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Effects of Nutritional Interventions during Pregnancy on Infant and Child Cognitive Outcomes: A Systematic Review and Meta-Analysis

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Abstract: Background: Epidemiological studies have demonstrated that folate, iodine and iron intake during pregnancy impacts on foetal brain development and cognitive function. However, in human studies, the relationship with other dietary nutrients is less clear. Objective: This systematic review aims to critically appraise the current literature and meta-analyses results from nutritional interventions during pregnancy that aimed to optimise infant and child cognitive outcomes. Design: Ten electronic databases were searched for articles published up to August 2017. The search was limited to articles published in English. Randomised controlled trials (RCTs) testing the impact of any nutritional intervention (dietary counselling, education, nutrient supplementation, fortified foods and/or foods) during pregnancy on cognitive outcomes of children (<10 years old). Two independent reviewers assessed study eligibility and quality using the American Dietetic Association quality criteria checklist for primary research. Standardised mean differences were used for nine cognitive domains to measure effects for meta-analyses. Results: A total of 34 RCTs were included (21 studies included children aged less than 35 months, 10 studies included children aged 36–60 months and 3 studies included children aged 61–119 months). The types of nutritional interventions included nutrient supplements, whole foods, fortified foods and nutrition education. The following nine cognition outcomes: attention, behaviour, crystallised intelligence, fluid intelligence, global cognition, memory, motor skills, visual processing, and problem solving were not significantly impacted by nutritional interventions, although 65% of studies conducted post-hoc data analyses and were likely to be underpowered. Although, long chain polyunsaturated fatty acids (LCPUFA) supplementation was associated with a marginal increase in crystallised intelligence (Effect size (ES): 0.25; 95% confidence interval (95% CI): −0.04, 0.53), the effect was not statistically significant (p = 0.09), with significant study heterogeneity (p = 0.00). Conclusions: LCPUFA supplementation may be associated with an...
improvement in child crystallised intelligence, however further research is warranted. The remaining eight cognition domains were not significantly impacted by maternal nutritional interventions.

**Keywords:** behaviour; child; cognition; cognitive function; infant; nutrition; pregnancy; supplement

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1. Introduction

Adequate nutrition during the prenatal period and early years of life is essential for brain development and cognitive function. Approximately 28 days after conception the neural plate folds and fuses, forming the neural tube, which gives rise to the development of the foetal brain [1]. Adequate folate from the maternal diet during this period is essential for the formation of the neural tube, a deficiency in this nutrient can adversely affect brain development, resulting in neural tube defects, spina bifida and encephalocoele [2–6]. Following the formation of the neural tube, neurodevelopmental processes including, cell proliferation and migration occur during gestation, while neurogenesis, synaptic formation and myelination continue until late adolescence [7]. Iodine is necessary for neural cell migration and differentiation, synaptogenesis and myelination [8], while dietary iron is necessary for neurogenesis and dopamine production [9–11]. Deficiency in these nutrients are known to compromise brain development and cause significant cognitive impairment in the offspring [8,12]. The importance of an adequate intake of folate, iodine and iron during pregnancy for foetal brain development has been well explored; however, the developmental role of other dietary nutrients (e.g., zinc, long-chain polyunsaturated fatty acids (LCPUFAs)) remains unclear from human studies.

The link between early-life nutrition and child cognitive function is difficult to establish because the brain is a heterogeneous organ consisting of multiple anatomic regions (e.g., the hippocampus, striatum, cortex) and neurodevelopmental processes (e.g., synaptogenesis, myelination) with distinct developmental trajectories that span and peak at different times [13,14]. For example, myelination commences at 12–14th week of gestation and occurs at a peak rate during the first two years of life, but continues until adulthood [15]. The prefrontal cortex, which has a prominent role in higher cognitive control, develops in growth spurts during the first two years of life, once again between seven and nine years of age and also at approximately 15 years of age [16,17]. The effect of nutrition on cognitive function depends on the nutrients involved and the maturation stage of the brain. These nutrient effects on brain structure and function may occur in the short-term, while others may not be apparent until full maturation is reached [18].

Optimising brain development and cognitive function holds major long-term consequences for individuals and societies. Cognitive function affects academic performance and the level of education attained [19]. Higher educational attainment is related to a lower burden of disease, due to greater health care access and healthy lifestyle behaviours [20]. Suboptimal cognition during childhood has been associated with an increased risk of adolescent delinquency [21–23] and adult violent criminality [21,24–26]. Incarceration can cause permanent cognitive damage, which holds major implications for future education, employment and health [27]. Strategies that aim to maximise the cognitive performance of children are therefore important for public health policy and nutrition during pregnancy.

