How do health professionals translate evidence on early childhood allergy prevention into health literacy-responsive practice? A protocol for a mixed-method study on the views of German health professionals

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ABSTRACT

Introduction Paediatricians, general practitioners (GPs) and midwives in primary care are important sources of information for parents on early childhood allergy prevention (ECAP). Research has shown that preventive counselling by health professionals can be effective in improving patients’ health literacy (HL) and health behaviour. Providing effective advice relies on two factors. First, health professionals need be up-to-date with research evidence on ECAP, to consider popular misconceptions and fears and to translate this knowledge into clear recommendations for parents (knowledge translation). Second, they need to know and apply counselling techniques and create a practice setting which accommodates parental HL needs (health literacy-responsive care). The objective of this study is to explore and assess how German health professionals take up and translate ECAP evidence into appropriate recommendations for parents, how they consider HL in counselling and practice organisation and what barriers and enablers they find in their performance of HL-responsive ECAP.

Methods and analysis The study has a sequential mixed-method design, in two phases. In the first phase, qualitative semi-structured expert interviews will be conducted with health professionals (paediatricians, GPs and midwives) at primary care level and professional policy level. Data collection is ongoing until January 2022. In the second phase, based on the qualitative results, a standardised questionnaire will be developed, and pilot-tested in a wider population of German health professionals. The findings of both phases will be integrated.

Ethics and dissemination The study has received ethical approval from the Ethics Committee of the University of Regensburg (18-1205-101). The results will be published in international peer-reviewed open access journals and via presentations at scientific conferences. The results will also be shared with German health professionals, decision-makers and potential funders of interventions.

INTRODUCTION

Health professionals’ role in early childhood allergy prevention

In Germany, paediatricians, general practitioners (GPs) and midwives in primary outpatient care are important sources of information on infant feeding, hygiene and early childhood allergy prevention (ECAP) for expectant and new parents.

These health professionals have regular contact with parents: more than 90% of families use the paediatric health screening programme (U-examinations) for children younger than 2 years, which is usually performed by paediatricians, but can also be conducted by GPs. In addition, German
mothers with statutory health insurance are entitled to prebirth and postbirth care by midwives, which in Bavaria is used by 65% of mothers prebirth and 94.9% postbirth.² Midwives also offer antenatal education courses about childbirth and parenthood for pregnant women and their partners. These courses cover breastfeeding and infant care,³ and therefore are relevant to ECAP.⁴

When counselling parents on ECAP, a health professional should advise them, for example, to keep the home smoke-free, to avoid mildew growth and to introduce solid food gradually after at least 4 months of exclusive breastfeeding.⁵ Children from high-risk families should be fed with hydrolysed formula if breastfeeding is not possible, they should be vaccinated in the same way as lower-risk children, and high-risk families should not have a pet cat.⁵ Health professionals are expected to help their patients adhere to these recommendations. However, allergy prevention may pose a special challenge to health professionals in terms of health literacy (HL)-responsive and evidence-based counselling of parents, for two reasons: First, the evidence on risk factors for childhood allergy changes quickly due to high research activity in the field. For example, while former guidelines have recommended avoidance of allergens (eg, nuts) and delayed introduction of some solid foods, the current guidelines emphasise that parents do not need to take any specific preventive action regarding diet.⁵ Second, scientific evidence on ECAP is socially contested by a variety of non-scientific myths and beliefs around infant feeding and allergy prevention. Studies have highlighted that feeding practices of new mothers are influenced by misinformation about breastfeeding, cultural expectations and stigma or by moralising ideologies.⁶⁻⁷ Likewise, gaps in knowledge about pathogenesis of allergy have allowed outdated ideas to persist, and new myths have emerged from misleading stories in the news and social media, and from product marketers taking advantage of uncertainty.⁸⁻¹¹ Health professionals may thus also need to dissuade parents taking (potentially harmful) actions based on misinformation and fears.

Research has shown that preventive counselling by health professionals can be effective in improving patients’ HL and health behaviour, for example, with regard to smoking cessation and physical activity.¹²⁻¹⁴ Primary care interventions which explicitly promote HL aimed at modifying lifestyle risk factors (eg, nutrition) were also successful in achieving behaviour changes, as demonstrated in a systematic review.¹² There is as yet no data available for the German healthcare context.

