Parents training programmes for managing infantile colic (Protocol)

Thomas, Megan, Gordon, Morris, Banks, Shel S C and Wallace, Chris

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**Parent training programmes for managing infantile colic**

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**ABSTRACT**

This is a protocol for a Cochrane Review (Intervention). The objectives are as follows:

1. To evaluate the effectiveness and safety of parent training programmes for managing colic in infants under four months of age.
2. To identify the educational content and attributes of such published programmes.

**BACKGROUND**

**Description of the condition**

Infantile colic can be defined as periods of inconsolable, unexplained and incessant crying in a seemingly healthy infant that, quite understandably, leads to exhausted, frustrated and concerned parents seeking to comfort their child (Landgren 2010a). The prevalence of excessive crying varies according to the definition used, although, most often, it peaks during the second month of life with a prevalence of 1.5% to 11.9% (Reijneveld 2001). Traditionally, the definition of the condition was based on the rule of three, that is, unexplained episodes of paroxysmal crying for more than three hours per day, for three days per week, for at least three weeks (Wessel 1954). More recently a new definition has been proposed. It refers to a clinical condition of fussing and crying for at least one week in an otherwise healthy infant (Hyman 2006). Rome III includes infantile colic, with diagnostic criteria including all of the following in infants from birth to four months of age: paroxysms of irritability, fussing or crying that starts and stops without obvious cause; episodes lasting three or more hours per day and occurring at least three days per week for at least three weeks; and no failure to thrive (Mostafa 2008). Colic is a symptom rather than a condition or diagnosis in and of itself. The incidence of colic is estimated to be between 10% and 30% of infants born (Clifford 2002; Rosen 2007). Paroxysms of inconsolable crying are often accompanied by flushing of the face, meteorism (excessive flatulence in the intestinal tract with distention of the abdomen), drawing up of the legs, and flatulence (Savino 2007). Symptoms have historically typically started in the second week of life in both breast-fed and formula-fed infants and resolved by three months of age (Lucas 1998). Generally speaking, these symptoms are not indicative of disease and thus hospital admission for these infants is generally unnecessary, detrimental, and should not be encouraged (Savino 2007). However, about 5% of colicky crying infants do have a serious, underlying medical
problem (Freedman 2009; Savino 2005; Savino 2007), and there is evidence that older children presenting with migraine are more likely to have been babies who have suffered colic (Romanello 2013). Therefore, all colicky infants should undergo a complete medical assessment in order to exclude underlying medical conditions which require investigation and treatment (Savino 2010). The etiopathogenesis of infantile colic as a symptom remains undefined and is most likely multifactorial. Despite the common nature of the condition, and the large amount of research investigating this area, there have been no breakthroughs in terms of the real mechanisms underlying infant colic.

It has been suggested that a number of behavioural factors (psychological and social) and biological components (food hypersensitivity, allergy, gut microflora, bloating from trapped gas and dysmotility) can contribute to its manifestation (Gupta 2007). These include the following.

First, lactose intolerance - due to a relative lactase deficiency - has been identified as a possible causative factor in infant colic. Carbohydrate malabsorption leads to the colonic fermentation of sugars and an increase in the levels of hydrogen gas (Infante 2011). The rapid production of hydrogen in the lower bowel distends the colon, sometimes causing pain, whereas the osmotic pressures generated by lactose and lactic acid in the colon cause an influx of water, leading to further distension of the bowel. Although studies evaluating the degree of hydrogen in the breath of colicky infants have produced inconsistent results, increases in breath hydrogen levels have been reported (Hyams 1989; Miller 1990; Moore 1998).

Second, the immunological model, which focuses on possible allergens, has been suggested as a cause of colic. A key allergen is cows’ milk proteins in breast milk or infant formula. Intact proteins from the mother’s diet can sometimes cross over into the breast milk and provoke an allergic response and symptoms of colic in her infants. Consequently, a low-allergen maternal diet, or hypoallergenic infant formula (Iacovou 2012), has been proposed as a form of treatment (Hill 2005; Schach 2002). The possibility that infantile colic could be related to allergens was first described by Shannon 1921. Since then a number of studies have evaluated the possible association between colic and food hypersensitivity (Heine 2013; Heine 2014; Hill 1995; Iacono 1991; Lothe 1982; Merras-Salnio 2013; Saps 2011).