This systematic review and meta-analysis aimed to determine whether nutritional intervention/s during pregnancy alter cognitive outcomes during infancy and late childhood. Larson et al. [28] analysed the impact of maternal nutritional interventions on child cognition, however this review focused on short-term cognitive outcomes in children under the age of two years in developing countries. Therefore, a systematic review that considers the impact of maternal nutritional interventions on long-term cognitive outcomes of children living in developed and developing countries is warranted to provide the most comprehensive evaluation of the literature in this area.
2. Methods

The review protocol was developed using The Cochrane Handbook for Systematic Reviews of Interventions. The Preferred Reporting Items for Systematic Review Meta-Analyses (PRISMA) [29] was applied. Similar methods have been followed previously by Gresham et al. [30]. The review protocol can be found in Supplementary Figure S1.

2.1. Search Strategy

A search strategy was devised with the assistance of a medical research librarian (DB). The final electronic literature search was conducted on August 2017, without date limits, using Medline, Pre-Medline, Embase, PsycInfo and Maternity and Infant Care via Ovid (http://www.ovid.com/), Scopus (http://www.scopus.com/), Proquest (http://www.proquest.com/), Web of Science (http://apps.webofknowledge.com) and Cumulative Index to Nursing and Allied Health Literature via EBSCO (http://www.ebsco.com/cinahl). The Cochrane Library was also searched separately to identify any similar systematic reviews that have been conducted previously. The following keywords were used: pregnan*, cognit*, neurodevelopment, infant*, child*, randomised control* trial, clinical trial, food* nutrition and supplement*. Full details of the search strategy tailored for each database can be found in Supplementary Table S1. The Boolean operation (i.e., AND, OR) was used to combine keywords that were searched as free text in the title, abstract or topics from all papers. All searches were limited to human studies published in English and citations were downloaded into the reference manager program ENDNOTE X6.v (New York, NY, USA: Thomson Reuters 2012). The keywords: mental development and motor development, were searched in all databases separately, to ensure that no eligible studies were missed from the initial search strategy.

2.2. Study Selection

Eligibility of the retrieved publications was assessed by two independent reviewers (RMT and SMF) as recommended by PRISMA [29]. Publication title and abstracts were evaluated against the inclusion and exclusion criteria (Table 1) Publications were excluded by a hierarchical approach based on the study design, population, intervention and outcomes. Publications meeting the initial eligibility screening were retrieved in full and were then subjected to a second round of screening by the same two reviewers to determine the final inclusion or exclusion. Assessment discrepancies between reviewers were resolved by discussion or assessment by a third party independent reviewer (AJH). The reference lists of the included publications were searched separately to identify any relevant articles that were not detected by the electronic search strategy. The abstract and then the full texts were retrieved for consideration. After the retrieval of the full text a final decision was made about the eligibility of the publication by the first reviewer (RMT).

<table>
<thead>
<tr>
<th>Criteria</th>
<th>Study Design</th>
<th>Population</th>
<th>Intervention</th>
<th>Outcome</th>
</tr>
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<tbody>
<tr>
<td><strong>Include</strong></td>
<td>Randomised or pseudorandomised controlled trials of any date</td>
<td>Pregnant women of any age or ethnicity</td>
<td>Dietary intervention/s, including dietary counselling and education as well as food/s, fortified foods or nutrient supplement/s</td>
<td>Measures cognitive outcomes of infants and children using cognitive assessment tests</td>
</tr>
<tr>
<td></td>
<td>Singleton pregnancies</td>
<td>Dietary intervention/s provided during pregnancy (exclusively) or intervention/s commencing during gestation and continued during lactation</td>
<td>Cognition is measured after birth in children ≤10 years of age</td>
<td></td>
</tr>
<tr>
<td><strong>Exclude</strong></td>
<td>All other designs including animal studies</td>
<td>Women that are not pregnant</td>
<td>Dietary intervention/s are not provided to pregnant women</td>
<td>Cognitive outcomes are not measured in children after pregnancy</td>
</tr>
<tr>
<td></td>
<td>Multiple births for the primary population</td>
<td>Dietary intervention/s that commence after pregnancy</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
2.3. Eligibility Criteria

