BMJ Open Costs associated with functional gastrointestinal disorders and related signs and symptoms in infants: a systematic review protocol

Julie Glanville,¹ Thomas Ludwig,² Carlos Lifschitz,³ James Mahon,¹ Mohamad Miqdady,⁴ Miguel Saps,⁵ Seng Hock Quak,⁶ Irene Lenoir-Wijnkoop,⁷ Mary Edwards,¹ Hannah Wood,¹ Hania Szajewska⁸

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JG and TL contributed equally.

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For numbered affiliations see end of article.

Correspondence to
Professor Hania Szajewska;
hania@ipgate.pl

ABSTRACT

Introduction: Functional gastrointestinal disorders (FGIDs) and FGID-related signs and symptoms have a fundamental impact on the psychosocial, physical and mental well-being of infants and their parents alike. Recent reviews and studies have indicated that FGIDs and related signs and symptoms may also have a substantial impact on the budgets of third-party payers and/or parents. The objective of this systematic review is to investigate these costs.

Methods and analysis: The population of interest is healthy term infants (under 12 months of age) with colic, regurgitation and/or functional constipation. Outcomes of interest will include the frequency and volume of reported treatments, the cost to third-party payers and/or parents for prescribed or over the counter treatments, visits to health professionals and changes in infant formula purchases, and the loss of income through time taken off work and out of pocket costs. Relevant studies will be identified by searching databases from 2005 onwards (including MEDLINE, EMBASE, PsycINFO, NEXIS, DARE, Health Technology Assessment database, National Health Service Economic Evaluation Database and others), conferences from the previous 3 years and scanning reference lists of eligible studies. Study selection, data extraction and quality assessment will be conducted by two independent reviewers and disagreements resolved in discussion with a third reviewer. Quality assessment will involve study design-specific checklists. Relevant studies will be summarised narratively and presented in tables. An overview of treatments and costs will be provided, with any geographical or other differences highlighted. An assessment of how the totals for cost differ across countries and elements that contribute to the differences will be generated.

Ethics and dissemination: This is a systematic review of published studies that will be submitted for publication to a peer-reviewed journal. Ethical committee approval is not required.

Trial registration number: CRD42016033119.

INTRODUCTION

Functional gastrointestinal disorders (FGIDs) and related signs and symptoms can have

Strengths and limitations of this study

- This is a systematic review with a registered protocol, following rigorous methods, including an extensive search, and data extraction and study quality assessment by two independent reviewers.
- The review seeks to identify direct and indirect costs from relevant populations: both studies with healthy term infants under 12 months of age who meet the Rome III criteria for functional constipation, infantile colic and regurgitation and studies of infants of the same age with gastrointestinal symptoms provided that these are linked to one of the aforementioned functional gastrointestinal disorders.
- The review is focused on more recent studies to ensure that currency and most recent practice are reflected in terms of care of functional gastrointestinal disorders and related signs and symptoms.
- The searches are limited to studies published since 2005 in English.

fundamental short-term and long-term impacts on the physical and mental well-being of children and parents.^{1–6} FGIDs include a variable combination of signs and symptoms which cannot be explained by obvious structural or biochemical abnormalities.^{7–9}

The literature suggests that more than 50% of all infants display at least one FGID or related sign and symptom during the first year after birth.^{10–12} A recent meta-review reported that the worldwide prevalence of the three most common FGIDs in infants, infantile regurgitation, colic and functional constipation, is ~30%, 20% and 15%, respectively.⁶

Experts broadly agree that the preferred treatment for infantile colic and regurgitation should be parental reassurance and nutritional advice.^{13–17} Nutritional management of regurgitation involves correcting the frequency and

volume of feeds and, in some cases of formula-fed infants, the use of antiregurgitation formulas.^{9 18–21} Thickened antiregurgitation formulas have been shown to reduce the volume and frequency of regurgitation and crying, to improve sleep and to support weight gain.^{18–21}

There is broad consensus in reviews that there is no indication for pharmacological treatment, such as proton pump inhibitors (PPIs) or prokinetic drugs, in infants with infantile colic or regurgitation, even if these are showing signs of distress. These reviews suggest that there is either no evidence of the effectiveness of pharmacological treatment or evidence of no effect, and there can be significant risks and negative side effects.^{14 18 20 22–25}

A recent survey of Italian general paediatricians on the implementation of the 2009 North American Society for Pediatric Gastroenterology, Hepatology, and Nutrition (NASPGHAN)-European Society for Pediatric Gastroenterology, Hepatology, and Nutrition (ESPGHAN) guidelines in children with GER symptoms²⁶ found that only 2% of participating paediatricians completely adhered to these guidelines. The study also found that 56% of paediatricians prescribed PPIs for infants with unexplained crying and/or distressed behaviour and 38% prescribed PPIs in infants with uncomplicated recurrent regurgitation and vomiting. Of the participating paediatricians, 79% were reported to overprescribe PPIs.

