

Neither patients nor the public will be involved in this study.

**Ethics and dissemination**

Ethical approval is not applicable for this study. We will share the findings from the study at national and/or international conferences and in a peer-reviewed journal in the fields of food allergies or food intolerances.

**DISCUSSION**

To our knowledge, this review will be the first PRISMA and COSMIN guidelines-guided systematic review of disease-specific HRQoL instruments for patients with food allergies and/or food intolerances and their caregivers. This review will identify, describe, evaluate and compare all eligible instruments. The methodological quality of all included studies on the measurement properties of these instruments and the psychometric properties of

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**Figure 1** Flow chart of literature selection process. PROM, patient-reported outcome measure.
all included instruments will be evaluated based on the COSMIN guideline for systematic reviews of PROMs. The findings will facilitate the formulation of the recommendations on the usage of the targeted instruments for clinical practice and research. We will also identify knowledge gaps associated with measurements of HRQoL for patients with food allergies and/or food intolerances and their caregivers. This review has the potential to clearly identify opportunities for further research, and therefore supports future studies on the development and improvement of disease-specific HRQoL instruments for these populations.

REFERENCES