To provide effective advice on ECAP, health professionals must:

- Be up-to-date with research evidence on ECAP, consider popular misconceptions and fears and translate this knowledge into clear recommendations for their patients’ families (knowledge translation).
- Know and use counselling techniques that respond to parental HL needs and create a HL-responsive practice setting (HL-responsive care).

The following paragraphs describe these two aspects in detail.

Knowledge uptake and translation
In order to understand how ECAP-specific evidence and recommendations affect daily practice, we need to know how health professionals learn about the current state of the scientific discussion, and how they translate this information into HL-responsive recommendations for parents. Research on the implementation of evidence-based care offers two theoretical perspectives on this process: some studies focus on identifying barriers to and enablers of knowledge dissemination in health professionals’ daily practice (medical and behavioural sciences perspective),¹⁵⁻¹⁷ while others focus on understanding how health professionals make sense of the evidence and how they engage in the social construction of practical expert knowledge (social sciences perspective).¹⁸⁻¹⁹ From the latter perspective, health professionals incorporate a variety of situational determinants, contextual conditions and tacit knowledge, as well as colleagues’ and patients’ experiences, when trying to make sense of the evidence available.¹⁹⁻²³ This process is also conceptualised as a ‘construction of mindlines’²⁰⁻²¹ in social ‘communities of practice’.²¹ In the current study, we embrace both theoretical approaches. A survey of German health professionals’ acceptance of recommendations on infant nutrition²⁵ seems to support the social constructivist view of knowledge translation: the midwives surveyed said that it was more important for them to consider families’ individual situations than to strictly adhere to guidelines.

Apart from this, we do not know how German health professionals gain access to, make sense of and translate into practice the evidence and recommendations available on ECAP.

Health literacy-responsive care
It is broadly recognised that HL should be addressed as part of each clinical encounter, to improve patient outcomes.²² A variety of guidelines on HL-responsive care aim to assist this process. Studies show that different communication techniques help to meet the needs of patients with low HL,²⁷⁻³⁰ for example, avoiding medical jargon, breaking down information to smaller, more manageable components and using the teach-back method, inviting patients to explain key information in their own words.²⁶⁻³¹ Organisations such as the American Medical Association endorse ‘universal HL precautions’, that is, using easy-to-understand terms with all patients instead of focusing only on those with low literacy.²⁶⁻³² As patients with low HL may be difficult to identify,³² Indeed, a systematic review found that many health professionals had a poor understanding of HL, and only a small percentage felt able to identify patients with low HL. Barriers cited to responding to low HL were time constraints and lack of educational resources.³⁴ US studies found that physicians often overlook HL in routine care, overestimating patients’ HL skills.³²⁻³⁴ In response, several studies suggest or employ
curricula for professional training. Beyond these individual-level approaches, care environments, that is, the paediatrician’s or midwife’s office, could be made more HL-responsive, for example, by fostering communication, inviting questions and by creating a ‘shame-free’ care environment, for example by improving the team’s attitudes to patient-centred care. Despite the proven relevance of HL-responsive care and care settings for improving HL and health outcomes in patients, addressing HL is still not a routine part of healthcare delivery.

For the German context, and for ECAP-related HL, it is still unclear how health professionals consider HL in routine care, and what their views are on responding to and supporting parental HL.

In summary, improved HL in ECAP requires the support of primary care health professionals to translate scientific evidence into understandable recommendations and support parents’ allergy-related HL. To make the most of this support, we need to better understand how evidence is taken up and translated by health professionals into practical knowledge, to understand their views and practices with regard to ECAP counselling and parental HL and to explore perceived barriers and enablers in their daily practice.

**Objectives**

This study aims to explore and assess how health professionals (figure 1):

- Inform themselves about ECAP.
- Make sense of the available research evidence.
- Translate this evidence into understandable recommendations for counselling parents.
- Experience HL challenges in ECAP counselling.
- Express their views on HL, HL-responsive care and their experiences with using HL-sensitive counselling techniques and with creating a HL-supportive care setting.
- Describe barriers to and enablers of ECAP knowledge translation and in HL-sensitive and HL-supportive patient counselling.