The Avon Longitudinal Study of Parents and Children (ALSPAC) found that 16% of infants with colic and flatulence were given activated dimethicone, and 13% were given gripe water.²⁷ Systematic reviews (SRs) have found both remedies to be not effective in the treatment of either of these conditions.^{17 28} The use of other medicinal and herbal products for the treatment of infantile colic is also common. Most of these products are either not licensed for use in this age group or their safety and tolerance have not been tested in clinical studies.^{14 27–33}

Although some of these FGIDs or related signs and symptoms may not require treatment, besides reassuring and educating parents, parents frequently seek medical attention or treatment.^{6 11 34} Infantile regurgitation, colic and constipation in infants are among the most common reasons for parents to consult a healthcare professional in the first months after birth.^{10 11} As such, the annual costs for sleeping and crying disorders in the UK are estimated to be in the range of £65 million.³⁵ Infants have the highest rate of emergency department visits for constipation in the USA³⁶ and the average costs of care per patient are US\$2306.³⁶

In summary, reviews and studies have indicated a high prevalence of FGIDs and related signs and symptoms in infants, and that these can cause relevant direct and indirect costs that may have a substantial impact on third-party payers and/or parents.

OBJECTIVES

The primary objective is to conduct an SR of the direct and indirect costs to third-party payers and/or parents of infants with FGIDs and related signs and symptoms.

The SR will seek data to answer the following questions:

- What is the international, national and regional cost of illness (COI) for FGIDs and related signs and symptoms reported in published studies?
- What are the costs of treating FGIDs and related signs and symptoms from the perspective of public payers, health insurers and parents of infants with FGIDs and related signs and symptoms (private purchase of remedies and interventions)?
- What are the indirect costs of FGIDs and related signs and symptoms in terms of parents' days lost from work/reduced productivity and from parental out of pocket expenses?

Using the evidence from the SR, at least one *de novo* exemplar COI calculation will be undertaken to estimate the total costs associated with the care of infants with FGIDs and related signs and symptoms in a developed economy.

The review protocol has been registered with PROSPERO (CRD42016033119).

METHODS AND ANALYSIS

The SR eligibility criteria are described below and are summarised in [table 1](#).

Participants

The population of interest is healthy term infants (under 12 months of age) with colic, regurgitation and/or functional constipation.

Studies that include infants who meet the Rome III criteria for the clinical diagnosis of infantile colic, regurgitation and/or constipation will be eligible for inclusion. Ideally, the cost of each of these conditions should be explored separately, but it is anticipated that this may not be possible due to the overlap in these conditions. FGID-related signs and symptoms that do not fulfil the Rome criteria will also generate direct and indirect costs ([figure 1](#)). Therefore, studies which include infants with symptoms, including hard stools or straining, vomiting, problematic feeding (eg, spitting) or inconsolable or persistent crying, will be eligible for inclusion, provided that these symptoms are linked to at least a suspicion of FGID in all cases ([figure 2](#)). Studies with these symptoms where no statement on the suspected underlying cause is made or where the symptoms could be the result of a non-FGID cause will not be eligible for inclusion ([figure 1](#)).

Studies involving exclusively preterm infants or where the majority of the infants are preterm will not be eligible.

In all cases, the definition used by authors for FGIDs and related signs and symptoms will be recorded.

Studies of participants from any country are eligible.

Interventions

Studies reporting on the costs of any intervention or no intervention are eligible for inclusion.

Table 1 Summary of inclusion/exclusion criteria

Criterion	Inclusion criteria	Exclusion criteria
Participants	Healthy term infants under 12 months who meet the Rome III criteria for <ul style="list-style-type: none"> ► Constipation ► Infantile colic ► Regurgitation Studies which include infants with symptoms, including hard stools or straining, vomiting, problematic feeding (eg, spitting) or inconsolable or persistent crying, will be eligible for inclusion, provided that these symptoms are linked to at least a suspicion of FGID in all cases	Infants without FGID. Infants with vomiting, problematic feeding, inconsolable or persistent crying and where no statement on the believed underlying gastrointestinal cause is made or where the symptoms could be the result of a non-FGID cause Preterm infants only or studies where preterm infants form >50% of the participants
Interventions	Any intervention (or no intervention for burden on parents or carers)	
Comparators	No comparator is necessary	
Outcomes	Reported treatments of FGIDs and related signs and symptoms (regardless of effectiveness) and the frequency and volume of use Costs to patients and third-party payers of <ul style="list-style-type: none"> ► Prescribed treatments ► Over the counter or home remedies ► Visits to orthodox and complementary health professionals and other providers of care Loss of income for parents/carers from infant FGIDs and related signs and symptoms through: <ul style="list-style-type: none"> ► Inability to return to work ► Time taken off work ► Out of pocket costs 	
Study types	<ul style="list-style-type: none"> ► RCTs ► Non-randomised controlled trials ► Cost of illness studies ► Economic evaluations ► Observational studies 	Non-systematic reviews Conference abstracts Letters Comments Editorial News stories Case reports Studies published before 2005 Studies in languages other than English
Limits	Studies published from 2005 onwards Studies in English Full publications	

Comparators

A study with any or no comparator will be eligible for inclusion in this SR.