Table 1 outlines the inclusion and exclusion criteria for this systematic review. Studies that delivered nutritional interventions to the mother during pregnancy (exclusively) or during pregnancy plus the post-natal period (<10 years) were included. This review included studies that measured cognition in children up to nine years of age because the effect of nutritional interventions may not be apparent until later life [18]. In addition, cognition assessment tests in older children (>7 years) are a more reliable predictor of adult cognitive function [31]. Nutritional interventions that commenced after pregnancy were excluded. This review included studies that measured cognition (as a primary or secondary outcome) in infants and young children using one or more cognition assessment tests or subtests. Cognition domains were based on the Cattell–Horn–Carroll (CHC) Theory of Intelligence [32]. The domains fluid and crystallised intelligence, specified in the CHC model [32], were combined to form the global cognition domain for cognition assessment tests that reported an overall global composite score. Verbal composite scores were grouped in the crystallised intelligence domain and non-verbal composite scores were grouped in the fluid intelligence domain. Short-term memory and long-term retrieval domains were combined to form an overall memory domain. The remaining six cognition domains of the CHC model [32] (quantitative reasoning, reading and writing ability, visual processing, auditory processing, processing speed and correct/decision speed) were included. Attention, motor skills and behaviour were also analysed because these domains are commonly analysed in cognition assessment tests. For example, the Intergrowth-21st Neurodevelopment Assessment reports child cognitive scores for the domains attention, motor skills and behaviour [33]. In addition, Bayley Scales of Infant development (third edition) includes a behaviour rating scale, which measures behavioural factors including orientation/engagement and emotional regulation [34]. This review included the outcome visual processing that focuses on both the response and comprehension of visual stimuli. Visual acuity which primarily focuses on vision clarity, was not reported as an outcome in this review. Similar methods have been described previously by Eilander et al. [35].

2.4. Quality Assessment

Articles considered eligible were assessed for methodological quality in their reporting using the American Dietetic Association (ADA) Quality Criteria Checklist for Primary Research [36] which is applicable to evaluating the validity of nutrition studies. The quality of the articles was assessed by two independent reviewers (RMT and SMF) and assessment discrepancies between reviewers, were managed through discussion or resolved by the third independent reviewer (AJH). The Quality Criteria Checklist [36] rates the study design and execution, as well as the risk of bias using ten validity questions, including four priority questions which must be satisfactory to gain a positive rating. Based on the responses to these questions, the checklist assigns a quality rating: positive (answered “yes” to six or more validity questions, including all four priority questions), negative (answered “no” to six or more validity questions) or neutral (answered “no” to one or more of the four priority criteria questions) [36]. No studies were given a negative rating in this review, therefore no study exclusions were made.

2.5. Data Extraction

The first reviewer (RMT) extracted relevant data from the included studies using an Excel spreadsheet. The Organisation for Economic Co-operation and Development (OECD) criteria [37] were used to classify country income into four categories: low, lower-middle, higher-middle and high. Nutritional interventions were divided into six categories; single nutrient supplement, multiple micronutrient supplement, fortified foods, foods, supplements with foods, supplements with foods and nutrition education or no nutrition intervention (control group). For the purpose of this review a multiple micronutrient supplement was defined as two or more nutrients which is consistent with
other authors [38–40]. The cognition outcomes data included the mean, standard deviation, confidence intervals, minimum and maximum range and odds ratio. Corresponding authors were emailed if additional data or clarification was required.

2.6. Data Synthesis

The nutritional interventions provided across the included studies were grouped together according to the nutrient supplied and the type of intervention (i.e., supplement, dietary changes). The cognition outcomes reported were organised into related cognition domains and the type of scoring (i.e., subtest or composite score). Thirteen cognition domain categories were derived: (i) fluid intelligence; (ii) crystallised intelligence; (iii) quantitative knowledge; (iv) memory; (v) visual processing; (vi) auditory processing; (vii) processing speed; (viii) correct decision speed; (ix) reading and writing; (x) global cognition; (xi) attention; (xii) motor skills; (xiii) behaviour.