The evidence shows that about 25% of infants with moderate or severe symptoms have cows’ milk protein-dependent colic (Axelsson 1986; Hill 2000; Lindberg 1999), which improves after some days of a hypoallergenic diet (Campbell 1989; Dupont 2010; Estep 2000; Iacono 1991; Iacono 2005; Jakobsson 1983; Jakobsson 2000; Lothe 1999; Savino 2001). For these infants, infantile colic could be the first manifestation of atopic disease and, for this reason, dietetic treatment should be the first therapeutic approach (Gupta 2007; Hall 2012; Savino 2010). Indeed, dietary changes, such as eliminating cows’ milk proteins, are particularly indicated in cases of suspected intolerance to cows’ milk proteins (e.g. in infants with a positive family history, eczema or onset after the first month of life, and colic associated with other gastrointestinal symptoms such as vomiting or diarrhea) (Hill 1995; Hill 2005; Jakobsson 1983; Lucassen 2000; Savino 2014). However, UpToDate 2016 grades the introduction of hydrolysate formula for formula-fed infants or hypoallergenic diet for mothers of breast-fed infants as a “2C”: “a very weak recommendation; other alternatives may be equally reasonable”; “benefits and risks may be finely balanced, or the benefits and risks may be uncertain” and “the evidence comes from observational studies, unsystematic clinical experience, or from randomised, controlled trials with serious flaws”. A Cochrane review is underway to firm up the evidence (Savino 2014).

Third, there is growing evidence that the intestinal microbiota in colicky infants differ from those in non-colicky controls, since higher levels of anaerobic bacteria, such as coliform and Escherichia coli, microaerophilic bacteria, such as Helicobacter pylori (Ali 2012), and a lower concentration of Lactobacilli have been reported in infants with colic (Savino 2010). Human milk naturally contains these prebiotics; they are defined as indigestible oligosaccharides that could selectively enhance the proliferation of certain probiotic bacteria in the colon, especially Bifidobacterium species (Thomas 2010). Some studies have failed to find a protective effect of breast feeding on the development of colic in breast-fed infants (Clifford 2002), however, it is unclear if these studies compared exclusively breast-fed-from-birth infants with exclusively artificially-fed-from-birth infants, and so it is still not known whether any breastfeeding has some protective effect or whether any artificial feeding compromises the infant gut microbiome in some way. Evidence suggests that oligosaccharide prebiotics (a mixture of galacto-oligosaccharides and fructo-oligosaccharides) to encourage growth of the positive bacteria in the gut may be effective treatments for allergy and food intolerance in general (Arslanoglu 2012), and for crying in formula-fed infants (de Wierth 2013). Evidence is building around the effectiveness of supplementing the infant’s diet with probiotics to prevent colic and other symptoms (Oozer 2015).

These three pathophysiological models indicate implicit treatment modalities, however, various therapeutic interventions have been used for infant colic that take a symptom-reduction, focused approach. These include pain relief (Savino 2002; Savino 2012); probiotic supplementation (Indiro 2014); complementary and alternative medicines and nutritional supplements such as fennel extract (Harb 2015) and chamomile (Perry 2011); sucrose and glucose solutions (Markestad 1997); and physical treatments such as manipulation (Dobson 2012; Olafsdottir 2001), massage and reflexology (Huhtala 2000; Perry 2011). Although systematic reviews have failed to provide evidence of its efficacy in reducing colicky symptoms, by reducing trapped gas in the liquid of the
stomach, simethicone is still often used (Metcalf 1994). Various other physical treatments have been studied to reduce symptoms, including carrying (Barr 1991), which may affect baby in psychological or social ways, or address mechanical aspects such as crib vibration (Huhtala 2000); and acupuncture (Landgren 2010a; Landgren 2010b; Reinthal 2008; Skjeie 2013). Evidence of efficacy is not comprehensive (Garrison 2000).

Description of the intervention

An alternate approach that has been investigated is to focus on training, support and psychologically underpinned interventions for parents of infants with colic.

There is recognition of the role of parental anxiety in the reported incidence of colic and evidence that parental reassurance is successful in reducing reports of distress (Furlong 2012; Hiscock 2014; Taubman 1984; Taubman 1988; Wolfe 1994; Zwi 2011). Guidance and informal education are often delivered by healthcare professionals to accompany any intervention for infantile colic so that parents and carers may better understand the potential aetiologies and pursue various management and treatment options (see, for example, Cook 2012). Parental behavioural modifications have often been suggested both for breast-fed and formula-fed infants, including advice to carry the infant (Barr 1991), not to carry the infant (McKenzie 1991), and to try to understand the infant’s needs (Taubman 1984).