**METHODS AND ANALYSIS**

**Study design**

This study has a sequential mixed-method design with two consecutive phases (see figure 2). The total duration of the study is 36 months, with the first study phase lasting 24 months from project start in February 2020, and the subsequent phase 2 another 12 months up until March 2023. Qualitative data collection is ongoing until 31 January 2022.

**Patient and public involvement:**

Health professionals did not take part in the study design, but are involved via informal consultations in further developing the interview guides and recruitment strategies during the research process. In the integration and dissemination phase, we are planning to discuss the result with health professionals in order to jointly develop future research projects and interventions.

**Phase 1**

Qualitative exploratory methods will be applied in phase 1, owing to the current lack of research in the area. Semi-structured expert interviews will be conducted at two levels: (a) with primary care health professionals (paediatricians, GPs and midwives) who work in private practices (primary care level), and (b) with (health) professionals who hold a functional position in a professional association or organisation dealing with HL and/or ECAP on a strategic level (professional policy level). We hope by considering both levels to gain greater insight.

**Phase 2**

The qualitative results of phase 1 will provide the basis for developing and pilot testing a standardised questionnaire with paediatricians, GPs and midwives. The questionnaire will systematically assess (a) ECAP and HL practices of the health professionals, and (b) perceived barriers and
enablers of the health professionals regarding HL-responsive counselling.

The qualitative and quantitative results of phase 1 and phase 2 will be integrated at the end of the study in order to generate recommendations for interventions to enhance HL-responsive counselling in ECAP, as well as to develop a study design for further implementation of the assessment tool.

The study is part of the multicentre research unit ‘Health Literacy in Early Childhood Allergy Prevention’ (HELICAP), which uses the example of ECAP to enhance the understanding of HL as a public health goal (https://www.helicap.org/en/home).

Phase 1: exploring health professionals’ views on ECAP knowledge translation and HL-responsive care

Primary care level: participants and sampling/recruitment

Participants

Eligible study participants are German health professionals in primary care who perform routine paediatric/infant healthcare in private practice — that is, prenatal and postnatal follow-up care (midwives) and the national child healthcare screening programme of ‘U-examinations’ (paediatricians/GPs). We intend to include health professionals from two German provinces, one in the northern part of the country (Lower Saxony) and one in the South (Bavaria). Health professionals with less than 2 years of professional experience are excluded, because we assume that is a necessary minimum to be able to report on routine counselling experiences and challenges in day-to-day care.

Sampling and recruitment

Initially, we will identify and recruit study participants via three routes: from pre-existing personal contacts of the research group, by cold calling and by recruiting agents in professional associations. We intend to use a purposive sampling strategy to obtain a great diversity of perspectives (for details cf. table 1). Subsequent snowballing recruitment will occur along with this purposive sampling strategy.44 45

In urban areas, the variety of patients’ background is expected to be greater in terms of socioeconomic status, ethnicity, and, since the prevalence of allergic conditions is higher in urban settings, parental information needs are expected to be different than in rural areas.46 47

Professional policy level: participants and sampling/recruitment

Participants

We will also conduct qualitative expert interviews with health professionals who hold a position in a professional association or political organisation, and who are responsible for (or have a leadership function in) either HL or ECAP in terms of policy, education and training of health professionals, guideline development or specific HL-related projects. These expert interviews will provide a strategic perspective on health professionals’ roles and potential in HL-responsive ECAP, which is valuable to contrast with the views of health professionals responsible for day-to-day healthcare.

Sampling and recruitment

We will identify relevant professional associations and organisations for physicians, midwives and public health in general (eg, State Associations of Physicians/Midwives, National Action Plan on Health Literacy), and subsequently identify and recruit representatives who have a special responsibility for either HL or ECAP. Recruitment will occur via cold calling and pre-existing contacts of the HELICAP research group members.

