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Carbohydrate supplementation of human milk to promote growth in preterm infants (Review)

Amissah EA, Brown J, Harding JE

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Carbohydrate supplementation of human milk to promote growth in preterm infants (Review)

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[Intervention Review]

Carbohydrate supplementation of human milk to promote growth in preterm infants

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ABSTRACT

Background

Preterm infants are born with low glycogen stores and require higher glucose intake to match fetal accretion rates. In spite of the myriad benefits of breast milk for preterm infants, it may not adequately meet the needs of these rapidly growing infants. Supplementing human milk with carbohydrates may help. However, there is a paucity of data on assessment of benefits or harms of carbohydrate supplementation of human milk to promote growth in preterm infants. This is a 2020 update of a Cochrane Review first published in 1999.

Objectives

To determine whether human milk supplemented with carbohydrate compared with unsupplemented human milk fed to preterm infants improves growth, body composition, and cardio-metabolic and neurodevelopmental outcomes without significant adverse effects.

Search methods

We used the standard search strategy of Cochrane Neonatal to search Cochrane Central Register of Controlled Trials (CENTRAL 2019, Issue 8) in the Cochrane Library and MEDLINE via PubMed on 22 August 2019. We also searched clinical trials databases and the reference lists of retrieved articles for randomised controlled trials and quasi-randomised trials.

Selection criteria

Published and unpublished controlled trials were eligible if they used random or quasi-random methods to allocate preterm infants in hospital fed human milk to supplementation or no supplementation with additional carbohydrate.

Data collection and analysis

Two review authors independently abstracted data and assessed trial quality and the quality of evidence at the outcome level using the Grading of Recommendations Assessment, Development and Evaluation (GRADE) method. We planned to perform meta-analyses using risk ratios (RRs) for dichotomous data and mean differences (MDs) for continuous data, with their respective 95% confidence intervals (CIs). We planned to use a fixed-effect model and to explore potential causes of heterogeneity via sensitivity analyses. We contacted study authors for additional information.

Main results

One unblinded, quasi-randomised controlled trial (RCT) assessing effects of carbohydrate supplementation of human milk in the form of a prebiotic in 75 preterm infants was eligible for inclusion in this review. We identified two publications of the same trial, which reported different methods regarding blinding and randomisation. Study authors confirmed that these publications pertain to the same trial, but they have not yet clarified which method is correct. Our analyses showed very low certainty evidence of an effect of carbohydrate

supplementation on weight by 30 days of age (MD 160.4 grams, 95% CI 12.4 to 308.4 grams), and no effect on risk of feeding intolerance (RR 0.64, 95% CI 0.36 to 1.15) or necrotising enterocolitis (NEC) (RR 0.2, 95% CI 0.02 to 1.3). Duration of hospital stay was shorter in the prebiotic group than in the control group (median difference 9 days). No data were available for assessing effects of carbohydrate supplementation on long-term growth and neurodevelopment.

Authors' conclusions

We are uncertain whether carbohydrate supplementation of human milk affects any outcomes in preterm infants. The only trial included in this review presented very low-quality evidence, and study authors provided uncertain information about study methods and analysis. The evidence may be limited in its applicability because researchers included a small sample of preterm infants from a single centre. Future trials could assess the safety and efficacy of different types and concentrations of carbohydrate supplementation for preterm infants fed human milk. However, we do not envisage that further trials of digestible carbohydrates will be conducted, as this is currently done as a component of multi-nutrient human milk fortification.

PLAIN LANGUAGE SUMMARY

Carbohydrate supplementation of human milk to promote growth in preterm infants

Review question

Does addition of extra carbohydrate to human milk fed to preterm infants compared with no additional carbohydrate improve growth, body fat, obesity, heart problems, high blood sugar, and brain development without causing significant side effects?

Background

Not enough carbohydrate intake in preterm infants may result in poor growth and development. Breast milk is the best food for preterm infants, but feeding them only breast milk may be nutritionally inadequate. Adding carbohydrate to breast milk may help. However, not enough data can be found on assessing the benefits and harms of adding carbohydrate to breast milk to promote growth in preterm infants.

Study characteristics

We found one trial involving 75 preterm infants with very low-quality evidence on the effects of adding extra prebiotics (a type of carbohydrate) to human milk in preterm infants. A second publication by the same study authors reported different methods regarding blinding and randomisation of the trial. Study authors confirmed that these publications describe the same trial but have not yet clarified which method is accurate. We were unable to reproduce the analyses from the data presented. The search is up to date as of August 2019.

Key results

Prebiotic carbohydrate supplementation increased the mean weight of preterm infants at day 30 and resulted in a shorter hospital stay compared with control. No evidence shows a clear difference in risk of feeding intolerance or necrotising enterocolitis between the prebiotic-supplemented and unsupplemented groups. No other data were available to show the effects of adding extra carbohydrate to human milk on short- and long-term growth, body fat, obesity, brain development, and heart problems.

Conclusions

Evidence on the short- and long-term effects of adding extra carbohydrate to human milk in preterm infants is lacking. This systematic review found very low-quality evidence on the effects of adding prebiotic carbohydrate to human milk in preterm infants, along with uncertainties about methods and analysis. The single trial included a small sample of Iranian preterm infants, and so the evidence may be considered as not generalisable. However, the outcomes assessed are common to all preterm infants, and the trial shows that adding prebiotic carbohydrate to human milk is possible in developing countries. Further research is needed to assess the benefits and harms of different types and concentrations of carbohydrate supplementation for preterm infants fed human milk. Currently, digestible carbohydrate supplementation in preterm infants is provided as a component of multi-nutrient human milk fortification.

SUMMARY OF FINDINGS

Summary of findings 1. Carbohydrate supplementation compared to control in preterm infants

Carbohydrate supplementation compared to control in preterm infants

Patient or population: preterm infants

Setting: tertiary neonatal units of Alzahra and Shahid Beheshti Hospital in Iran

Intervention: carbohydrate (prebiotic) supplementation

Comparison: no carbohydrate (prebiotic) supplementation

| Outcomes | Anticipated absolute effects* (95% CI) | | Relative effect (95% CI) | N° of participants (studies) | Certainty of the evidence (GRADE) | Comments |
|-----------------------------|--|--|--------------------------|------------------------------|-----------------------------------|--|
| | Risk with control | Risk with carbohydrate supplementation | | | | |
| Growth/Weight at day 30 | Mean weight 1542.4 g | Mean weight increased by 160.4 g (12.4 to 308.4) | - | 75 (1 RCT) | ⊕⊕⊕⊕ Very low ^{a,b,c} | No other data were reported on growth except for weight at day 30. |
| Neurodevelopmental outcomes | - | - | - | - | - | No data were reported for this outcome in the included trial. |
| Duration of hospital stay | - | - | - | 75 (1 RCT) | ⊕⊕⊕⊕ Very low ^{a,b,c} | The trial reported data on duration of hospital stay using median (range) for the prebiotic-supplemented and unsupplemented groups as 16 (9 to 45) (95% CI 15.34 to 24.09 days) and 25 (11 to 80) (95% CI 25.52 to 34.39 days), respectively. We have reported this outcome in the text of the review. |
| Feeding Intolerance | 560 per 1000 | 358 per 1000 (202 to 644) | RR 0.64 (0.36 to 1.15) | 75 (1 RCT) | ⊕⊕⊕⊕ Very low ^{a,b,c} | Study authors defined feeding intolerance as "gastric residue, i.e. the presence of milk in the stomach two hours after completion of a feeding". However another reported outcome was "requiring to cut off milk", which was similar to our pre-specified definition of feeding intolerance, i.e. resulting in cessation or reduction in enteral feeding. Thus, we used "requiring to cut off milk" in the analysis of feeding intolerance. |

| | | | | | | |
|---------------------------|--------------|---------------------------|--------------------------|---------------|-----------------------------------|--|
| Necrotising enterocolitis | 220 per 1000 | 40 per 1000 (4 to 293) | RR 0.18 (0.02 to 1.3) | 75 (1 RCT) | ⊕○○○ Very low ^{a,b,c} | Definition was suspected NEC, which was based on clinical assessment |
|---------------------------|--------------|---------------------------|--------------------------|---------------|-----------------------------------|--|

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: confidence interval; RCT: randomised controlled trial; RR: risk ratio.

GRADE Working Group grades of evidence

High certainty: we are very confident that the true effect lies close to that of the estimate of the effect.