Whilst there have been some attempts to synthesise evidence-informed pathways for infantile colic that do include some parental resources, for example, NICE (National Institute for Health and Care Excellence) Clinical Knowledge Summaries, UpToDate Patient Information Tips/health professional information and Map of Medicine, it is well documented that the evidence base is poor and inconclusive. Indeed, NICE CKS 2014 states that their recommendations are pragmatic. There is currently no national or international consensus on best practice for such interventions.

Why it is important to do this review

There is no clarity as to the extent these components contribute to the overall efficacy of symptom reduction strategies or to parental anxiety levels. Some interventions have been ineffective such as the McRury 2010 study based on techniques found in a popular parenting book - The Happiest Baby - by Karp 2003. Given the clinical and methodological heterogeneity of studies on these interventions, the efficacy of these interventions in reducing infant colic remains inconclusive, at present. Established studies and reports may now be outdated (e.g. Schmitt 1987; Taubman 1984), and more recent reported approaches are based on different approaches (e.g. Hiscock 2014 is based on an intervention described in Cook 2012; Keefe 2005), and so an up-to-date systematic review using Cochrane methodology is required. It is also important to note that focus within the published body of work often seeks to assess ‘whether’ such training is effective (e.g. Hiscock 2014), and this can be considered of limited educational research value (Norcini 2011). This is important as, in this context, the intervention being considered is educational, and if this cannot be defined, it cannot be reliably and validly reproduced and disseminated in a systematic fashion. Therefore, equally relevant questions are ‘how’ it achieves this outcome, ‘why’ the teaching is effective and ‘for whom and when’ such training can be effective. A review and synthesis of the evidence must also address these items and, from an educational stance, identify a relevant theory from this evidence base (Haji 2013). This will support future professionals in understanding and delivering such an intervention in a reliable and reproducible manner. Even if such data are not explicit within primary studies, synthesis can highlight such outcomes, as has been increasingly shown in the field of health education (Gordon 2011; Gordon 2013).
This review sets out to consider the effectiveness of parent training programmes (when compared to other interventions), the safety of such programmes, and to identify the content and attributes underpinning such programmes.

**OBJECTIVES**

1. To evaluate the effectiveness and safety of parent training programmes for managing colic in infants under four months of age.
2. To identify the educational content and attributes of such published programmes.

**METHODS**

**Criteria for considering studies for this review**

**Types of studies**
Randomised controlled trials (RCTs) and quasi-RCTs.

**Types of participants**
Infants younger than four months of age who are already suffering from infantile colic as defined by the study, for example, Rome III (Mostafa 2008) or Wessel (Wessel 1954). Both breast-fed and formula-fed infants will be eligible.

**Types of interventions**
Any form of parental training programmes, alone or in combination, versus another intervention(s) or placebo. Examples of programme content may include:

1. normalisation material in any form;
2. soothing techniques; and
3. feeding management advice.

In the teaching forms of:

1. face-to-face courses;
2. online and e-learning;
3. printed materials;
4. home visits and coaching; and
5. remote support and counselling.

**Types of outcome measures**
For all proposed outcomes, we will use the final outcomes from the end of the trials, and we will record the timings of these outcomes as they may guide the subgroup analysis (see Subgroup analysis and investigation of heterogeneity).

**Primary outcomes**

1. A reduction in the duration of crying (post-treatment versus baseline). Data may be continuous (e.g. hours per day), or dichotomous (e.g. reduction under a predefined threshold, as determined by the trial authors).
2. Adverse effects, including parental depression and mental illness, choking, apparent life-threatening events (dichotomous outcome).

**Secondary outcomes**

1. The number of responders in each group after treatment: reduction in frequency of crying episodes per 24 hours (post-treatment versus baseline) (dichotomous outcome, as defined by the primary studies).
2. Parental or family quality of life, including measures of parental stress, anxiety or depression, as proposed by the primary studies and so no single scale may be possible (continuous outcome).
3. Infant sleep duration per 24 hours at seven, 14, and 21 days (post-treatment versus baseline) (continuous outcome).

**Search methods for identification of studies**
We will identify relevant trials by searching the sources described below.