For all interviews, financial incentives for participation will be offered. Recruitment will be continued until thematic saturation is achieved in analysis, which we expect to be achieved by a sample size of about n=20 in each of the health professional groups in primary care, and by a sample size of about n=10 on the professional policy level, respectively.48

Data collection

Conducting expert interviews is a common qualitative method in public health research, including studies on HL.49 50 The interviews will draw on the experts’ specific experience and knowledge resulting from their daily practice and professional position. When planning the study, the interviews were originally intended to be conducted face to face; however, due to the COVID-19 pandemic and infection prevention measures (social distancing and travel restrictions), we switched to telephone interviews. All qualitative interviews in phase 1 will be led by two experienced members of the research team and will last about 45 min.

Primary care level

The interview guide for the expert interviews on primary care level has been developed based on literature search as well as informal consultations with paediatricians/GPs, and midwives (see online supplemental appendix).

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Table 1 Purposive sampling categories for qualitative data collection

<table>
<thead>
<tr>
<th>Inclusion criteria</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Catchment area</td>
<td>Rural vs urban</td>
</tr>
<tr>
<td>Specialisation</td>
<td>Specialised in allergies vs no specialisation</td>
</tr>
<tr>
<td>Form of practice</td>
<td>Single vs group</td>
</tr>
<tr>
<td>Gender</td>
<td>Male vs female</td>
</tr>
<tr>
<td>Professional experience</td>
<td>15 year +vs less than 15 years</td>
</tr>
</tbody>
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The interview guide starts with an open prompt, asking the interviewee to provide a narrative account of their personal experience with counselling parents on ECAP, and then continues with semi-structured questions covering the following topics:

- Personal practice of searching for, studying and making sense of available evidence on ECAP.
- ECAP knowledge translation into recommendations, knowledge transfer to parents.
- Views on and experiences with counselling parents on ECAP in daily care.
- Awareness of and views on HL-responsive care and counselling (techniques).
- Awareness of and views on creating a HL-responsive practice setting.
- Perceived barriers and enablers regarding knowledge translation and HL-responsive counselling on ECAP.

The first draft of the interview guide will be pretested with paediatricians/GPs and midwives (n=2 each) and revised accordingly. A standardised data sheet will be developed to collect relevant sociodemographic data of the interviewees (ie, age, gender, length of professional experience) as well as characteristics of the practice setting (ie, single vs group practice, specifics about the catchment area, specifications in practice profile).

Professional policy level
The interview guides for the interviews on professional policy level will be developed based on further literature searches, field knowledge of the research team and insights from the interviews on primary care level. Depending on the interviewees’ position and expertise, the interview guide will focus either on HL or on ECAP, and/or on training of physicians or midwives, etc. The guides will encompass semi-structured questions covering the following topics:

- Role and responsibility of health professionals (midwife or physician, depending on sample) in ECAP or HL, respectively.
- Challenges of knowledge translation in ECAP/challenges of HL-responsive care.
- View on feasible HL practices/ECAP counselling in daily care.
- Needs for action (policies, education/training, structures, resources).

Data analysis
All interviews will be audio-recorded and transcribed verbatim with participants’ permission. Data will be analysed using computer-assisted qualitative data analysis software (ATLAS.ti, V.7), based on structuring qualitative content analysis. This approach allows for discovering newly emerging themes via open coding (‘inductive category formation’). Two experienced researchers will repeatedly read, code and analyse the transcripts, structuring the data into overarching themes as predefined in the interview guide, while at the same time identifying emergent themes in the data. Differences in coding between the researchers will be discussed until consensus is reached. Themes will then be grouped into higher-level categories.

Analysis of the expert interviews on the professional policy level will be performed as for the interviews with health professionals in primary care.

Finally, we will contrast the views of professionals in a leadership role with the views of those who report from day-to-day interaction with patients. Thereby, we intend to compare and contrast the positions in order to elicit potential consensus, (mis)understandings and mutual expectations.

All results of the qualitative study phase will be prepared for publication according to the consolidated criteria for reporting qualitative research-standards for reporting on qualitative research results.

Phase 2: developing and pilot testing a standardised questionnaire and study design
The subsequent study phase depends on the findings from phase 1 (sequential-dependent study design). Following a ‘building approach’ for integrating qualitative and quantitative data at methods level in the mixed-method design, the results of the qualitative study phase will inform the proceedings and content of the second study phase, that is, the development and testing of a standardised questionnaire. The pilot test aims to evaluate the usability of the questionnaire and the process of data collection. We aim to apply the results of phase 2 for a representative survey in a follow-up project.