2.7. Statistical Analysis

The aim of meta-analysis was to present a single effect size for the 13 specified cognition scores. The standardised mean difference (SMD) was used to measure the effect considering that all outcomes were measured using different cognition assessment tests and reporting units. The SMD, also known as Cohen’s d, is defined as the difference between the mean of the intervention and control group divided by the standard deviation of the data. Methods to convert other units (e.g., Odds ratio) to SMD are described in Sanchez-Meca et al. [41]. Studies that presented multiple cognition outcomes within the same cognition domain, multiple time-points, or multiple interventions for multiple micronutrient supplementation with only a single control group, the respective method described by Borenstein et al. [42] was used to calculate a summary SMD and standard error which accounts for non-independence within each study. As the correlations between outcomes were not reported by the studies, the correlation was assumed to be 0.5, and sensitivity analyses were performed using a higher (0.8) or lower (0.2) correlation to assess the changes in the standard error.

Supplementary Table S2 shows an example of how the composite effect size was created. A random effects meta-analytic model (using the method of DerSimonian and Laird [43]) was conducted for each cognitive outcome, where the observed effect sizes are regarded to vary due to sampling variance. A fixed effects model using inverse variance weights was also conducted for secondary analysis. The heterogeneity and total variation of studies were analysed using the I$^2$ statistic [44]. Funnel plots were used to assess publication bias. Subgroup analysis was performed for each outcome by the nutrient of interest. A sensitivity analysis with a random effects model was conducted in which the studies were stratified by country income. The statistical analysis was conducted using the metan command in the statistical software package STATA 13 (College Station, TX, USA: StataCorp LP 2013).

3. Results

3.1. Search Results

A total of 1195 publications (excluding duplicates, n = 439) were assessed against the inclusion and exclusion criteria (Figure 1). Seven included publications were identified from hand searching the reference lists of included publications detected from the initial search strategy. These articles all used the keyword mental development, which was not included separately. This systematic review resulted in 34 included publications after inclusion and exclusion criteria were applied to full texts.
3.2. Quality

A summary of the ADA quality assessment including participant selection, handling of withdrawals and the use of study blinding for each trial is presented in Supplementary Table S3. Direct (e.g., blood, urine tests) intervention compliance measures were used in 12 out of 34 studies [45–56]. Indirect (e.g., pill count, self-reported) intervention compliance measures were used in 15 out of 34 studies [57–71]. The attrition rate ranged from 0–99% indicating the potential for bias with a loss to follow-up. Post-hoc data analysis was used in 22 studies [46,47,49–51,53,55–58,60,61,63–67,69,72–75].
3.3. Description of Studies

Table 2 summarises the characteristics of the included studies. The earliest publication was from 1979 [76] and the latest from 2017 [72,75]. A summary of the studies from the OECD is provided in Supplementary Table S5. In summary, 50% of studies were from high-income countries, 29% of studies were from middle-income countries and 21% of studies were from low-income countries.

3.4. Participants

The study populations were mostly healthy pregnant women, except one publication [61] which included pregnant women with human immunodeficiency virus type 1 (HIV-1), although pregnant women were excluded if they were diagnosed with acquired immunodeficiency syndrome (AIDS) according to the World Health Organisation (WHO) definition [77].

3.5. Intervention

A summary of the nutritional interventions of the studies is provided in Supplementary Table S4. In summary, 76% of studies provided a nutrient supplement exclusively for the study intervention, no studies provided nutrition counselling or education as a stand-alone intervention.

3.6. Cognitive Outcomes

Table 2 summarises the cognition assessment tools used in each study. Further details about the cognition assessment tests used is provided in Supplementary Table S5. The age of child cognitive testing ranged from less than two weeks [76] to nine years old [71], however 62% of studies (n = 13 trials) reported cognition outcomes in children less than 35 months, and 46% of these studies (n = 6 trials) provided LCPUFA interventions in high-income countries. Meta-analysis was completed for nine cognition outcomes measured in the 34 included studies. The analyses for each cognition domain is described below, however in summary all cognition domains were not significantly affected by nutritional interventions.
Table 2. Overview of the characteristics of the randomised controlled trials (RCTs).