Outcomes

The outcomes of interest are:

- Reported treatments of FGIDs and related signs and symptoms (regardless of their effectiveness), or the specific symptom combinations described in the Population description above, and the frequency and volume of use.
- Costs to third-party payers and/or parents of infants with FGIDs and related signs and symptoms of:
 - prescribed treatments;
 - over the counter or home remedies;
 - visits to orthodox and complementary health professionals and other providers of care;
 - change in infant formula purchases.
- Loss of income for parents/carers from infants with FGIDs and related signs and symptoms, or the specific

symptom combinations described in the Population section above, through:

- inability to return to work;
- time taken off work;
- out of pocket costs.

Study types

Studies of the following designs will be eligible for inclusion in the review:

- randomised controlled trials (RCTs);
- non-RCTs;
- COI studies;
- economic evaluations;
- observational studies.

Previously published SRs will not be eligible for inclusion, but their reference lists will be checked to identify any additional relevant studies.

Studies published as abstracts or conference presentations will not be included in the review but will be listed for information and used as clues to full publications.

Figure 1 Defining the population of interest. Infants who present in daily paediatric practice or receive treatment for functional gastrointestinal symptoms may not always fulfil the stringent criteria for a functional gastrointestinal disorder (FGID) as defined by the Rome criteria (dark centre). However, these FGID-related symptoms do still cause direct and indirect costs and need to be considered (light circles).

Any study of a design not listed above will not be eligible for inclusion in the SR. This includes, but is not limited to, case studies and non-SRs. Opinion pieces, letters and editorials will not be eligible for inclusion.

Limits

The searches are limited to English language studies published since 2005 to focus on the most current costs and most recent practice in terms of care of FGIDs and related signs and symptoms.

Identifying relevant studies

The search strategy to identify studies for this review is shown in [figure 2](#). The search excludes publication types that are unlikely to contain research results. Records of publications published prior to 2005 are excluded and only studies in English will be retrieved.

The searches will be conducted in the following databases:

- ▶ MEDLINE and MEDLINE in process (via OvidSP);
- ▶ PubMed;
- ▶ EMBASE (via OvidSP);
- ▶ PsycINFO (via OvidSP);

- ▶ NEXIS (media database useful for commercial/market reports on over the counter sales of interventions such as gripe water);
- ▶ Database of Abstracts of Reviews of Effects (DARE) in the Cochrane Library;
- ▶ Health Technology Assessment Database (HTA Database) in the Cochrane Library;
- ▶ NHS Economic Evaluation Database (NHS EED) in the Cochrane Library;
- ▶ CEA Registry (via <https://research.tufts-nemc.org/cear4/>);
- ▶ Evidence Search (<https://www.evidence.nhs.uk/>);
- ▶ OAISTER (<http://www.oclc.org/oaister.en.html>);
- ▶ RePEc (<http://repec.org/>).

The search strategy ([figure 2](#)) will be adapted suitably to perform efficiently in these databases.

The following conferences will be searched via Embase or via conference websites for the past 3 years, to identify recently completed research and to provide contact details, so that the authors can be contacted for full text:

- ▶ ISPOR;
- ▶ European Society for Paediatric Gastroenterology, Hepatology and Nutrition;