Participants, eligibility criteria, recruitment and sampling
The participants will be chosen by random sampling from all health professionals registered in Bavaria (Southern German province). As of 2019, there are n=1140 paediatricians, n=2850 midwives and n=9343 GPs registered in Bavaria. We will sample only primary care professionals who work in private practice, and only GPs who perform infant screening examinations (cf. eligibility of participants of phase 1).

Assessment tool development and data collection in pilot test
When developing the assessment tool, items will be generated based on results of phase 1, in order to capture barriers and enablers. The questionnaire will be checked for face validity and tested for comprehensibility, comprehensiveness and relevance in a pretest with three representatives of each health professional group (paediatricians, midwives and GPs). The think-aloud method will be applied when the interviewed person fills in the questionnaire.

The (mail) questionnaire will be distributed among n=100 participants of each of the three health professional groups (paediatricians, GPs and midwives). Special efforts will be needed to ensure high response rates, since survey return rates for health professionals, above all physicians, tend to be very low. We will therefore draw on Dillman’s total design method, which recommends...
to create a relationship of reciprocity with respondents, to lower their investment and maximise their rewards (eg, by offering incentives) and to include repeated non-responder follow-up mailings.\textsuperscript{62-63} The researchers will document each step in a semi-standardised documentation sheet, including time, effort, effectiveness and feasibility of procedures. The data collected will be converted into a data set using the statistical software SPSS.

**Data analysis and revision of study design and questionnaire**

The collected data of the pilot surveys will be analysed, for example, regarding completeness and frequency of missing values using statistical software (SPSS, V.25). Free-text responses will be analysed for incomprehensible or misleading items and variance of the responses. The implementation of the questionnaire will be evaluated regarding response rates (overall and within the three groups of health professionals), the duration from first contact to response and the number of contacts needed. Based on the statistical data analysis and the documentation sheet, strengths and weaknesses of the study design and questionnaire will be assessed. A revised study protocol and questionnaire will be generated. These tools are intended to guide a representative questionnaire survey, which is planned for a second funding phase after the end of this 3-year research study, and which is meant to test the generalisability of the findings of this research study.

**Integration of results**

The research team will integrate the findings of the qualitative and quantitative study phases at two points. First, the design and contents of the quantitative study, for example, questionnaire items, will be developed according to the results of the qualitative study phase (‘building approach’, see methods section). Second, at the end of the study, we will integrate the qualitative and quantitative findings at the interpretation and reporting level.\textsuperscript{54} While parts of the qualitative and quantitative results will be published and presented separately during the course of the study, theme-based parts of the qualitative and quantitative results will be reported jointly at the end, using three types of outputs. First, a deeper understanding of themes and challenges, barriers and enablers. Second, practice recommendations for interventions aimed at enhancing health professionals’ skills. Third, based on the qualitative and quantitative insights, we will finalise the pilot-tested assessment tool and study design in preparation of a larger-scale representative survey.

**ETHICS AND DISSEMINATION**

Ethical considerations and informed consent:

The study has received ethical approval from the Ethics Committee of the University of Regensburg (18-1205-101). This study complies with the Declaration of Helsinki.\textsuperscript{62,63}

Participation in the study will only be possible after providing informed consent to the audio recording and scientific use of the interviews. All of the study information and informed consent documents which will be handed out to study participants have been approved by the ethics committee of the University of Regensburg.

**Confidentiality and data protection**

The transcripts will be deidentified with regard to names and cities: names will be replaced by an ID number, and personal identifiers will be removed early on during transcription and analysis. All persons involved in collecting, transcribing and analysing the data will be trained in procedures to protect the participants’ privacy and confidentiality and are bound by the Data Protection Act (DSGVO). When disseminating findings from the research study, direct quotes from respondents may be used as an example and will remain anonymous.