Moderate certainty: we are moderately confident in the effect estimate: the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different.

Low certainty: our confidence in the effect estimate is limited: the true effect may be substantially different from the estimate of the effect.

Very low certainty: we have very little confidence in the effect estimate: the true effect is likely to be substantially different from the estimate of effect.

^aDowngraded one level for risk of bias: the trial lacked methodological details and caregivers were not masked.

^bDowngraded one level for indirectness: we did not get any response from study authors for clarification on dosage, frequency, and duration of administration of the intervention.

^cDowngraded two levels for serious imprecision: small sample size, few events, and wide confidence intervals.

BACKGROUND

Description of the condition

Infants born preterm have low glycogen reserves because they are born before the phase of rapid glycogen accumulation in the third trimester of pregnancy (Velaphi 2011). To compensate for prenatal and postnatal deficits, and to match fetal accretion rates, preterm infants require higher glucose intakes and have higher glucose synthetic rates than full-term infants (Fenton 2013). For example, the rate of glucose synthesis among preterm infants at 28 weeks' gestation is 6 to 8 mg/min/kg compared to 3 to 5 mg/min/kg in full-term infants (Hay 2008).

Lactose, the most abundant carbohydrate in human milk, is the least variable among milk macronutrients (Ballard 2013). However, its concentration decreases in human milk with decreasing gestational age (Mahajan 2017). Lactose facilitates the absorption of two essential minerals - calcium and magnesium (Gregory 2005; Martin 2016), and it provides about 40% of the caloric intake of preterm infants (Elzouki 2012). Lactose is a disaccharide that is metabolised in the small intestine by brush border lactase to glucose and galactose (Cowett 2012). Of these two metabolites, glucose is the main endogenous substrate for energy production in the brain, and galactose is essential for the production of galactolipids, which are required for the infant's brain development (Elzouki 2012).

Human milk oligosaccharides (HMOs), the second most abundant carbohydrate in human milk, function as immunological agents (prebiotics) to promote the growth of selective commensal gut bacteria (Ballard 2013). They exhibit anti-infective properties by serving as soluble decoys that alter bacterial adhesion to intestinal walls (Jantscher-Krenn 2012), thereby reducing the risk of sepsis. Specific HMOs act to enhance gastrointestinal immunity and decrease the risk of necrotising enterocolitis (NEC) (Bode 2012). Insufficient consumption of these carbohydrates by the preterm infant may, therefore, have adverse effects on growth, intestinal colonisation, immune maturation, and neurological functioning (Morrow 2011).

Human milk, the ideal enteral nutrition for preterm infants, optimises neurodevelopmental outcomes (Belfort 2016). It decreases the incidence of non-specific gastrointestinal tract infections by 64% (Duijts 2010), and it reduces the incidence of NEC by 77%, compared with cow's milk-based infant formula (Sullivan 2010). Human milk feeding is associated with fewer re-hospitalisations in the first year of life (Vohr 2006), and unlike infant formula, human milk contains a wide range of HMOs (Bode 2012).

However, in spite of its numerous benefits, unsupplemented human milk may not meet the nutritional needs of preterm infants. First, its HMO content is variable. It shows intraindividual and interindividual variations (Blank 2012), and it varies according to maternal genetics (i.e. secretor and Lewis blood group status) and by stage of lactation. HMO concentration is highest in colostrum and decreases with lactational changes from transitional to mature milk (Bode 2012; Gabrielli 2011).

Second, the developmental deficiency of the lactase enzyme in preterm infants may interfere with complete digestion of lactose from breast milk (Ayede 2011). For instance, infants born at between 28 and 34 gestational weeks have only 30% of the lactase

activity of term infants (Fanaroff 2012). This deficiency could limit their ability to maximally derive energy from lactose in breast milk for optimal growth (Blackburn 2017). Although this inefficiency of dietary energy utilisation is salvageable in the colon through fermentation of undigested lactose, some of the energy is lost as heat (Erasmus 2002).

Therefore, for preterm infants who need larger amounts of energy and carbohydrate, carbohydrate supplements are sometimes added to human milk.

Description of the intervention

Carbohydrate supplements occur as monosaccharides, disaccharides, oligosaccharides, or polysaccharides. They may be derived from cow's milk, human milk, or soy milk or may be synthetically made. They are commercially available as modular supplements or as components of multi-nutrient supplements in liquid or powder form. Carbohydrate supplements may also be found in mixed forms designed to achieve an intended effect (e.g. a disaccharide/polysaccharide mix of glucose polymers and lactose to enhance carbohydrate absorption because of concerns about limited lactase activity in preterm infants) (Duggan 2008). Glucose polymers provide high caloric densities without increasing the osmotic load (Hay 2017), and unlike lactose, they are better absorbed, as they do not require lactase for digestion but are dependent on glucoamylase, which is available in sufficient quantities (Cowett 2012). However, in preterm infants, it is recommended that lactose should account for 40% to 100% of the carbohydrate intake (Klein 2002), as it is needed to aid mineral absorption and to foster prebiotic and lactase activities (Blackburn 2017).

Carbohydrate supplements are fed enterally to preterm infants once they begin to tolerate breast milk feeds. Like other macronutrients, they are commonly administered as a fixed dose per unit volume of breast milk, also known as 'standardised fortification' (Mangili 2017).

How the intervention might work

Carbohydrate digestion in the preterm infant is dependent on the composition of the carbohydrate ingested, the functional maturity of intestinal hydrolytic enzymes, and the gastrointestinal system (Elzouki 2012). Unlike other hydrolytic enzymes such as glucoamylase, which are embedded deeper in villus enterocytes, lactase is located at the tips of intestinal villi, making it vulnerable to intestinal mucosal injury (Duggan 2008). Thus, carbohydrate supplementation is expected to increase concentrations of carbohydrate in human milk and may increase mucosal uptake in cases of intestinal injury (Blackburn 2017). Higher concentrations of carbohydrate fed to preterm infants increase caloric density and thus could contribute to faster growth (Brown 2014).

Why it is important to do this review

Carbohydrate supplementation of human milk could enhance the optimal growth and health of preterm infants. However, it confers potential risks. For example, excessive intake may result in hyperglycaemia and transient symptoms of malabsorption including diarrhoea, flatulence, abdominal distension, and pain (Heine 2017). Thus, a systematic review of available evidence on the efficacy and safety of carbohydrate supplements in preterm infants is needed to clarify the uncertainties surrounding this intervention.

This is an update of a previously published Cochrane Review ([Kuschel 2000](#)).

OBJECTIVES

To determine whether human milk supplemented with carbohydrate compared with unsupplemented human milk fed to preterm infants improves growth, body composition, and cardio-metabolic and neurodevelopmental outcomes without causing significant adverse effects.

METHODS

Criteria for considering studies for this review

Types of studies

We considered published and unpublished controlled trials utilising random or quasi-random patient allocation for inclusion in this review. We excluded cross-over trials.

Types of participants

Preterm infants (< 37 weeks' gestation) receiving enteral feeding of human milk within a hospital setting.

Types of interventions

Human milk with or without additional carbohydrate supplementation.

Types of outcome measures

Primary and secondary outcomes for this review were aligned with the outcomes of the Cochrane Review titled "Multi-nutrient fortification of human milk for preterm infants" ([Brown 2016](#)).

Primary outcomes

- Growth: weight, length, head circumference, skinfold thickness, body mass index, and measures of body composition (lean/fat mass) and growth restriction (proportion of infants who remain < 10th percentile for the index population distribution for weight, length, or head circumference)
 - Researchers assessed growth parameters from birth to hospital discharge, at or after two years' corrected age, during adolescence, and during adulthood
- Neurodevelopmental outcomes
 - Neurodevelopmental outcomes after 12 months post term included neurological evaluations, developmental scores, and classifications of disability, including auditory and visual disability. We defined neurodevelopmental impairment as the presence of one or more of the following: non-ambulant cerebral palsy, developmental quotient greater than two standard deviations below the population mean, blindness (visual acuity < 6/60), or deafness (any hearing impairment requiring or unimproved by amplification)

Secondary outcomes

- Duration of hospital admission
- Feeding intolerance that results in cessation of or reduction in enteral feeding
- Necrotising enterocolitis (NEC)
- Hyperglycaemia

- Diarrhoea
- Gastrointestinal disturbance
- Long-term measures of cardio-metabolic health such as insulin resistance, obesity, diabetes, and hypertension

Search methods for identification of studies

We used the criteria and standard methods of Cochrane and Cochrane Neonatal (see the Cochrane Neonatal [search strategy](#) for specialised register).