**Electronic searches**
We will search the following electronic databases and trial registers from inception onwards:

1. Cochrane Central Register of Controlled Trials (CENTRAL; current issue) in the Cochrane Library, which includes Cochrane Developmental, Psychosocial and Learning Problems Specialised Register;
2. MEDLINE Ovid (1946 to date);
3. MEDLINE In-Process and Other Non-Indexed Citations Ovid (current issue);
4. MEDLINE Epub Ahead of Print Ovid (current issue);
5. Embase Ovid (1980 to date);
6. CINAHL Plus EBSCOhost (Cumulative Index to Nursing and Allied Health Literature; 1937 to date);
7. PsycINFO Ovid (1967 to date);
8. Science Citation Index - Expanded Web of Science (SCI; 1970 to date);
9. Social Sciences Citation Index Web of Science (SSCI; 1970 to date);
10. Conference Proceedings Citation Index - Science Web of Science (CPCI-S; 1990 to date);
11. Conference Proceedings Citation Index - Social Science and Humanities Web of Science (CPCI-SSH; 1990 to date);
12. LILACS (Latin American and Caribbean Health Science Information database; lilacs.bvsalud.org/en);
13. Cochrane Database of Systematic Reviews (CDSR; current issue) in the Cochrane Library;
14. Database of Abstracts of Reviews of Effects (DARE; current issue) in the Cochrane Library;
15. Epistemonikos (limited to systematic reviews; epistemonikos.org);
16. WorldCat (limited to theses; worldcat.org);
17. US National Institutes of Health Ongoing Trials Register ClinicalTrials.gov (clinicaltrials.gov); and
18. World Health Organization International Clinical Trials Registry Platform (WHO ICTRP; apps.who.int/trialsearch).

We will search MEDLINE using the search strategy in Appendix 1, which uses the sensitivity maximizing version of the Cochrane Highly Sensitive Search Strategy for identifying randomised trials (Lefebvre 2008). We will adapt this strategy for other databases. We will not impose any date or language restrictions. Studies published in languages other than English will be professionally translated in full.

Searching other resources

Grey literature

To assess more contemporaneous studies that have not yet been published in full, we will handsearch abstracts presented at relevant international meetings, including the European Society for Paediatric Gastroenterology, Hepatology and Nutrition (ESPGHAN) and the North American Society for Paediatric Gastroenterology, Hepatology and Nutrition (NASPGHAN), published from 2010 onwards. There is some evidence that data from abstracts can be inconsistent with data in published articles (Pitkin 1999). Therefore, we will only include abstract publications if sufficient data are presented to judge inclusion and assess quality. Where such data are not presented, we will attempt to contact authors for more information, and meanwhile will place the studies in 'Awaiting classification'.

Reference searching

We will inspect the references of included studies for more trials. We will search the bibliographies of included studies to identify any other potentially relevant articles.

Personal contacts

We will contact leaders in the field to try to identify other published and unpublished studies.

Data collection and analysis

Selection of studies

Two review authors (Morris Gordon (MG) and Shel Banks (SB)) will independently screen titles, abstracts, and full reports for eligibility against the inclusion criteria (see Criteria for considering studies for this review). Specifically, they will:
1. merge search results using reference management software and remove duplicate records of the same report;
2. examine titles and abstracts to remove irrelevant records;
3. retrieve full texts of potentially relevant reports;
4. link together multiple reports of the same study;
5. examine full-text reports for studies that meet the eligibility criteria;
6. correspond with investigators, when appropriate, to clarify study eligibility;
7. at all stages, note reasons for inclusion and exclusion of reports on a study flow spreadsheet, resolving any disagreements through consensus;
8. make final decisions on study inclusions and resolve any discrepancies through a process of consensus; and
9. proceed to data collection.

We will record our selection process in a PRISMA diagram (Moher 2009).

Data extraction and management

We will develop data extraction forms a priori, as per the recommendations in the Cochrane Handbook for Systematic Reviews of Interventions (Higgins 2011a). We will extract the information described below.
1. Characteristics of participants: source of participants, inclusion and exclusion criteria, total number at baseline, total number at completion, setting, definition of ‘colic’ applied, diagnostic criteria applied, type of feeding (breast feeding, formula feeding), age at onset of colic, age at commencement of intervention, and evaluation of potential effect modifiers (e.g. age, gender).
2. Characteristics of intervention: content of training, pedagogical methods employed, context, resources and educator details, any theoretical underpinning described.
3. Interventions and controls: number of groups, intervention(s) applied, frequency and duration of treatment, total number of treatments, permitted cointerventions.
4. Outcomes of interest: primary and secondary outcomes, measurement and assessment.

We will conduct data analysis using Review Manager (RevMan) 5.3 (Higgins 2014) by the Cochrane Collaboration. Where possible, we will contact study authors to obtain additional or unreported data. We will conduct meta-analyses using random-effects models, unless there is evidence of homogeneity, in which case we will use a fixed-effects model.
4. Methods: study design, duration, sequence generation, allocation concealment, blinding of outcome assessors, evaluation of success of blinding.