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1 "cost of illness"/ (20176)
2 (costing adj3 (illness$ or disease$ or sickness$)).ti,ab,kf. (36)
3 (burden adj3 (illness$ or disease$ or sickness$)).ti,ab,kf. (16032)
4 (burden adj3 (family or families or human$1 or mother$ or father$ or parent$ or caregiver$ or care-giver$)).ti,ab,kf. (1827)
5 ((economic or human$) adj3 consequence$1).ti,ab,kf. (4195)
6 "costs and cost analysis"/ or cost-benefit analysis/ (106732)
7 exp health care costs/ (51343)
8 (cost or costs or economic evaluation or pharmacoeconomic).ti,ab,kf. (324226)
9 (resource$1 adj4 use$1).ti,ab,kf. (17903)
10 (resource$1 adj4 usage).ti,ab,kf. (348)
11 (resource$1 adj4 utili$).ti,ab,kf. (8949)
12 (visit or visits or hospitalization$1 or hospitalisation$1 or admission$1 or admitted or emergency room or rescue).ti,ab,kf. (456371)
13 quality-adjusted life years/ or "quality of life"/ (140039)
14 (quality adjusted life or qol).ti,ab,kf. (27230)
15 (qaly$ or qald$ or qale$ or qtime$).ti,ab,kf. (5643)
16 (sf36 or sf 36 or sf thirty six or sf thirty six).ti,ab,kf. (14109)
17 (sf6 or sf 6 or short form or shortform or sf six or sfsix).ti,ab,kf. (19557)
18 (sf12 or sf 12 or sf twelve or sftwelve).ti,ab,kf. (2473)
19 (sf16 or sf 16 or sf sixteen or sfsixteen).ti,ab,kf. (18)
20 (sf20 or sf 20 or sf twenty or sftwenty).ti,ab,kf. (301)
21 (euroqol or eq5d or eq 5d).ti,ab,kf. (4554)
22 (hql or hqol or hrqol or hrql or hr ql).ti,ab,kf. (10731)
23 (hye or hyes).ti,ab,kf. (60)
24 health$1 year$1 equivalent$1).ti,ab,kf. (38)
25 (hui or hui1 or hui2 or hui3).ti,ab,kf. (943)
26 disutili$.ti,ab,kf. (242)
27 (quality adj3 (wellbeing or well being)).ti,ab,kf. (1438)
28 qwb.ti,ab,kf. (179)
29 (willingness adj3 pay).ti,ab,kf. (2566)
30 standard gamble$.ti,ab,kf. (698)
31 (time trade off$1 or time tradeoff$1 or tto or timetradeoff).ti,ab,kf. (1261)
32 ((valu$ or measur$) adj3 (health or outcome$1 or effect$1 or change$1 or state$1)).ti,ab,kf. (283373)
33 (preference$ adj3 (patient$1 or public or valu$ or measur$)).ti,ab,kf. (12015)
34 ((quality adj3 life) or qol).ti,ab,kf. (160969)
35 (index adj3 wellbeing).ti,ab,kf. (81)
36 (multiattribute$ health or multi attribute$ health).ti,ab,kf. (53)
37 (multiattribute$ theor$ or multi attribute$ theor$ or multiattribute$ analys$ or multi attribute$ analys$).ti,ab,kf. (10)
38 (multiattribute$ utilit$ or multi attribute$ utilit$).ti,ab,kf. (198)
39 (utilit$ adj3 (valu$ or measur$ or health or life or estimat$ or elicit$ or disease)).ti,ab,kf. (6614)
40 (euro qual or euroqual).ti,ab,kf. (15)
41 (visual analog$ or vas).ti,ab,kf. (46648)
42 (prom or proms or patient reported outcome$1 or pro or pros).ti,ab,kf. (127679)
43 functional assessment.ti,ab,kf. (6235)
44 (symptom$1 adj4 (score$1 or scale$ or instrument$1 or measur$)).ti,ab,kf. (39358)
45 exp patient satisfaction/ (68374)
46 (satisfaction or dissatisf$ or unsatisf$).ti,ab,kf. (104301)
47 (anxiety or depression or anxious or depressed).ti,ab,kf. (348460)
48 exp emotions/ (182971)
49 exp fatigue/ or absenteeism/ or presenteeism/ (22534)
50 stress,psychological/ (95491)
51 (Gastrointestinal rating scale or GSRS or (Gastrointestinal Quality adj3 Index) or GIQLI or (Severity adj2 Dyspepsia Assessment) or SODA).ti,ab,kf. (3202)
52 ((parent$ or family or families or mother$ or father$ or caregiver$ or care-giver$) adj5 (concern$1 or perception$1 or view$1 or worry or worrie$1)).ti,ab,kf. or parents/px
53 or/1-52 (1988310)
54 (colic/ or exp Diarrhea/ or Colonic Diseases, Functional/ or exp abdominal pain/) and (exp infant/ or child, preschool/) (19125)
55 diarrhea, infantile/ (6857)
56 gastrointestinal diseases/ and pain/ and (exp infant/ or child, preschool/) (52)
57 (constipation/ or vomiting/) and (exp infant/ or child, preschool/) (5474)
58 ((infantile or infant$1 or baby or babies or neonat$ or newborn$1 or new born or toddler$1 or child or children or pediatric or paediatric) adj5 (colic or constipation or constipated or regurgitat$ or spitting or spit)).ti,ab,kf. (2389)
59 ((infantile or infant$1 or baby or babies or neonat$ or newborn$1 or new born or toddler$1 or child or children or pediatric or paediatric) adj5 (colicky or defecat$ or stool$1 or bowel movement$1)).ti,ab,kf. (2830)
60 ((fgid or fgids) and (infantile or infant$1 or neonat$ or baby or babies or newborn$1 or new born or toddler$1 or child or children or pediatric or paediatric)).ti,ab,kf. (93)
61 (crying adj5 (infantile or infant$1 or neonat$ or baby or babies or newborn$1 or new born or toddler$1 or child or children or pediatric or paediatric)).ti,ab,kf. (1012)
62 (gastrointestinal adj5 (infantile or infant$1 or neonat$ or baby or babies or newborn$1 or new born or toddler$1 or child or children or pediatric or paediatric)).ti,ab,kf. (4024)
63 ((dyschezia or colonic inertia or diarrhea or diarrhoea or cramp$ or reflux or functional abdominal pain or bowel symptom$1 or irritable bowel or IBS) adj5 (infantile or infant$1 or neonat$ or baby or babies or newborn$1 or new born or toddler$1 or child or children or pediatric or paediatric)).ti,ab,kf. (14827)
64 or/54-63 (38774)
65 53 and 64 (5973)
66 limit 65 to (english language and yr="2005-Current") (2623)

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Figure 2 Search strategy for Ovid MEDLINE® in-process and other non-indexed citations and Ovid MEDLINE® <1946 to present>.