Electronic searches

We conducted a comprehensive search including: Cochrane Central Register of Controlled Trials (CENTRAL 2019, Issue 8) in the Cochrane Library and MEDLINE via PubMed (2018 to 22 August 2019). We have included the search strategies for each database in [Appendix 1](#). We did not apply language restrictions.

We searched clinical trial registries for ongoing or recently completed trials (ISRCTN Registry). The World Health Organization's International Clinical Trials Registry Platform (ICTRP) (www.who.int/ictip/search/en/) and the U.S. National Library of Medicine's ClinicalTrials.gov (clinicaltrials.gov) were searched via Cochrane CENTRAL.

This search updates the searches conducted for previous versions of the review ([Amissah 2018](#), [Kuschel 1999](#)).

Searching other resources

We searched the reference lists of articles included in this review to identify additional relevant articles. We did not search any additional conference proceedings.

Data collection and analysis

We used the guidelines and standardised methods of Cochrane and its Neonatal Review Group to assess the methodological quality of the included trial. Two review authors (EA and JH) independently extracted data, compared data, and resolved differences. We used the standard method of Cochrane Neonatal to synthesise data and expressed results as relative risk and weighted mean difference.

Selection of studies

For the 2018 update, review authors carried out the study selection process independently as follows: two review authors (EA and JB) independently screened the titles and abstracts of records identified by the searches. We resolved conflicts through discussion or by consultation with a third review author (JH). We retrieved the full text of all potentially relevant articles and linked together reports of the same study. Two review authors (EA and JB) independently assessed full-text articles for inclusion or exclusion using the review eligibility criteria and resolved conflicts by discussion. We used Covidence during the study selection and data collection process.

For the 2020 update, Cochrane Neonatal screened the titles and abstracts identified by the search independently and in duplicate in consultation with a review author (JH).

Data extraction and management

We developed a data extraction form before gathering data to enable two review authors (EA and JH) to independently extract

information from study reports. We extracted data such as source details, study eligibility, study design, participant characteristics, intervention and control details, and outcomes. We resolved conflicts in the data extraction and management process by discussion. We then exported the data into Cochrane's statistical software, Review Manager 3 (RevMan 2014).

Assessment of risk of bias in included studies

Two review authors (EA and JH) independently assessed the risk of bias (low, high, or unclear) of all included trials using the Cochrane 'Risk of bias' tool for the following domains (Higgins 2017).

- Sequence generation (selection bias).
- Allocation concealment (selection bias).
- Blinding of participants and personnel (performance bias).
- Blinding of outcome assessment (detection bias).
- Incomplete outcome data (attrition bias).
- Selective reporting (reporting bias).
- Any other bias.

We resolved disagreements by discussion. See [Appendix 2](#) for a detailed description of risk of bias for each domain. We contacted the primary author of the included trial for confirmation that the two publications describe the same trial, and for clarification of study methods, specifically, methods related to randomisation and blinding during the trial.

Measures of treatment effect

We used the numbers of events in control and intervention groups of each study to calculate risk ratios (RRs) with 95% confidence intervals (CIs) for dichotomous data. We planned to calculate mean differences (MDs) between treatment groups when outcomes were measured in the same way for continuous data. However, this was not possible because data for our pre-defined outcomes were scarce. Trial investigators measured duration of hospital stay using median and range values. We opted to report their findings narratively rather than convert values to mean and standard deviation because of the skewness of the data. We did not need to use standardised mean differences (SMDs) in this update as there was only one included trial. We reported 95% CIs for all outcomes. We did not calculate number needed to treat for an additional beneficial outcome or number needed to treat for an additional harmful outcome because data were insufficient.

Unit of analysis issues

We did not identify any unit of analysis issues.

Dealing with missing data

We noted levels of attrition. We carried out analyses on an intention-to-treat basis, when possible, for all outcomes. We analysed all participants, when possible, in the treatment group to which they were randomised, regardless of the treatment received. We did not contact the primary author regarding missing data. We were unable to conduct sensitivity analyses and were unable to address the potential impact of missing data on findings of the review because data were insufficient.

Assessment of heterogeneity

We planned to assess whether the clinical and methodological characteristics of included studies were sufficiently similar for meta-analysis to provide a clinically meaningful summary. We planned to do this by assessing statistical heterogeneity using the Chi² test and the I² statistic. An I² measurement greater than 50% and a low P value (< 0.10) in the Chi² test for heterogeneity were taken to indicate moderate to high heterogeneity. When we detected moderate to high heterogeneity, we planned to explore possible explanations through sensitivity and/or subgroup analyses. We planned to take statistical heterogeneity into account when interpreting trial results, especially if we noted any variation in the direction of effect. We were unable to perform any of these assessments, as we included only one trial.

Assessment of reporting biases

Some types of reporting bias (e.g. publication bias, multiple publication bias, language bias) reduce the likelihood that all studies eligible for a review will be retrieved. If all eligible studies are not retrieved, the review may be biased. We aimed to conduct a comprehensive search for eligible studies and were alert for duplication of data. We were unable to assess publication bias, as we found insufficient studies for any of the outcomes (10 or more trials required).

Data synthesis

We planned to use the GRADE approach, as outlined in the [GRADE Handbook](#) (Schünemann 2013), to assess the quality of evidence for the following clinically relevant outcomes: growth, neurodevelopment, duration of hospital admission, feeding intolerance that results in cessation or reduction in enteral feeding, and NEC. However, because of lack of data, we could assess quality using GRADE only for feeding intolerance and NEC.

Two review authors (EA and JB) independently assessed the quality of the evidence for each of the outcomes above. We considered evidence from RCTs as high quality but downgraded the evidence one level for serious (or two levels for very serious) limitations based upon the following: design (risk of bias), consistency across studies, directness of evidence, precision of estimates, and presence of publication bias. We used the [GRADEpro GDT](#) Guideline Development Tool to create a 'Summary of findings' table to report the quality of the evidence.

The GRADE approach yields an assessment of the quality of a body of evidence by one of four grades.

- High: we are very confident that the true effect lies close to that of the estimate of the effect.
- Moderate: we are moderately confident in the effect estimate: the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different.
- Low: our confidence in the effect estimate is limited: the true effect may be substantially different from the estimate of the effect.
- Very low: we have very little confidence in the effect estimate: the true effect is likely to be substantially different from the estimate of effect.

Subgroup analysis and investigation of heterogeneity

We planned to perform subgroup and sensitivity analyses if we noted moderate to high heterogeneity. We planned to consider whether an overall summary was meaningful, and if it was, we planned to use a random-effects model to analyse it. We planned to carry out the following subgroup analyses to evaluate differences in outcomes: gestational age subgroups (< 30 vs 30 to < 34 vs 34 to < 37 completed weeks), birth weight subgroups (< 1 kg vs ≥ 1 kg), male versus female sex, and types of carbohydrate supplements (lactose vs other forms). However, data were insufficient for us to conduct any subgroup analyses.

Sensitivity analysis

We planned to conduct sensitivity analysis by examining only trials considered to have low risk of bias for allocation concealment and

randomisation. We were unable to do this, as we included only one trial in this review.