5. Outcomes: list of outcomes assessed, definitions used, values of means and standard deviations (SDs) at baseline and at time points as defined by the study protocol (or change from baseline measures, if given).

6. Results: measures at end of protocol, follow-up data (including means and SDs, standard errors, or confidence intervals (CI) for continuous data, and summary tables for dichotomous data), withdrawals, and losses to follow-up.

7. Other: references to other relevant studies, points to follow up with authors, comments from the authors, key conclusions from the study (by the authors), other comments from review authors.

Two review authors (MG and SB) will independently extract the data using the data extraction form. A third review author (Megan R Thomas (MRT)) will resolve any disagreements. We will collate data in the latest version of Review Manager 2014.

Assessment of risk of bias in included studies

Two review authors (MG and SB) will independently evaluate each study for risk of bias using the criteria recommended in the Cochrane Handbook for Systematic Reviews of Interventions (Deeks 2011; Higgins 2011b) for the following domains: sequence generation; allocation concealment; blinding of parents and health professionals; blinding of outcome assessment; incomplete outcome data; selective outcome reporting; and other potential threats to validity. We will judge each domain as being at ‘low’, ‘high’, or ‘unclear’ risk of bias. We will compare the judgments, and discuss and resolve any inconsistencies in the assessments. A third review author (MRT) will resolve any persisting disagreements. We will complete a ‘Risk of bias’ table for each included study and present the ‘Risk of bias’ summary graphically.

Sequence generation

We will include only RCTs or quasi-RCTs in this review. We will assess randomisation as being at low risk of bias if the procedure of sequence generation was explicitly described to confirm it was random; examples include computer-generated random numbers, a random numbers table or coin-tossing. If no description is given, we will contact the authors for further information and if we fail to receive a response, we will assign a judgment of unclear risk of bias. We will consider studies that use non-randomised procedures to be at high risk of bias.

Allocation concealment

We will assess concealment of treatment allocation as being at low risk of bias if the procedure was explicitly described and adequate efforts were made to ensure that intervention allocations could not have been foreseen in advance of, or during, enrolment; examples include centralized randomisation, numbered or coded containers, or sealed envelopes. Procedures considered to have a high risk of bias include alternation or reference to case record numbers or dates of birth. We will also assign a high risk of bias if allocation concealment did not occur, as intervention allocations could have been foreseen and thus introduced potential bias. If no description is given, we will contact the study authors and if no response is received, we will assign a judgment of unclear risk of bias.

Blinding of parents and health professionals

In this context, the intervention is administered by parents and so, in effect, they will be considered the target of the blinding procedures. Indeed, as the participants will be less than four months of age by the defined inclusion criteria, it is deemed that this item is not applicable to them. Furthermore, parents often act as outcome assessors. We will primarily assess the risk of bias associated with the blinding of parents of participants based on the likelihood that such blinding was sufficient to ensure that parents had no knowledge as to which intervention they, on behalf of the infant, received.

Blinding of outcome assessment

We will describe, for each included study, the methods used, if any, to blind the outcome assessors from knowledge of which intervention a participant received. We will judge studies to be at low risk of bias if they were blinded or if we consider that the lack of blinding could not have affected the results. If blinding was not done, or not possible due to the nature of intervention, we will judge the study to be at high risk of bias because it is possible that the lack of blinding influenced the results. If no description is given, we will contact the study authors for more information, and if we do not receive a response, we will assign a judgment of unclear risk of bias. We will note the blinding of health professionals, if reported.

Incomplete outcome data

Incomplete outcome data essentially include attrition, exclusions, and missing data.

We will assign a judgment of low risk of bias if:
1. participants included in the analysis are exactly those who were randomised into the trial, if missing outcome data are balanced in terms of numbers across intervention groups with similar reasons for missing data across groups, or if there are no missing outcome data;
2. for dichotomous outcome data, the proportion of missing outcomes compared with observed event risk is not sufficient to have a clinically relevant impact on the intervention effect estimate;
3. for continuous outcome data, the plausible effect size (standardized mean differences (SMD)) among missing outcomes is not sufficient to have a clinically relevant impact on observed effect size; or
4. missing data have been imputed using appropriate methods.

We will assess the reporting of outcomes as being at low risk of bias if:
1. there is sufficient reporting of attrition or exclusions, or both, to permit a judgment of low or high risk of bias; or
2. the study reported incomplete outcome data; or
3. the numbers randomised to intervention and control groups are not clearly reported.