- ▶ North American Society for Pediatric Gastroenterology, Hepatology and Nutrition;
- ▶ American Academy of Pediatrics National Conference;
- ▶ World Congress of Pediatric Gastroenterology, Hepatology and Nutrition.

The reference lists of any included studies and SRs will be checked to identify any studies that may have been missed by the database searches.

Details of all search strategies and search results will be provided in the SR report. The search and record

selection process will be presented in a PRISMA flow diagram (<http://www.prisma-statement.org/statement.htm>).

Incidence/prevalence data would be required to perform a *de novo* calculation: these values will be sought from the published literature using a pragmatic focused epidemiology search in MEDLINE (figure 3). This strategy will be combined with specific country terms depending on the exemplar country.

Figure 3 Pragmatic focused epidemiology search in MEDLINE.

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1 (colic/ep or exp Diarrhea/ep or Colonic Diseases, Functional/ep or exp abdominal pain/ep) and (exp infant/
  or child, preschool/)
2. (colic/ or exp Diarrhea/ or Colonic Diseases, Functional/ or exp abdominal pain/) and (exp infant/ or child,
  preschool/) and (incidence/ or prevalence/)
3 diarrhea, infantile/ep or (diarrhea, infantile/ and (incidence/ or prevalence/))
4 gastrointestinal diseases/ep and pain/ and (exp infant/ or child, preschool/)
5 gastrointestinal diseases/ and pain/ and (exp infant/ or child, preschool/) and (incidence/ or prevalence/)
6 (constipation/ep or vomiting/ep) and (exp infant/ or child, preschool/)
7 (constipation/ or vomiting/) and (exp infant/ or child, preschool/) and (incidence/ or prevalence/)
8 ((infantile or infant$1 or baby or babies or neonat$ or newborn$1 or new born or toddler$1 or child or children
  or pediatric or paediatric) adj5 (colic or constipation or constipated or regurgitat$ or spitting or spit)).ti,ab,kf.
  AND (incidence or prevalence or epidemiol$).ti.
9 ((infantile or infant$1 or baby or babies or neonat$ or newborn$1 or new born or toddler$1 or child or children
  or pediatric or paediatric) adj5 (colicky or defecat$ or stool$1 or bowel movement$1)).ti,ab,kf. AND
  (incidence or prevalence or epidemiol$).ti.
10 ((fgid or fgids) and (infantile or infant$1 or neonat$ or baby or babies or newborn$1 or new born or toddler$1
  or child or children or pediatric or paediatric)).ti,ab,kf. AND (incidence or prevalence or epidemiol$).ti.
11 (crying adj5 (infantile or infant$1 or neonat$ or baby or babies or newborn$1 or new born or toddler$1 or child
  or children or pediatric or paediatric)).ti,ab,kf. AND (incidence or prevalence or epidemiol$).ti.
12 (gastrointestinal adj5 (infantile or infant$1 or neonat$ or baby or babies or newborn$1 or new born or toddler$1
  or child or children or pediatric or paediatric)).ti,ab,kf. AND (incidence or prevalence or epidemiol$).ti.
13 ((Dyschezia or "Colonic Inertia" or diarrhea or diarrhoea or cramp$ or reflux or "functional abdominal pain" or
  "bowel symptom$1" or "irritable bowel" or IBS) adj5 (infantile or infant$1 or neonat$ or baby or babies or
  newborn$1 or new born or toddler$1 or child or children or pediatric or paediatric)).ti,ab,kf. AND (incidence
  or prevalence or epidemiol$).ti.
14 or/1-13
15 {country search terms}
16 14 AND 15

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Study selection

The results of the searches will be loaded into an EndNote library and deduplicated. Following the removal of obviously irrelevant records by a single experienced information specialist, studies will be added to Covidence software. (The Alfred Hospital, Monash University, National ICT Australia, *et al.* Covidence. Secondary Covidence [reviewing software]. <https://www.covidence.org>.)

Study selection will be undertaken by two reviewers independently, based first on information in the titles and abstracts (when available). Disagreements will be resolved by discussion or involvement of a third reviewer. Studies that are not excluded based on title and abstract screening will be retrieved as full-text documents. Records excluded based on an assessment of the full text will be recorded in an excluded studies table with a reason for exclusion.