RESULTS

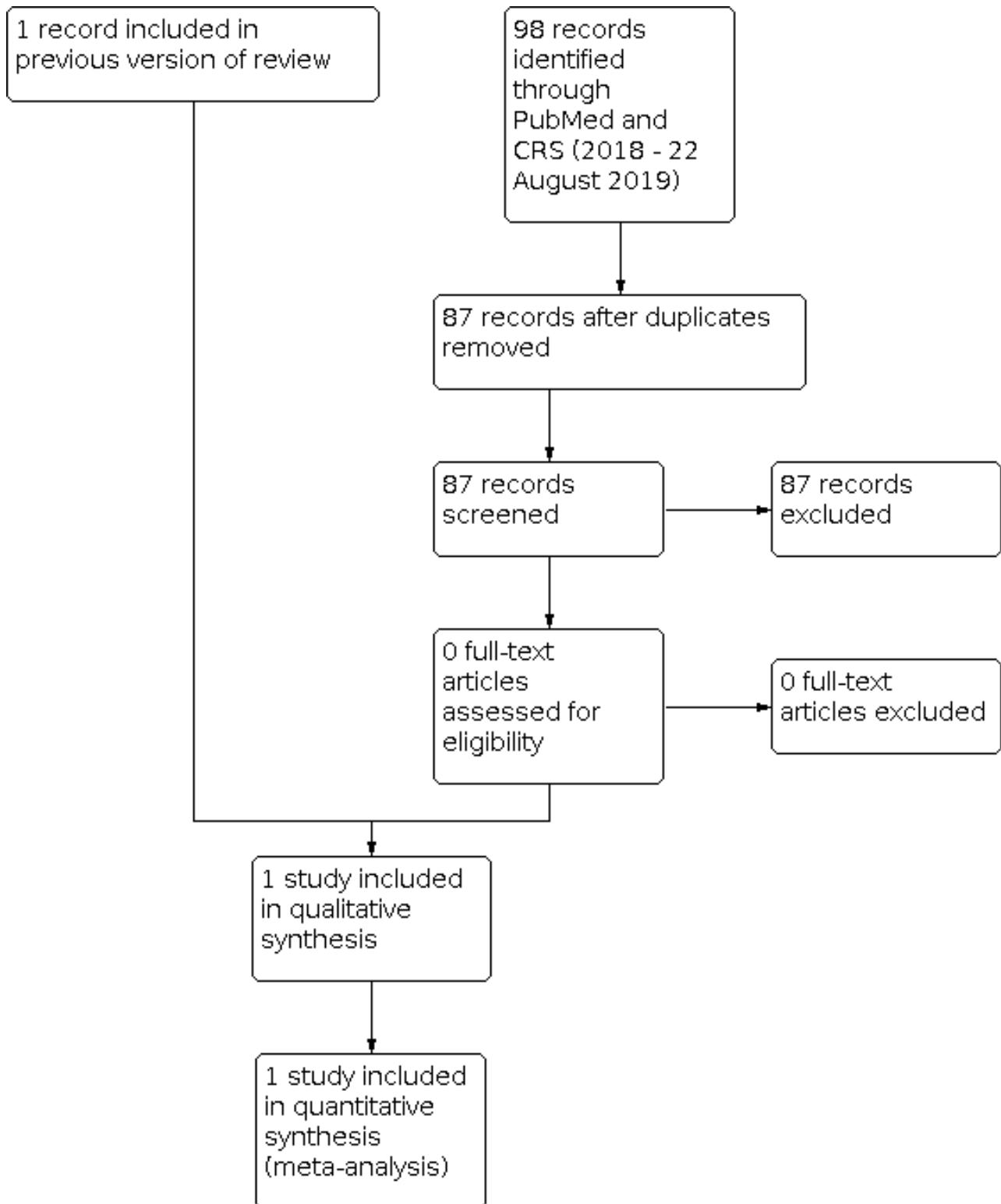
Description of studies

Please refer to the tables sections for study details ([Included studies](#); [Excluded studies](#)).

Results of the search

Using search methods, we identified 548 records after duplicates had been removed. After trial and abstract screening, we excluded 545 records and retrieved three full-text articles for further analysis ([Figure 1](#)). We identified one trial (two publications) as eligible for inclusion in this review ([Armanian 2014](#)).

Figure 1. Study flow diagram: review update



Included studies

One trial published in English met our inclusion criteria (Armanian 2014). This two-armed, single-centre study was carried out at a tertiary neonatal intensive care unit in Iran and included a total of 75 preterm infants. One publication stated that the trial was

quasi-randomised and unblinded (Armanian 2014), but another publication by the same study authors stated that the study was randomised and blinded (Armanian 2016). It was conducted between December 2012 and November 2013 and reported on effects of prebiotic supplementation of human milk on preterm

infants. We have summarised details of the included study in the [Characteristics of included studies](#) table.

Participants

Preterm infants involved in this trial were at ≤ 34 gestational weeks with birth weight ≤ 1500 grams. They had no asphyxia, major congenital anomalies, congenital cyanotic heart disease, gastrointestinal system anomalies, proven sepsis, or infection before the start of the study and were not transferred to other departments. Preterm infants in both intervention and control groups entered the study when their milk intake reached 30 mL/kg/d.

Interventions

Researchers used a non-human short-chain galacto-oligosaccharides/long-chain fructo-oligosaccharides (GOS/ FOS) supplement in a 9:1 mixture. Trialists initially gave incremental doses of the supplement until the infant’s milk intake reached 150 mL/kg/d. However, it was not clear if these initial doses were given separately from breast milk. They then mixed a single dose of 1.5 g/kg/d of supplement with breast milk and fed this to preterm infants in the intervention group for a day or two. We sought clarification from study authors on dosing and mode of administration of the intervention, but we have not yet received a response. Investigators fed only human milk to preterm infants in both intervention and control groups throughout the study. However, it is not clear if the milk was maternal or donor human milk or both.

Comparators

The control group received unsupplemented human milk.

Outcomes

The trial evaluated weight at day 30 but did not report our pre-defined outcome of weight gain in g/kg/d. Study authors also reported duration of hospital stay, feeding intolerance, and NEC, sepsis, intraventricular haemorrhage, patent ductus arteriosus, time to full enteral feeds, and death. They provided no data on short- and long-term growth, body mass index, body composition, or neurodevelopmental and cardio-metabolic outcomes.

Excluded studies

We excluded one trial, for which the intervention was not relevant to our review, as it involved comparing a synbiotic versus no intervention ([Nandhini 2016](#)). We were unable to identify any ongoing trials involving carbohydrate supplementation of human milk in preterm infants. See [Characteristics of excluded studies](#) for details on exclusions.

Risk of bias in included studies

Please see [Characteristics of included studies](#) and the risk of bias graph and summary ([Figure 2](#); [Figure 3](#)) for details.

Figure 2. Risk of bias graph: review authors' judgements about each risk of bias item presented as percentages across all included studies.

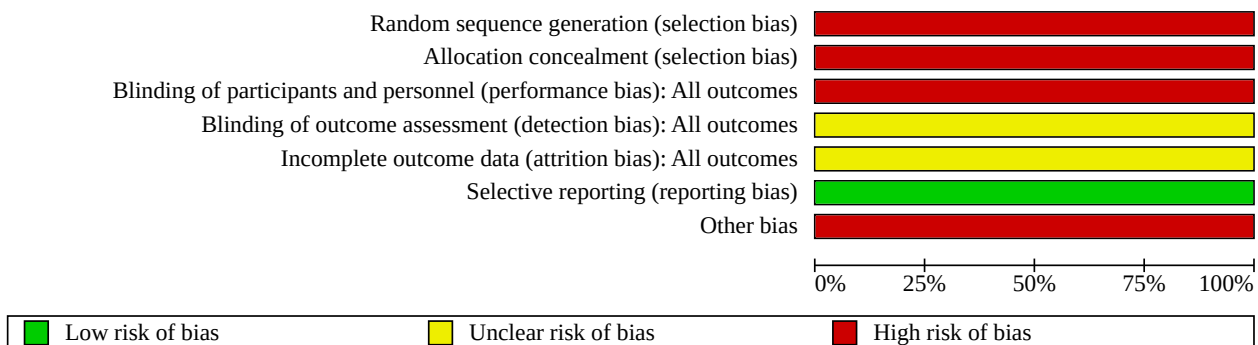
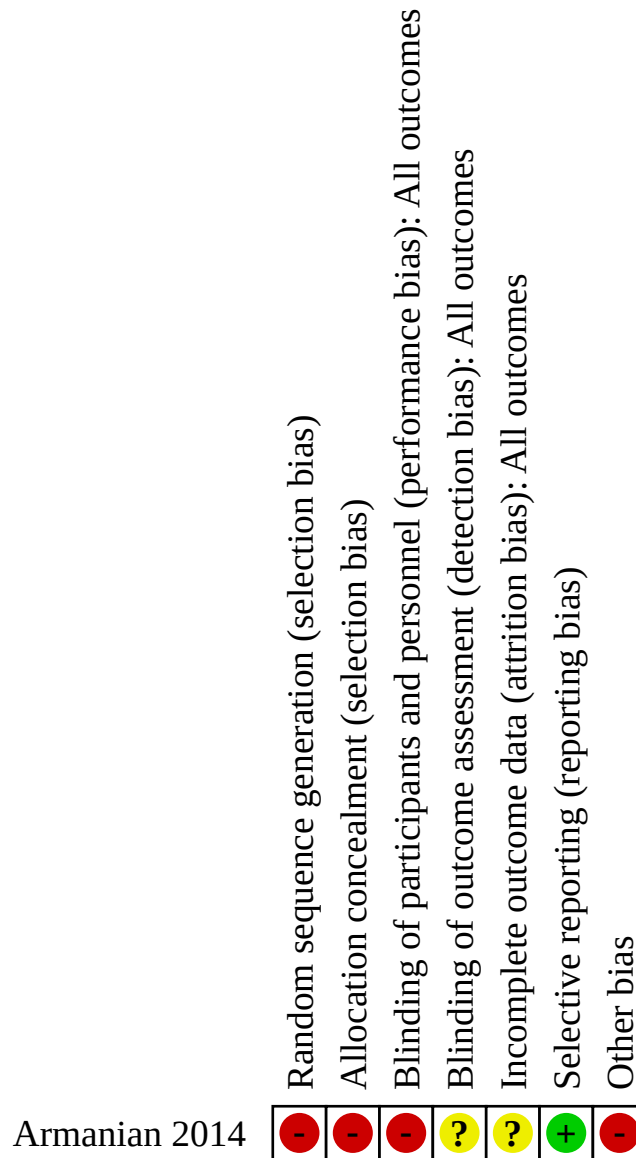