Selective outcome reporting

We will assess the reporting of outcomes as being at low risk of bias if all the study outcomes declared in the methods section have been reported in the results. We will also evaluate whether different reports of the study are available, including protocols, and examine them to ensure there is no suggestion of selective outcome reporting. If no description is given, we will contact the authors for more information and if no response is received, we will assign a judgment of unclear risk of bias. If there is evidence of selective reporting (such as a significant departure from the protocol or key outcomes that were due to be investigated are not reported), we will assign a judgment of high risk of bias.

Other potential threats to validity

If the study is at risk of other sources of bias, we will assess it as being at high risk of bias. For instance, if it was stopped early due to a data-dependent process, having a baseline imbalance between the groups, or sources of sponsorship or funding. We will assess the study as being at low risk of bias if it appears to be free from such threats to validity. When the risk of bias is unclear from published information, we will attempt to contact authors for clarification. If this is not forthcoming, we will assess these studies as being at unclear risk of bias.

Measures of treatment effect

Dichotomous data

We will present dichotomous data as risk ratios (RR), since the effects of the RR are readily understood (Walter 2000). We will report all dichotomous data with their associated 95% CI and probabilities of control and treatment groups (where possible).

Continuous data

If all studies use the same measurement scale, we will calculate mean differences (MD) for change scores. Where studies use different scales, we will calculate the SMD using Hedges’ (adjusted) g. If necessary, we will calculate effect estimates from P values, t-statistic, analysis of variance (ANOVA) tables, or other statistics as recommended in the Cochrane Handbook for Systematic Reviews of Interventions (Deeks 2011).

For this analysis we will use, according to need, either change scores or final values, without combining them: we will not combine these two different indices in meta-analysis and will only meta-analyse homogeneous data sets.

If both continuous and dichotomous data are available for an outcome, we will include only the continuous outcome in the primary analysis. If some studies report an outcome as a dichotomous measure, and others use a continuous measure of the same construct, we will convert the results for the former from the dichotomous measure to a SMD, provided that we can assume the underlying continuous measure has approximately a normal or logistic distribution (otherwise we will carry out two separate analyses).

Unit of analysis issues

Cluster-randomised trials

For each included study, we will determine whether the unit of analysis is appropriate for the unit of randomisation and the design of that study (that is, whether the number of observations matches the number of ‘units’ that were randomised (Deeks 2011)). It is unlikely that we will find cluster-randomised trials because this design is uncommon in this field. However, if we do, we will use the intraclass correlation coefficient (ICC) to convert trials to their effective sample size before incorporating them into the meta-analysis, as recommended in the Cochrane Handbook for Systematic Reviews of Interventions (Higgins 2011b). When the ICC cannot be used, we will use values available in the published literature as an
Dealing with missing data
Where data are missing, we will contact the corresponding authors of included studies to supply any unreported data. For all outcomes, in all studies, we will carry out analyses as far as possible on an intention-to-treat basis; that is, we will attempt to include all participants randomised to each group in the analyses, and we will analyse all participants in the group to which they were allocated, regardless of whether or not they received the allocated intervention. For missing statistics (missing SDs), we will estimate SDs from other available data, such as standard errors, or we will impute them using the methods suggested in Higgins 2011b. We will make no assumptions about loss to follow-up for continuous data, and we will base analyses on those participants completing the trial. If there is a discrepancy between the number randomised and the number analysed in each treatment group, we will calculate and report the percentage lost to follow-up in each group.

Assessment of heterogeneity
We will assess clinical heterogeneity by comparing the distribution of important participant characteristics between trials (e.g. age) and trial characteristics (e.g. randomisation, concealment, blinding of outcome assessment, losses to follow-up, treatment type, cointerventions). We will assess statistical heterogeneity by examining the I² statistic (Deeks 2011), a quantity that describes the proportion of variation in point estimates that is due to variability across studies rather than sampling error. We will interpret the I² statistic as suggested in the latest version of Deeks 2011:

1. 0% to 40%: might not be important;
2. 30% to 60%: may represent moderate heterogeneity;
3. 50% to 90%: may represent substantial heterogeneity; or
4. 75% to 100%: suggests considerable heterogeneity.

We will employ a Chi² test of homogeneity, with a 10% level of significance, to determine the strength of evidence that heterogeneity is genuine. We will also report Tau². Once data have been extracted, clinical and methodological heterogeneity will be judged by discussion between authors (see Subgroup analysis and investigation of heterogeneity).