Data extraction

Double independent reviewer data extraction will be undertaken into an Excel spreadsheet. Any disagreement between the two reviewers will be resolved by discussion and, if necessary, by a third reviewer.

The following data will be extracted from all eligible included publications:

- ▶ bibliographic data;
- ▶ publication type;
- ▶ country;
- ▶ patient population (number, age, gender, ethnicity);
- ▶ definition of FGID and, if applicable, the definition of the symptoms related to FGID.

Topic specific information will also be extracted (table 2).

Quality assessment

Double independent reviewer quality assessment will be undertaken. Any disagreement between the two reviewers will be resolved by discussion and, if necessary, a third reviewer will be involved to adjudicate. The quality of COI studies will be assessed using the following questions:

- ▶ Was the type of COI method clearly described?
- ▶ Was the quality of the data used described?
- ▶ Were the data sources and dates reported clearly?
- ▶ Were data gaps described?
- ▶ Were data extrapolations reasonable?
- ▶ Were reasonable methods employed to avoid double counting?
- ▶ Were the calculations of cost clearly described?
- ▶ Were the methods used to handle uncertainty appropriate?
- ▶ Have the researchers offered assessments of the limitations of the study approach?

The quality of other studies will be assessed using instruments listed in table 3.

Data synthesis

Studies identified for the SR and the extracted data will be summarised in tables.

Results will be presented by geographical region and/or country and by specific FGID conditions. A distinction will be made in the tables to highlight interventions that are reported in the literature to be common or accepted practice and those that have been trialled.

An overview of treatments in use and costs incurred will be provided, with any geographical or other differences highlighted. An assessment of how the totals for

Table 2 Data extraction elements

Systematic review question	Data extraction
What are the current treatment options (medicinal and otherwise) in use?	<ul style="list-style-type: none"> ▶ Study aim and type ▶ Treatment option (description/dosage) ▶ Prescription only/parent purchase/fees ▶ Timescales/length of treatment ▶ Number of treatments ▶ Management of adverse effects ▶ Management of comorbidities
What are the direct and indirect costs for parents?	<ul style="list-style-type: none"> ▶ Study aim and type ▶ Burden components (eg, over the counter treatments, changes in infant formula purchases and the loss of income through time taken off work/inability to return to work due to caring for the infant or attend clinical appointments) ▶ Dates of data collection ▶ Currency ▶ The value of any costs identified
What are the current estimates of the COI of FGIDs and related signs and symptoms?	<ul style="list-style-type: none"> ▶ Study aim and type ▶ Definition of COI ▶ Burden components (eg, prescribed treatments, visits to health professionals) ▶ Dates of data collection ▶ Currency ▶ Methods of data identification/collection ▶ Results: COI ▶ Limitations

cost differ across countries and the elements that contribute to the differences—such as the difference in costs for treatments or the different ways healthcare is provided between countries—will be undertaken as far as the detail in the identified literature allows. If applicable, gaps and limitations of this approach will be highlighted and discussed in detail.

If data permit, one exemplar of what the COI would be in total across all cost areas considered for a developed country will be produced (*de novo* calculation) using data from the review where possible. Following data synthesis, a recommendation will be made regarding which country should be chosen, based on the completeness of evidence from the literature for making a COI calculation, and the accessibility of unpublished data to fill any gaps.

Table 3 Quality assessment instruments

Study design	Tool
RCTs	Cochrane Risk of Bias tool ³⁷
Economic Evaluations	Drummond and Jefferson checklist ³⁸
Case-control studies	Case-control Checklist (CRD report 4) ³⁹ Newcastle Ottawa Scale ⁴⁰
Case series	Case Series Checklist (CRD report 4) ³⁹
Cohort studies	Cohort Study Checklist (CRD report 4) ³⁹
Cross-sectional studies	Adapted Newcastle-Ottawa for Cross-Sectional studies, adapted by Herzog <i>et al</i> (2013) ⁴¹
Decision models	Philips Checklist ⁴²

All costs identified will be adjusted for inflation and converted to 2015 US\$.

DISSEMINATION

The high incidence of FGIDs and related signs and symptoms in infants has been reported and reviewed.^{6 11} There is a gap in knowledge on the international, regional and national economic impact of infant FGIDs and related signs and symptoms which this SR seeks to close.

CONCLUSION

This SR will provide specific insight into the direct and indirect costs associated with FGIDs and related signs and symptoms to parents and third-party payers.