All databases and the software used are stored on secure servers of the University of Regensburg. Audiotapes, transcriptions and notes will be kept in a secure, locked location to which only authorised persons will have access. A numbered hard copy of the transcriptions will be retained, to provide for easy follow-up in the case of questions, and an additional electronic copy will be stored so that it can be used for data confirmation and/or audits. In accordance with the rules of good scientific practice, the qualitative research data (audio data and transcripts) will be archived in the Department of Epidemiology and Preventive Medicine of the University of Regensburg for at least 10 years.

**Output, dissemination and outlook**

The results of this study will be published in international peer-reviewed open access journals and via oral and poster presentations at national and international scientific conferences. We will also disseminate results via presentations and workshops with German health professionals, decision-makers and potential funders of interventions (eg, health insurance providers).

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**Contributors**

JC conceptualised the study. JvS, JL and MLD contributed to further development of the study design. JC drafted the first version of the manuscript and finalised it. JvS, E-MG, JL and MLD revised the manuscript. All authors approved the final version.

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**Competing interests**

None declared.

**Patient consent for publication**

Not required.

**Provenance and peer review**

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Supplemental material

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**Topics/questions:**

**Introduction**

1. How is early childhood allergy prevention relevant in your routine counselling?
   a. When and with whom do you address this topic?
   b. What is the most important message you want to convey to parents concerning allergy prevention? Do you make a difference between low- and high-risk groups?
   c. Which topics do also matter in your counselling on early childhood allergy prevention (e.g. nutrition, living environment, early exposure to allergens)?

**Information and evidence**

1. What are your main sources to keep yourself informed about health topics (especially allergy prevention)?
2. Are you satisfied with the available information?
   a. Do you feel well informed? Why? Why not?
   b. Is there anything that would help you to keep up to date?
3. How do you handle inconsistent and changing information? (e.g. avoidance of allergens vs. early confrontation with allergens)
   a. How did/do you feel about it when a key message of your consultation changes due to new research findings?

**Knowledge translation and implementation**

1. We have just talked about how you keep yourself up to date on health-related recommendations: What are you doing with this knowledge (in your head) to make it applicable/use it for your practice?
2. How do you incorporate this knowledge into your daily work?
   a. Do you pass on specific scientific information to your patients (on ECAP)?
      i. If yes, could you give an example how you pass on this information to your patients?
      ii. What helps you to pass on your knowledge to parents?
3. When you think of scientific evidence and recommendations, etc.: Do you experience any barriers that make it difficult for you to share the information with parents in an understandable way? What exactly makes that difficult?
   a. Can you describe a particularly difficult counselling situation?
4. Is there anything that could be improved to help you in counselling parents on health-related issues?
Promotion of health literacy

A. Counselling of parents and health literacy

1. How do you deal with different patients (level of knowledge, education, migration background) in counselling and transfer of knowledge? Do you differentiate? Could you give an example?

2. **How do you assess** the level of knowledge and information demand of parents?
   a. How do you notice that you have to explain a lot/in a way that is easy to understand?
   b. Do you consciously use strategies to assess what kind of information and support needs parents have? If yes, which ones do you use?

3. Do you think parents are well informed regarding allergy prevention?
   a. How do you recognize that?

4. What previous knowledge or lay conceptions do parents bring up in counselling?

5. What opportunities do you see to support parents with regard to accessing and appraising information?
   a. Could you give some examples from your everyday practice?

6. Do you provide your patients with information sources so they can inform themselves about health topics (e.g. allergy prevention)?
   a. Which sources do you consider (not) helpful?

7. Do you use certain strategies during counselling to make sure the parents understand everything? (e.g. plain language, drawing pictures)
   a. If yes, what are your experiences with these strategies?
   b. Do you encourage parents to ask questions? How?

8. How do you ensure that parents apply your health behavior recommendations?

B. Attitudes towards and experiences with health literacy/health literacy-sensitive care

9. How well do you feel educated/trained to counsel your patients according to their needs?

10. Do you personally see a need of further information/training on how to convey health knowledge to parents in a more understandable manner and/or on how you can take greater account of the parent’s health literacy or even strengthen it?

11. How should academic results/recommendations be prepared and made available so that you can use them efficiently in consultations?

**HL concept awareness**

1. Have you heard of the term “health literacy” in the context of your work or in advanced training courses?