Assessment of reporting biases
In order to minimize publication bias, we will attempt to obtain the results of any unpublished studies in order to compare the results extracted from published journal reports with the results obtained from other sources (including correspondence).
In addition, if there are more than 10 studies grouped in a comparison, we will evaluate whether reporting biases are present by using funnel plots to investigate any relationship between effect estimates and study size or precision, or both, as recommended in the Cochrane Handbook for Systematic Reviews of Interventions (Sterne 2011).

**Data synthesis**

Where interventions are similar in type of parental training programme and type of outcome assessed, we plan to group the studies and synthesize their results in a meta-analysis. We will present results for each combination of parental training programme, assessed outcome and colic type, with the exception of those studies for which no data are observed. For instance, if two or more studies assess the effects of a parental training programme for parents of otherwise healthy children with colic and both measure the daily crying, we will perform a meta-analysis of the results. Because we assume that clinical heterogeneity is very likely to impact on our review results, given the wide breadth and types of interventions included, we will combine the studies using a random-effects model, regardless of evidence of statistical heterogeneity. We will calculate all overall effects using inverse variance methods. We will carry out statistical analysis using Review Manager 2014. Where data are insufficient to allow meta-analysis or qualitative analysis, we will provide a narrative synthesis and descriptive summary of the study outcomes.

While there may be heterogeneity in the interventions, as well as the comparisons, we consider that the consensus on definitions of symptoms for eligibility manages the risk of ‘blurring’ the results. However, we remain vigilant and if a risk is perceived upon evaluation of our findings, a sensitivity analysis removing such trials to provide more definite findings may be required.

**Subgroup analysis and investigation of heterogeneity**

Large numbers of subgroup analyses may lead to misleading conclusions (Oxman 1992; Yusuf 1991). We plan to carry out the following subgroup analyses:

1. type of feeding (bottle-fed versus breast-fed);
2. short-term and long-term follow-up (four weeks or less versus more than four weeks of treatment);
3. type of parental training (face-to-face versus distance or written materials); and
4. crying time in consideration with infant’s age.

These analyses will be exploratory as they will involve non-experimental (cross-study) comparisons on primary outcomes only. We will treat any conclusions with caution.

**Sensitivity analysis**

We will conduct sensitivity analyses to determine whether findings are sensitive to:

1. trials with low levels of potential bias versus trials with high levels of potential bias;
2. missing information or missing results (using 20% attrition as ‘high risk’ of bias, see above);
3. the definition of colic used, by conducting analyses on studies using the stringent Wessel definition of infant colic (Wessel 1954), the more recent definition given by Hyman 2006, and a non-recognized definition; and
4. the choice of model used, by comparing results from the random-effects model, which we are using, with those from the fixed-effects model.

**Presentation of main results**

We will assess the overall quality of evidence using the GRADE approach (Guyatt 2008). We will present the results of our assessment in a ‘Summary of findings’ table, per comparison, created using GRADEpro GDT software (GRADEpro GDT 2014). The comparisons used will be those outlined in the studies, and so may be another intervention(s) or placebo. Our table will include information on the type of participants, the interventions and comparisons used in each case, and the outcomes and their measurements for each study, as well as the setting and the length of follow-up. The GRADE approach appraises the quality of a body of evidence based on the extent to which one can be confident that an estimate of effect, or association, reflects the item being assessed. RCTs start as high-quality evidence but may be downgraded due to: risk of bias (methodological quality), indirectness of evidence, unexplained heterogeneity, imprecision (sparse data), and publication bias. Intention-to-treat data would be of better quality than per protocol results. Two review authors (SB and MG) will independently assess and agree the overall quality of the evidence for each outcome after considering each of these factors, and will grade them as:

1. high quality: further research is very unlikely to change confidence in the estimate of effect;
2. moderate quality: further research is likely to have an important impact on confidence in the estimate of effect, and may change the estimate;
3. low quality: further research is very likely to have an important impact on confidence in the estimate of effect, and is likely to change the estimate; or
4. very low quality: any estimate of effect is very uncertain.

We will include any rationale for the downgrading of the quality of the evidence in the footnotes of the table.

We will take the assumed risk from the median of the quality of the evidence in the footnotes of the table. We will use the following outcomes:

1. a reduction in the duration of crying (post-treatment versus baseline);
2. adverse effects, including parental depression and mental illness, choking, apparent life-threatening events;
3. the number of responders in each group after treatment: reduction in frequency of crying episodes per 24 hours (post-treatment versus baseline);
4. parental or family quality of life, including measures of parental stress, anxiety or depression, as proposed by the primary studies and so no single scale may be possible; and
5. infant sleep duration per 24 hours at seven, 14, and 21 days (post-treatment versus baseline).