Figure 3. Risk of bias summary: review authors' judgements about each risk of bias item for each included study.



Allocation

Random sequence generation was performed through the use of odd or even file numbers; therefore we judged this as high risk. In addition, although unequal allocation of two controls to one case, as was done in this trial, may be scientifically desirable (Hey 2014), study authors did not report how this was done by using odd and even file numbers. We also judged allocation concealment as high risk because caregivers were not blinded and the allocation sequence was therefore easy to predict.

Blinding

This trial was not blinded, as study authors stated, "care providers were not blinded to an infant's protocol". We judged blinding of allocated participants and personnel as high risk, as knowledge of the allocated intervention was not concealed from clinicians nor participants. We judged blinding of outcome assessors as unclear risk owing to insufficient methodological detail.

Incomplete outcome data

Study authors clearly reported reasons for withdrawals and dropouts and provided the missing numbers for each group. One reason given for missing numbers in each group was transfer to other departments. However, numbers transferred differed between groups, and study authors did not report why infants were transferred, as this potentially could be related to study outcomes. Study authors also reported no comparison of baseline characteristics between included and excluded participants. It is unclear whether researchers used an intention-to-treat approach, as they included four participants lost from both groups in the analysis and excluded the 13 randomised infants who were transferred. We judged attrition bias to be at unclear risk owing to insufficient methodological details.

Selective reporting

We viewed no protocol. However, we included in the [Results](#) section all outcomes mentioned in the [Methods](#) section. We judged this to be of low risk.

Other potential sources of bias

We judged this as high risk owing to the publication of two reports for the same trial that reported different methods and different numbers of included infants. We were unable to reproduce the reported analyses from published data.

Effects of interventions

See: [Summary of findings 1 Carbohydrate supplementation compared to control in preterm infants](#)

1 Carbohydrate supplementation versus control

1.1 Growth/Weight

1.1.1 Weight at day 30

One randomised controlled trial including 75 infants contributed data ([Armanian 2014](#)). Prebiotic supplementation of human milk led to an increase in weight at day 30 compared with unsupplemented human milk (MD 160.4 grams, 95% CI 12.4 to 308.4 grams; one RCT, N = 75 infants; very low-quality evidence). We downgraded the evidence for risk of bias, as methodological information provided was insufficient for judgement of the risk of bias, participants and events were few, and confidence intervals were wide.

1.2 Feeding intolerance

[Armanian 2014](#) contributed data providing no clear evidence that prebiotic supplementation increased the risk of feeding intolerance (RR 0.64, 95% CI 0.36 to 1.15; one RCT, N = 75 infants; very low-quality evidence). We downgraded the evidence for risk of bias as methodological information was insufficient for judgement of risk of bias and participants and events were few.

1.3 Necrotising enterocolitis

One trial reported data on the incidence of necrotising enterocolitis ([Armanian 2014](#)), which showed no evidence of a clear difference in risk between prebiotic-supplemented and unsupplemented groups (RR 0.2, 95% CI 0.02 to 1.3; one RCT, N = 75 infants; very low-quality evidence). We downgraded the quality of evidence for risk of bias as methodological information was insufficient for judgement

of risk of bias, participants and events were few, and confidence intervals were wide.

1.4 Duration of hospital stay

One trial reported on duration of hospital stay ([Armanian 2014](#)), noting that the prebiotic-supplemented group had a shorter hospital stay than the unsupplemented group. The median (range) length of hospital stay was 16 (9 to 45) days (95% CI 15.34 to 24.09) and 25 (11 to 80) days (95% CI 25.52 to 34.39), respectively. We downgraded the evidence for risk of bias to very low quality, as methodological information was insufficient for judgement of risk of bias, patients and events were few, and confidence intervals were wide.

Study authors also reported on sepsis, intraventricular haemorrhage (IVH), patent ductus arteriosus (PDA), time to full feeds, and death, which were not among our pre-defined outcomes. However, they provided no data for any of our other pre-specified primary or secondary outcomes, including long-term growth, body mass index, body composition, and neurodevelopmental and cardio-metabolic outcomes. We were unable to conduct our pre-specified subgroup analysis owing to insufficient data.

DISCUSSION

Summary of main results

We identified one trial for inclusion in this review ([Armanian 2014](#)). In this trial, the prebiotic-supplemented group showed an increase in mean weight at 30 days of age and shorter length of hospital stay compared to the control group. Study authors provided no evidence of a clear difference in risk of feeding intolerance and necrotising enterocolitis (NEC) between intervention and control groups. Except for weight at day 30, no data were available for assessment of effects of carbohydrate supplementation on short- and long-term growth, body mass index (BMI), body composition, and neurodevelopmental and cardio-metabolic outcomes.

Overall completeness and applicability of evidence

The only trial included in this review is limited in applicability because it was conducted among a small sample of preterm infants in Iran. However, the outcomes assessed are common to all preterm infants. This trial shows that prebiotic carbohydrate supplementation of human milk may be feasible in developing countries.

Quality of the evidence

The included trial had high risk of selection and performance bias owing to quasi-randomisation and lack of blinding of caregivers. In addition, two reports of this trial were inconsistent regarding methods and numbers of included infants, and we were unable to reproduce analyses of findings for NEC and weight at day 30, as reported in the publication. The overall quality of evidence for outcomes assessed according to GRADE was very low owing to insufficient methodological details, high risk of bias, small sample size, few events, and variable precision.

Potential biases in the review process

The comprehensive search strategy, use of appropriate search terminology, and lack of language restriction in this review

minimised bias and increased the likelihood of identifying all relevant articles. Additionally, involvement of multiple authors in the review process to independently screen records for inclusion and extract data for analysis limited the introduction of bias into the review. However, there is always potential for publication bias. Unfortunately, we were unable to create funnel plots and evaluate this potential risk because we included only one trial in the review.

Agreements and disagreements with other studies or reviews

To the best of our knowledge, this is the only systematic review covering this topic, and our previous review found no trials eligible for inclusion.

AUTHORS' CONCLUSIONS

Implications for practice

The only included trial shows very low-quality evidence of the effects of prebiotic carbohydrate supplementation of human milk in preterm infants, and provided no data on the short- and long-term health benefits and harms of digestible carbohydrate supplementation of human milk in this population. Therefore, we are unable to make any clinical suggestions on the basis of the single included trial.

Implications for research

Currently, we have found little evidence to support or refute the practice of carbohydrate supplementation of human milk in preterm infants. If further trials seek to examine prebiotic or digestible carbohydrate supplementation done as a component of multi-nutrient fortification, trialists should assess the optimum concentrations of carbohydrate supplements, adverse effects, and short- and long-term growth and health benefits for preterm infants.