Author affiliations

¹York Health Economics Consortium, University of York, York, UK

²Nutricia Research, Utrecht, The Netherlands

³Hospital Italiano, Buenos Aires, Argentina

⁴Pediatric Gastroenterology, Hepatology and Nutrition Division, Sheikh Khalifa Medical City, Abu Dhabi, United Arab Emirates

⁵Nationwide Children's Hospital, Columbus, Ohio, USA

⁶National University of Singapore, Singapore, Singapore

⁷University of Utrecht, Utrecht, The Netherlands

⁸Department of Paediatrics, Medical University of Warsaw, Warsaw, Poland

Contributors All authors gave input on the design and aim of the systematic review. HW, JG and TL designed the search strategy; HS, MM, SHQ, IL-W, MS and CL gave input to the search strategy and the inclusion and exclusion criteria; JM and JG defined the data extraction elements; JM, JG, ME and TL wrote the protocol and this manuscript and its revisions; HS, MM, SHQ, IL-W, MS and CL revised the protocol and this manuscript.

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Competing interests HS, MS, CL, MM and SHQ have served as consultants, advisory board members and/or speakers for companies manufacturing infant formulas, foods and probiotics or prebiotics. TL is an employee of Nutricia Research.

Ethics This is an SR of published studies and ethical committee approval is not required. The planned output for this SR is publication in a peer-reviewed journal.

Provenance and peer review Not commissioned; externally peer reviewed.