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Parent training programmes for managing infantile colic (Protocol)

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St James-Roberts 1991

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Taubman 1988
APPENDICES

Appendix 1. MEDLINE search strategy

1 colic/
2 colic$.tw.
3 ((stomach or abdominal or abdomen$) adj3 (spasm$ or pain$ or cramp$)).tw.
4 ((gastric or gastro$) adj3 (spasm$ or pain$ or cramp$)).tw.
5 crying/
6 (cry or crying or cries).tw.
7 or/1-6
8 exp Infant/
9 (baby or babies or infant$ or child$ or newborn$ or neonat$).tw.
10 or/8-9
11 7 and 10
12 exp Parents/
13 Parenting/
14 exp Parent-Child Relations/
15 family relations/
16 exp maternal behavior/
17 maternal deprivation/
18 paternal behavior/

* Indicates the major publication for the study

Wessel 1954

Wolfe 1994

Wright 2003

Yusuf 1991

Zwi 2011
CONTRIBUTIONS OF AUTHORS

Shel Banks and Morris Gordon wrote the protocol. Megan R Thomas reviewed the protocol and is lead author. Chris Wallace reviewed the protocol. Morris Gordon has overall responsibility for managing the review.

DECLARATIONS OF INTEREST

Megan R Thomas - none known.

Morris Gordon (MG) - has received travel grants in the last three years from Ferring Pharmaceuticals and Biogaia to attend scientific meetings to present the results of previous and ongoing Cochrane systematic reviews on probiotics amongst other therapies. These companies have had no involvement in this or any other research or synthesis project by MG in terms of conception, planning, design, carrying out or writing up works. They have had no editorial input or control in this or any other previous works by MG. MG has received travel grants from Ferring Pharmaceuticals, Vifor Pharma, Danone/Nutricia, Abbott and Tillots Pharma AG to attend meetings to present results of previous works. They have had no input or involvement in any aspect of the review process during this or previous systematic reviews carried out by MG. MG declares that whilst Ferring and Biogaia produce products that can be used for colic, from the author team’s scope of the literature they do not expect these to be used in any of the trials in this review. MG also declares that Danone/Nutricia produce formulas that may be competing treatments for colic in this review, but Abbott, Tillots and Vifor have no products of interest. MG has had involvement as an author on Cochrane reviews, including ‘Bowel preparation for paediatric colonoscopy’ (Gordon 2012) and ‘Probiotics for maintenance of remission in ulcerative colitis’ (Naidoo 2011). MG is a Paediatrician with an interest in gastroenterology. This involves seeing patients referred with infantile colic and managing them in line with current accepted evidence-based practice.
Shel SC Banks (SSCB) - is being paid as a Research Assistant for this review from Blackpool Teaching Hospitals NHS Foundation Trust. SSCB is Chair of the Local Infant Feeding Information Board (LIFIB) - providing syntheses of evidence-based information on infant feeding topics for health professionals. SSCB works as a Consultant for LIFIB and the Sudden Unexpected Death of a Child Prevention Team. Until March 2016, this post was funded by money from the local authority via Breastfeeding Network and SSCB was paid by the hour. SSCB also works as an Internationally Board Certified Lactation Consultant in private practice. SSCB declares that neither she personally nor any of the entities she represents take funding of any kind from any commercial interests in infant feeding or early years and that she works completely within the professional code of ethics as an Internationally Board Certified Lactation Consultant.

Chris Wallace (CW) - was involved in a review looking at probiotics as a treatment for chronic constipation, which was accepted for poster presentation at the European Society for Paediatric Gastroenterology, Hepatology and Nutrition (ESPGHAN) annual meeting. CW received funding for travel and accommodation from the Young Investigators Award (YIA) to attend the meeting. Chris has been an author on a Cochrane review entitled probiotics for chronic constipation in childhood, and as a member of ESPGHAN, he applied for a young investigator award to present findings at the annual meeting in 2016. ESPGHAN had no involvement in the project.

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**Internal sources**
- Blackpool Teaching Hospitals NHS Foundation Trust, UK.

Blackpool Teaching Hospitals is the employer of all four review authors: Morris Gordon and Megan R Thomas are employed in the medical team for the hospital, Chris Wallace is employed as a Foundation Year 2 Doctor and Shel SC Banks is employed as a Project Manager in infant feeding, with a 12-month sabbatical as Research Assistant to complete this review.

The views and opinions expressed therein are those of the review authors and do not necessarily reflect those of Blackpool Teaching Hospitals NHS Foundation Trust.

**External sources**
- None, Other.