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* Indicates the major publication for the study

CHARACTERISTICS OF STUDIES
Characteristics of included studies [ordered by study ID]
Armanian 2014
Study characteristics

| | |
|---------------|---|
| Methods | Quasi-randomized controlled trial, single centre |
| Participants | Inclusion criteria: infants with birth weight \leq 1500 grams, gestational age \leq 34 weeks, who had no formula feeds. Infants were included when their milk feeding volume reached 30 mL/kg/d. Exclusion criteria: infants with asphyxia, major congenital anomalies, congenital cyanotic heart disease, gastrointestinal system anomalies, proven sepsis, or infection immediately before the start of the study, transferred to other departments, refused to participate Setting: NICUs of the Isfahan University of Medical Sciences (Alzahra and Shahid Beheshti Hospital) Timing: December 2012 to November 2013 |
| Interventions | Breast milk supplemented with 9:1 mixture of short-chain galacto-oligosaccharide/long-chain fructo-oligosaccharide (n = 25) vs unsupplemented breast milk (n = 50) Not clear when the intervention ceased |
| Outcomes | Primary outcomes: incidence of suspected NEC Secondary outcomes: milk volumes, feeding intolerance (presence of milk in the stomach 2 hours after a meal, i.e. gastric residue), abdominal distension, postnatal age when full enteral feeding was attained, death, length of hospital stay, weight at day 30, associated patent ductus arteriosus, and intraventricular haemorrhage |
| Notes | Conflicts of interest: none declared, but Nutricia MMP, Mashad, Iran is acknowledged. Source of support: This paper is derived from a residency thesis (No. 392237) at Isfahan University of Medical Sciences. No details of source of funding are provided. Study authors published 2 reports of the same trial but reported different methods regarding blinding and randomisation, and included different numbers of infants. The first publication reported quasi-randomisation and lack of blinding of caregivers, and the second publication reported computer-generated randomisation and blinding of the investigator and nurse. Study authors confirmed via email that publications described the same trial, but they have not yet clarified the different methods reported. We chose to include details from the first publication, as it involved a larger sample size and assessed outcomes relevant to this review. |

Risk of bias

Armanian 2014 (Continued)

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | High risk | Infants were randomised based on files ending in odd or even numbers. It is not clear how the 2:1 allocation was achieved. |
| Allocation concealment (selection bias) | High risk | Randomisation was done by an independent employee, but allocation was not blinded and could be predicted. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Study authors stated, "care providers were not blinded to an infant's protocol". |
| Blinding of outcome assessment (detection bias) All outcomes | Unclear risk | Suspected NEC diagnosis was clinical and therefore was potentially subject to bias. However, it is unclear whether outcome assessors were among the care providers who were not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Unclear risk | Losses to follow-up were disclosed as follows: of 131 eligible infants, 24 were excluded for having major congenital anomalies, gastrointestinal system anomalies, asphyxia, or sepsis, and parents of 19 infants refused to participate or were transferred to other wards. Of 88 infants randomised, 13 (9 in the prebiotic group and 4 in the control group) were transferred to other departments. The 13 excluded after randomisation appear to be attributable to reasonable attrition. However, the reasons for transfer, which could be related to outcomes assessed, were not stated. It also was not clear whether study authors compared baseline characteristics between participants included and excluded from the study. One loss to follow-up and 1 death occurred in each group, but these appear to be included in the analysis, although the 13 transferred to other departments were excluded, making it difficult to determine whether an intention-to-treat approach was used. |
| Selective reporting (reporting bias) | Low risk | The study protocol was not available, but researchers reported results for all outcomes listed in the methods. |
| Other bias | High risk | Study authors published 2 reports of the same trial that described different methods and different numbers of included infants. We were unable to reproduce their analyses from the published data. |

NEC: necrotising enterocolitis.

NICU: neonatal intensive care unit.

Characteristics of excluded studies [ordered by study ID]

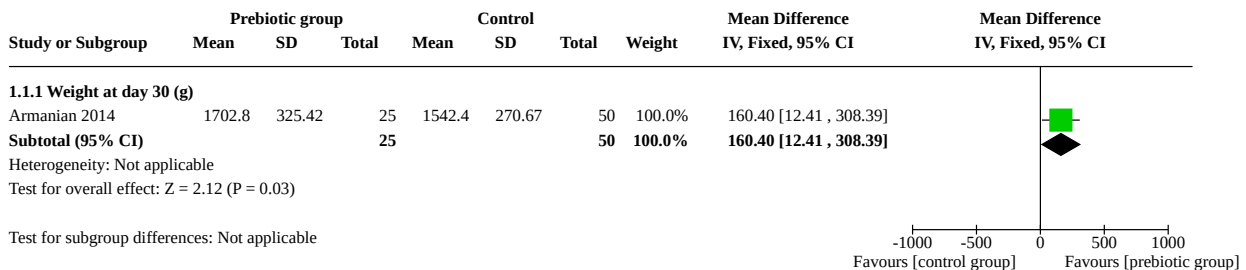
| Study | Reason for exclusion |
|-------------------------------|---|
| Nandhini 2016 | Wrong intervention - synbiotic vs no intervention |

DATA AND ANALYSES

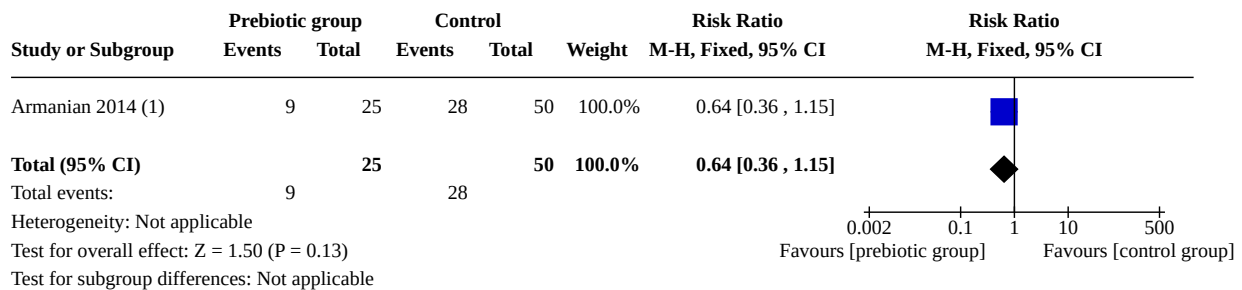
Comparison 1. Carbohydrate supplementation versus control

| Outcome or subgroup title | No. of studies | No. of participants | Statistical method | Effect size |
|-------------------------------|----------------|---------------------|-------------------------------------|------------------------|
| 1.1 Growth/Weight | 1 | | Mean Difference (IV, Fixed, 95% CI) | Subtotals only |
| 1.1.1 Weight at day 30 (g) | 1 | 75 | Mean Difference (IV, Fixed, 95% CI) | 160.40 [12.41, 308.39] |
| 1.2 Feeding intolerance | 1 | 75 | Risk Ratio (M-H, Fixed, 95% CI) | 0.64 [0.36, 1.15] |
| 1.3 Necrotising enterocolitis | 1 | 75 | Risk Ratio (M-H, Fixed, 95% CI) | 0.18 [0.02, 1.33] |

Analysis 1.1. Comparison 1: Carbohydrate supplementation versus control, Outcome 1: Growth/Weight



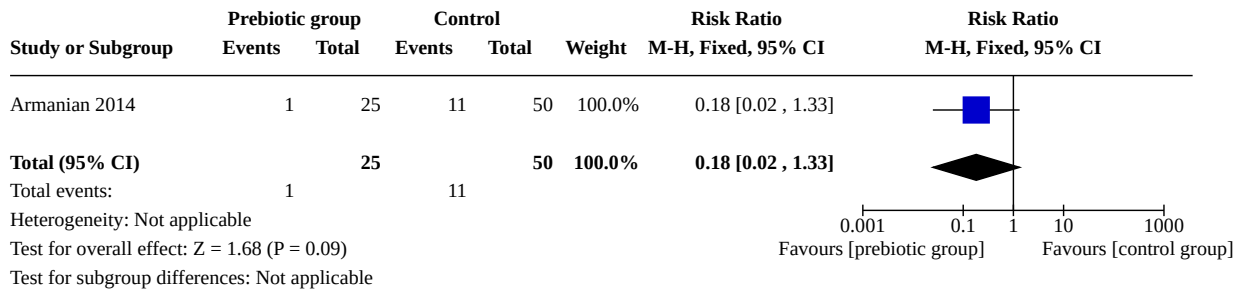
Analysis 1.2. Comparison 1: Carbohydrate supplementation versus control, Outcome 2: Feeding intolerance



Footnotes

(1) The authors defined feeding intolerance as: gastric residue, i.e. the presence of milk in the stomach 2 h after completion of a feeding. They also assessed th