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REFERENCES

1. Brown M, Heine RG, Jordan B. Health and well-being in school-age children following persistent crying in infancy. *J Paediatr Child Health* 2009;45:254–62.
2. Rautava P, Lehtonen L, Helenius H, *et al.* Infantile colic: child and family three years later. *Pediatrics* 1995;96(Pt 1):43–7.
3. Vik T, Grote V, Escribano J, *et al.* Infantile colic, prolonged crying and maternal postnatal depression. *Acta Paediatr* 2009;98:1344–8.
4. Miller-Loncar C, Bigsby R, High P, *et al.* Infant colic and feeding difficulties. *Arch Dis Child* 2004;89:908–12.
5. Indrio F, Di Mauro A, Riezzo G, *et al.* Infantile colic, regurgitation, and constipation: an early traumatic insult in the development of functional gastrointestinal disorders in children? *Eur J Pediatr* 2015;174:841–2.
6. Vandenplas Y, Abkari A, Bellaiche M, *et al.* Prevalence and health outcomes of functional gastrointestinal symptoms in infants from birth to 12 months of age. *J Pediatr Gastroenterol Nutr* 2015;61:531–7.
7. Mitchell CM, Drossman DA. Survey of the AGA membership relating to patients with functional gastrointestinal disorders. *Gastroenterology* 1987;92(Pt 1):1282–4.
8. Drossman DA. The functional gastrointestinal disorders and the Rome III process. *Gastroenterology* 2006;130:1377–90.
9. Hyman PE, Milla PJ, Benninga MA, *et al.* Childhood functional gastrointestinal disorders: neonate/toddler. *Gastroenterology* 2006;130:1519–26.
10. van Tilburg MA, Hyman PE, Walker L, *et al.* Prevalence of functional gastrointestinal disorders in infants and toddlers. *J Pediatr* 2015;166:684–9.
11. Iacono G, Merolla R, D'Amico D, *et al.* Gastrointestinal symptoms in infancy: a population-based prospective study. *Dig Liver Dis* 2005;37:432–8.
12. Shamir R, St James-Roberts I, Di Lorenzo C, *et al.* Infant crying, colic, and gastrointestinal discomfort in early childhood: a review of the evidence and most plausible mechanisms. *J Pediatr Gastroenterol Nutr* 2013;57(Suppl 1):S1–45.
13. Lucassen P. Colic in infants. *BMJ Clin Evid* 2015;2015:pii:0309.
14. Garrison MM, Christakis DA. A systematic review of treatments for infant colic. *Pediatrics* 2000;106(Pt 2):184–90.
15. Drug and Therapeutics Bulletin. Management of infantile colic. *BMJ* 2013;347:f4102.
16. Harb T, Matsuyama M, David M, *et al.* Infant Colic—what works: a systematic review of interventions for breast-fed infants. *J Pediatr Gastroenterol Nutr* 2016;62:668–86.
17. Hall B, Chesters J, Robinson A. Infantile colic: a systematic review of medical and conventional therapies. *J Paediatr Child Health* 2012;48:128–37.
18. Vandenplas Y, Alarcon P, Alliet P, *et al.* Algorithms for managing infant constipation, colic, regurgitation and cow's milk allergy in formula-fed infants. *Acta Paediatr* 2015;104:449–57.
19. Vandenplas Y, Gutierrez-Castrellon P, Velasco-Benitez C, *et al.* Practical algorithms for managing common gastrointestinal symptoms in infants. *Nutrition* 2013;29:184–94.
20. Vandenplas Y, Rudolph CD, Di Lorenzo C, *et al.* Pediatric gastroesophageal reflux clinical practice guidelines: joint recommendations of the North American Society for Pediatric Gastroenterology, Hepatology, and Nutrition (NASPGHAN) and the European Society for Pediatric Gastroenterology, Hepatology, and Nutrition (ESPGHAN). *J Pediatr Gastroenterol Nutr* 2009;49:548–547.
21. Lightdale JR, Gremse DA, Section on Gastroenterology Hepatology, and Nutrition. Gastroesophageal reflux: management guidance for the pediatrician. *Pediatrics* 2013;131:e1684–95.
22. Gieruszczak-Bialek D, Konarska Z, Skórka A, *et al.* No effect of proton pump inhibitors on crying and irritability in infants: systematic review of randomized controlled trials. *J Pediatr* 2015;166:767–70 e3.
23. Vandenplas Y. Management of paediatric GERD. *Nat Rev Gastroenterol Hepatol* 2014;11:147–57.
24. Czinn SJ, Blanchard S. Gastroesophageal reflux disease in neonates and infants: when and how to treat. *Paediatr Drugs* 2013;15:19–27.
25. Lucassen PL, Assendelft WJ, van Eijk JT, *et al.* Systematic review of the occurrence of infantile colic in the community. *Arch Dis Child* 2001;84:398–403.
26. Quitadamo P, Miele E, Alongi A, *et al.* Italian survey on general pediatricians' approach to children with gastroesophageal reflux symptoms. *Eur J Pediatr* 2015;174:91–6.
27. Headley J, Northstone K. Medication administered to children from 0 to 7.5 years in the Avon Longitudinal Study of Parents and Children (ALSPAC). *Eur J Clin Pharmacol* 2007;63:189–95.
28. Lucassen P. Colic in infants. *BMJ Clin Evid* 2010;2010:pii:0309.
29. Illingworth C, Timmins J. Gripe water: what is it? Why is it given? *Health Visit* 1990;63:378.
30. Chinawa JM, Ubesie AC, Adimora GN, *et al.* Mothers' perception and management of abdominal colic in infants in Enugu, Nigeria. *Niger J Clin Pract* 2013;16:169–73.
31. Sas D, Enrione MA, Schwartz RH. *Pseudomonas aeruginosa* septic shock secondary to 'gripe water' ingestion. *Pediatr Infect Dis J* 2004;23:176–7.
32. Snyder J, Brown P. Complementary and alternative medicine in children: an analysis of the recent literature. *Curr Opin Pediatr* 2012;24:539–46.
33. Kheir AE. Infantile colic, facts and fiction. *Ital J Pediatr* 2012;38:34.
34. Barr RG. The normal crying curve: what do we really know? *Dev Med Child Neurol* 1990;32:356–62.
35. Morris S, James-Roberts IS, Sleep J, *et al.* Economic evaluation of strategies for managing crying and sleeping problems. *Arch Dis Child* 2001;84:15–19.
36. Sommers T, Corban C, Sengupta N, *et al.* Emergency department burden of constipation in the United States from 2006 to 2011. *Am J Gastroenterol* 2015;110:572–9.
37. Higgins JPT, Altman DG, Sterne JAC. Assessing risk of bias in included studies. In: Higgins JPT, Green S, eds. *Cochrane Handbook for Systematic Reviews of Interventions*, 2011.
38. Drummond MF, Jefferson TO. Guidelines for authors and peer reviewers of economic submissions to the BMJ. The BMJ economic evaluation working party. *BMJ* 1996;313:275–83.
39. Centre for Reviews and Dissemination. *Undertaking systematic reviews of research on effectiveness: CRD's guidance for those carrying out or commissioning reviews*. Report 4. 2nd edn. York: University of York, 2001.
40. Wells GA, Shea B, O'Connell D, *et al.* The Newcastle-Ottawa Scale (NOS) for assessing the quality of nonrandomised studies in meta-analyses. Secondary The Newcastle-Ottawa Scale (NOS) for assessing the quality of nonrandomised studies in meta-analyses [webpage]. 2014. http://www.ohri.ca/programs/clinical_epidemiology/oxford.asp
41. Herzog R, Álvarez-Pasquin MJ, Díaz C, *et al.* Are healthcare workers' intentions to vaccinate related to their knowledge, beliefs and attitudes? A systematic review. *BMC Public Health* 2013;13:154.
42. Philips Z, Ginnelly L, Sculpher M, *et al.* Review of guidelines for good practice in decision-analytic modelling in health technology assessment. *Health Technol Assess* 2004;8:iii–iv, ix–xi, 1–158. http://www.journalslibrary.nihr.ac.uk/_data/assets/pdf_file/0015/65121/FullReport-ha8360.pdf

Correction: *Costs associated with functional gastrointestinal disorders and related signs and symptoms in infants: a systematic review protocol*

Glanville J, Ludwig T, Lifschitz C, *et al*. Costs associated with functional gastrointestinal disorders and related signs and symptoms in infants: a systematic review protocol. *BMJ Open* 2016;6:e011475. The name segmentation of the 7th author is incorrect. The author's surname is Quak; first names are Seng Hock. This author should be cited as 'Quak SH'.

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