Analysis 1.3. Comparison 1: Carbohydrate supplementation versus control, Outcome 3: Necrotising enterocolitis



APPENDICES

Appendix 1. Search methods 2020 update

MEDLINE via PubMed

((Dietary Carbohydrates[Mesh] OR Carbohydrate*[tiab] OR Glucans[Mesh] OR Sucrose[Mesh] OR glucan*[tiab] OR monosaccharide*[tiab] OR disaccharide*[tiab] OR oligosaccharide*[tiab] OR polysaccharide*[tiab] OR polycose[tiab] OR lactose[tiab] OR sucrose[tiab] OR corn syrup[tiab]) AND (((Milk, Human[MeSH] OR breastmilk*[tiab])) OR (((human[tiab] OR breast[tiab] OR expressed[tiab] OR mother*[tiab] OR maternal[tiab] OR donor*[tiab])) AND milk*[tiab]))) AND (((infant, newborn[MeSH] OR newborn OR neonate OR neonatal OR premature OR low birth weight OR VLBW OR LBW or infan* or neonat*) AND (randomized controlled trial [pt] OR controlled clinical trial [pt] OR randomized [tiab] OR placebo [tiab] OR drug therapy [sh] OR randomly [tiab] OR trial [tiab] OR groups [tiab]) NOT (animals [mh] NOT humans [mh])))

Cochrane Library

| | |
|----|---|
| 1 | MESH DESCRIPTOR Dietary Carbohydrates EXPLODE ALL AND CENTRAL:TARGET |
| 2 | MESH DESCRIPTOR Sucrose EXPLODE ALL AND CENTRAL:TARGET |
| 3 | MESH DESCRIPTOR Glucans EXPLODE ALL AND CENTRAL:TARGET |
| 4 | (Carbohydrate* or glucan* or monosaccharide* or disaccharide* or oligosaccharide* or polysaccharide* or polycose or lactose or sucrose or corn syrup):TI,AB,KW AND CENTRAL:TARGET |
| 5 | #1 OR #2 OR #3 OR #4 |
| 6 | MESH DESCRIPTOR Milk, Human EXPLODE ALL AND CENTRAL:TARGET |
| 7 | ((human OR breast OR expressed) NEAR2 milk*):TI,AB,KW AND CENTRAL:TARGET |
| 8 | ((mother* or maternal or donor*) NEAR2 milk*): TI,AB,KW AND CENTRAL:TARGET |
| 9 | #6 OR #7 OR #8 AND CENTRAL:TARGET |
| 10 | (infan* or newborn or neonat* or premature or preterm or very low birth weight or low birth weight or VLBW or LBW) AND CENTRAL:TARGET |
| 11 | #5 AND #9 AND #10 |

Appendix 2. Risk of bias tool

We used the standard methods of Cochrane and Cochrane Neonatal to assess the methodological quality (to meet the validity criteria) of trials. For the included trial, we sought information from trial investigators regarding the intervention, the method of randomisation, and blinding, but to date we have received no response. We assessed each criterion as low, high, or unclear risk. Two review authors separately assessed the study. We resolved any disagreement by discussion. We added this information to the table 'Characteristics of included studies'. We evaluated the following issues and entered the findings into the risk of bias table.

1. Sequence generation (checking for possible selection bias). Was the allocation sequence adequately generated?

For the included study, we categorised the method used to generate the allocation sequence as:

- low risk (any truly random process, e.g. random number table; computer random number generator);
- high risk (any non-random process, e.g. odd or even date of birth; hospital or clinic record number); or
- unclear risk.

2. Allocation concealment (checking for possible selection bias). Was allocation adequately concealed?

For the included study, we categorised the method used to conceal the allocation sequence as:

- low risk (e.g. telephone or central randomisation; consecutively numbered sealed opaque envelopes);
- high risk (e.g. open random allocation; unsealed or non-opaque envelopes, alternation; date of birth); or
- unclear risk.

3. Blinding of participants and personnel (checking for possible performance bias). Was knowledge of the allocated intervention adequately prevented during the study?

For the included study, we categorised the methods used to blind study participants and personnel from knowledge of which intervention a participant received. Blinding was assessed separately for different outcomes or classes of outcomes. We categorised the methods as:

- low risk, high risk, or unclear risk for participants; and
- low risk, high risk, or unclear risk for personnel.

4. Blinding of outcome assessment (checking for possible detection bias). Was knowledge of the allocated intervention adequately prevented at the time of outcome assessment?

For the included study, we categorised the methods used to blind outcome assessment. Blinding was assessed separately for different outcomes or classes of outcomes. We categorised the methods as:

- low risk for outcome assessors;
- high risk for outcome assessors; or
- unclear risk for outcome assessors.

5. Incomplete outcome data (checking for possible attrition bias through withdrawals, dropouts, protocol deviations). Were incomplete outcome data adequately addressed?

For the included study and for each outcome, we described the completeness of data including attrition and exclusions from the analysis. We noted whether attrition and exclusions were reported, numbers included in the analysis at each stage (compared with the total randomised participants), reasons for attrition or exclusion when reported, and whether missing data were balanced across groups or were related to outcomes. When sufficient information was reported or supplied by the trial authors, we re-included missing data in the analyses. We categorised the methods as:

- low risk (< 20% missing data);
- high risk (\geq 20% missing data); or
- unclear risk.

6. Selective reporting bias. Are reports of the study free of suggestion of selective outcome reporting?

For the included study, we described how we investigated the possibility of selective outcome reporting bias and what we found. We assessed the methods as:

- low risk (when it is clear that all of the study's pre-specified outcomes and all expected outcomes of interest to the review have been reported);

- high risk (when not all of the study's pre-specified outcomes have been reported; one or more reported primary outcomes were not pre-specified outcomes of interest and are reported incompletely and so cannot be used; study fails to include results of a key outcome that would have been expected to have been reported); or
- unclear risk.

7. Other sources of bias. Was the study apparently free of other problems that could put it at high risk of bias?

For the included study, we described any important concerns we had about other possible sources of bias (e.g. whether a potential source of bias was related to the specific study design, whether the trial was stopped early owing to some data-dependent process). We assessed whether the study was free of other problems that could put it at risk of bias as:

- low risk;
- high risk; or
- unclear risk.

If needed, we planned to explore the impact of the level of bias by undertaking sensitivity analyses.

Appendix 3. Previous search methods

MEDLINE via PubMed

```
((Dietary Carbohydrates[Mesh] OR Carbohydrate*[tiab] OR Glucans[Mesh] OR Sucrose[Mesh] OR glucan*[tiab] OR monosaccharide*[tiab] OR disaccharide*[tiab] OR oligosaccharide*[tiab] OR polysaccharide*[tiab] OR polycose[tiab] OR lactose[tiab] OR sucrose[tiab] OR corn syrup[tiab]) AND (((Milk, Human[MeSH] OR breastmilk*[tiab])) OR (((human[tiab] OR breast[tiab] OR expressed[tiab] OR mother*[tiab] OR maternal[tiab] OR donor*[tiab])) AND milk*[tiab]))) AND (((infant, newborn[MeSH] OR newborn OR neonate OR neonatal OR premature OR low birth weight OR VLBW OR LBW or infan* or neonat*) AND (randomized controlled trial [pt] OR controlled clinical trial [pt] OR randomized [tiab] OR placebo [tiab] OR drug therapy [sh] OR randomly [tiab] OR trial [tiab] OR groups [tiab]) NOT (animals [mh] NOT humans [mh])))
```

Embase

| | |
|----|--|
| 1 | exp carbohydrate diet/ |
| 2 | exp glucan/ |
| 3 | exp sucrose/ |
| 4 | (Carbohydrate* or glucan* or monosaccharide* or disaccharide* or oligosaccharide* or polysaccharide* or polycose or lactose or sucrose or corn syrup).ti,ab. |
| 5 | 1 or 2 or 3 or 4 |
| 6 | exp breast milk/ |
| 7 | ((human or breast or expressed) adj milk\$).ti,ab. |
| 8 | ((mother\$ or maternal or donor\$) adj milk\$).ti,ab. |
| 9 | 6 or 7 or 8 |
| 10 | (infan* or newborn or neonat* or premature or very low birth weight or low birth weight or VLBW or LBW).mp. |
| 11 | exp infant/ |
| 12 | 10 or 11 |
| 13 | (human not animal).mp. |

(Continued)

| | |
|----|--|
| 14 | (randomized controlled trial or controlled clinical trial or randomized or placebo or clinical trials as topic or randomly or trial or clinical trial).mp. |
| 15 | 12 and 13 and 14 |
| 16 | 5 and 9 and 15 |

CINAHL

| | |
|-----|--|
| S1 | (MH "Dietary Carbohydrates+") |
| S2 | (MH "Glucans+") |
| S3 | (MH "Sucrose+") |
| S4 | Carbohydrate* OR glucan* OR monosaccharide* OR disaccharide* OR oligosaccharide* OR polysaccharide* OR polycose OR lactose OR sucrose OR corn syrup |
| S5 | S1 OR S2 OR S3 OR S4 |
| S6 | (human OR breast OR expressed OR mother* OR maternal OR donor*) AND milk* |
| S7 | breastmilk* |
| S8 | S6 OR S7 |
| S9 | (infan* OR newborn OR neonat* OR premature OR low birth weight OR VLBW OR LBW) AND (randomized controlled trial OR controlled clinical trial OR randomized OR placebo OR clinical trials as topic OR randomly OR trial OR PT clinical trial) |
| S10 | S5 AND S8 AND S9 |

Cochrane Library

| | |
|---|---|
| 1 | MESH DESCRIPTOR Dietary Carbohydrates EXPLODE ALL AND CENTRAL:TARGET |
| 2 | MESH DESCRIPTOR Sucrose EXPLODE ALL AND CENTRAL:TARGET |
| 3 | MESH DESCRIPTOR Glucans EXPLODE ALL AND CENTRAL:TARGET |
| 4 | (Carbohydrate* or glucan* or monosaccharide* or disaccharide* or oligosaccharide* or polysaccharide* or polycose or lactose or sucrose or corn syrup):TI,AB,KW AND CENTRAL:TARGET |
| 5 | #1 OR #2 OR #3 OR #4 |
| 6 | MESH DESCRIPTOR Milk, Human EXPLODE ALL AND CENTRAL:TARGET |
| 7 | ((human OR breast OR expressed) NEAR2 milk*):TI,AB,KW AND CENTRAL:TARGET |

(Continued)

| | |
|----|---|
| 8 | ((mother* or maternal or donor*) NEAR2 milk*): TI,AB,KW AND CENTRAL:TARGET |
| 9 | #6 OR #7 OR #8 AND CENTRAL:TARGET |
| 10 | (infan* or newborn or neonat* or premature or preterm or very low birth weight or low birth weight or VLBW or LBW) AND CENTRAL:TARGET |
| 11 | #5 AND #9 AND #10 |

WHAT'S NEW

| Date | Event | Description |
|-----------------|---------|---|
| 20 October 2020 | Amended | <p>Abstract results amended to better align with summary of findings table.</p> <p>In future, this review will no longer be updated, as it will be superseded by Cochrane Review Multi-nutrient fortification of human milk for preterm infants (Brown 2020).</p> |

HISTORY

Protocol first published: Issue 3, 1997

Review first published: Issue 2, 1999

| Date | Event | Description |
|------------------|--|--|
| 29 June 2020 | New search has been performed | <ul style="list-style-type: none"> We updated the search for eligible studies to August 2019. We found no new randomised controlled trials. |
| 29 June 2020 | New citation required but conclusions have not changed | A supplementary search was carried out in August 2019. No new studies were identified. The main conclusions of the original review remain unchanged. |
| 26 February 2018 | New citation required and conclusions have changed | We made changes to the main conclusions of the original review. |
| 26 February 2018 | New search has been performed | <ul style="list-style-type: none"> We updated the search for eligible studies to February 2018. We found one randomised controlled trial (two publications) that was eligible for inclusion in this review (Armanian 2014). We included body mass index and measures of body composition in the growth parameters of the primary outcome. We also included new secondary outcome measures: long-term measures of cardio-metabolic health (such as insulin resistance, obesity, diabetes, and hypertension). These outcomes were aligned with those of the Cochrane Review titled "Multi-nutrient fortification of human milk for preterm infants" (Brown 2016). We applied the Grading of Recommendations Assessment, Development and Evaluation (GRADE) approach to assess the quality of evidence. |

| Date | Event | Description |
|------------------|--|--|
| 10 May 2002 | New search has been performed | This is an update of the existing review: "Carbohydrate supplementation of human milk to promote growth in preterm infants" (Cochrane Library; 1999, Issue 2). No new trials were identified in the search done in April 2002; as a result, we made no substantive changes to the review. We made no change to the conclusion that the addition of carbohydrate supplements to human milk in preterm infants has not been studied sufficiently to inform recommendations for practice. |
| 29 November 1998 | New citation required and conclusions have changed | Substantive amendments were made. |

CONTRIBUTIONS OF AUTHORS

For the 2018 update:

Emma Amissh assessed study eligibility, extracted data, assessed risk of bias of included studies, analysed data, interpreted results of the analysis, and updated the review. She wrote all drafts and addressed comments from co-authors.

Julie Brown assessed study eligibility, reviewed data extracted, assessed risk of bias of included studies, assisted in interpretation of analyses, and provided comments on drafts.

Jane Harding extracted data, answered queries on trial eligibility, performed risk of bias assessments of included studies, assisted in interpretation of analyses, and provided comments on all drafts of the review.

All review authors read and approved the final version of the review.

For the 2020 update:

Only the search was updated, with no new trials found. The search was screened by Cochrane Neonatal in consultation with Jane Harding. All previous author contributions remain the same, as the text of the review remains unchanged.

DECLARATIONS OF INTEREST

EA receives a scholarship in the form of a stipend from the University of Auckland as a PhD student.

JB is currently employed by a medical writing company. The preparation of this review took place prior to this employment and her current work is not related to the topic of this review.

JH has partial salary support from research grants from the Health Research Council of New Zealand. The Council has no role in the production of this review. An undirected research grant is pending from Biomed Ltd, Auckland, New Zealand. This company makes dextrose gel.

Core editorial and administrative support for the 2020 update of this review has been provided by a grant from The Gerber Foundation. The Gerber Foundation is a separately endowed, private foundation, independent from the Gerber Products Company. The grantor has no input on the content of the review or the editorial process (see [Sources of support](#)).

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External sources

- Liggins Institute, University of Auckland, New Zealand

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- National Institute for Health Research, UK

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- Vermont Oxford Network, USA

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- The Gerber Foundation, USA

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DIFFERENCES BETWEEN PROTOCOL AND REVIEW

For the 2018 update:

The original protocol was published in 1997, and formed the basis of the last version of this review, written in 1999. The 2018 update aligned review outcomes with those of the Cochrane Review titled "Multi-nutrient fortification of human milk for preterm infants" (Brown 2016). We added body mass index and measures of body composition to growth parameters of the primary outcome. We also included new secondary outcome measures such as long-term measures of cardio-metabolic health (e.g. insulin resistance, obesity, diabetes, hypertension). We added 'Summary of findings' tables and GRADE recommendations, which were not included in the original protocol.

For the 2020 update:

As of July 2019, Cochrane Neonatal no longer searches Embase for its reviews. RCTs and controlled clinical trials (CCTs) from Embase are added to the Cochrane Central Register of Controlled Trials (CENTRAL) via a robust process (see [How CENTRAL is created](#)). Cochrane Neonatal has validated their searches to ensure that relevant Embase records are found while searching CENTRAL.

Also starting in July 2019, Cochrane Neonatal no longer searches for RCTs and CCTs on the following platforms: ClinicalTrials.gov or from The World Health Organization's International Clinical Trials Registry Platform (ICTRP), as records from both platforms are added to CENTRAL on a monthly basis (see [How CENTRAL is created](#)). Comprehensive search strategies are executed in CENTRAL to retrieve relevant records. The ISRCTN (at www.isrctn.com/, formerly Controlled-trials.com), is searched separately.

We did not search CINAHL for this update. The 2018 search methods are listed in [Appendix 3](#).

INDEX TERMS

Medical Subject Headings (MeSH)

Body Weight; Dietary Carbohydrates [*administration & dosage]; *Dietary Supplements; Enterocolitis, Necrotizing [epidemiology]; Food Intolerance [epidemiology]; *Infant Nutritional Physiological Phenomena; Infant, Premature [*growth & development]; Length of Stay; *Milk, Human [chemistry]; Oligosaccharides [administration & dosage]; *Prebiotics

MeSH check words

Humans; Infant, Newborn