<table>
<thead>
<tr>
<th>Source, Year, Country, OECD</th>
<th>Infant Sample (n)</th>
<th>Intervention Group/s</th>
<th>Control Group/s</th>
<th>Intervention Duration</th>
<th>Cognitive Testing Age</th>
<th>Cognitive Assessment Tests</th>
<th>Main Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Choline Intervention vs. Control</strong></td>
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</tr>
<tr>
<td>Cheatham et al. [45] 2012, US, 1</td>
<td>99</td>
<td>Supplement: Vitamins: 750 mg/day choline</td>
<td>Supplement: corn oil capsules</td>
<td>18 weekends gestation to 3 months postnatally</td>
<td>10 and 12 month</td>
<td>Visuospatial Memory Delayed Response Task [78–80], Imitation paradigms [81], CDI [82] &amp; MSEL [83]</td>
<td>There were no significant differences between the intervention and control groups for global development, language, short-term visuospatial memory, or long-term episodic memory test scores (p &gt; 0.05) at 10 and 12 months</td>
</tr>
<tr>
<td><strong>Iodine Intervention vs. Control</strong></td>
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<tr>
<td>Santiago et al. [54], Spain, 1</td>
<td>102</td>
<td>Supplement: Mineral: I1: 200 µg/day iodine I2: 300 µg/day iodine</td>
<td>Fortified food: used iodised salt</td>
<td>Before 10 weekends gestation to delivery</td>
<td>6 to 18 month</td>
<td>BSID-III [34]</td>
<td>There were no significant differences between the groups for MDI and PDI scores (p &gt; 0.05) at 6 to 18 months</td>
</tr>
<tr>
<td>Zhou et al. [70], Australia, 1</td>
<td>53</td>
<td>Supplement: Mineral: 150 µg/day iodine</td>
<td>No supplement</td>
<td>Less than 20 weekends gestation to delivery</td>
<td>18 month</td>
<td>BSID-III [34]</td>
<td>There was no significant difference between the intervention compared to the control group for child cognition (p = 0.42), language (p = 0.83) and motor skills (p = 0.61) at 18 months of age</td>
</tr>
<tr>
<td>Zhou et al. [65], Australia, 1</td>
<td>302</td>
<td>Supplement: Mineral: 20 mg/day Fe</td>
<td>No supplement</td>
<td>20 weekends gestation to delivery</td>
<td>48 months</td>
<td>SB4 [84] &amp; SDQ [85]</td>
<td>There were no significant differences between groups for child cognition and behaviour test scores (p &gt; 0.05) at 48 months</td>
</tr>
<tr>
<td><strong>Iron and Folic Acid or Multiple Micronutrient Intervention vs. Control</strong></td>
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<tr>
<td>Chang et al. [73], China, 3</td>
<td>850</td>
<td>Supplement: Vitamins: I1: 400 µg/day FA I2: 400 µg/day FA, 800 µg/day vit A, 1.4 mg/day vit B1, 1.4 mg/day vit B2, 18 mg/day vit B3, 1.9 mg/day vit B6, 2.6 µg/day vit B12, 70 µg/day vit C, 5 µg/day vit D, 10 mg/day vit E Minerals: I1: 60 mg/day Fe I2: 2 mg/day Cu, 150 µg/day iodine, 30 mg/day Fe, 65 µg/day Se, 15 mg/day Zn</td>
<td>Supplement: Vitamin: 400 µg/day FA</td>
<td>Less than 28 weekends gestation to delivery</td>
<td>3, 6, 12, 18 and 24 months</td>
<td>BSID-II (Chinese translation) [86,87]</td>
<td>The prenatal IDA intervention groups had a lower MDI (p = 0.046 in folic acid; p = 0.034 in multiple micronutrient supplements) compared to the prenatal-non-IDA group. There was no significant difference in MDI scores in the prenatal-IDA group and prenatal-non-IDA group that received iron and folic acid supplements (p = 0.641)</td>
</tr>
</tbody>
</table>
Table 2. Cont.

<table>
<thead>
<tr>
<th>Source, Year, Country, OECD</th>
<th>Infant Sample (n)</th>
<th>Intervention Group/s</th>
<th>Control Group/s</th>
<th>Intervention Duration</th>
<th>Cognitive Testing Age</th>
<th>Cognitive Assessment Tests</th>
<th>Main Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>Li et al. [60], China, 2</td>
<td>1305</td>
<td>Supplement: Vitamins: I1: 800 µg/day vit A, 1.4 mg/day vit B1, 1.4 mg/day B2, 18 mg/day vit B3, 1.9 mg/day vit B6, 2.6 mg/day vit B12, 70 mg/day vit C, 10 µg/day vit D, 10 mg/day vit E, 400 µg/day FA I2: 400 µg/day FA Minerals: I1: 2 mg/day Cu, 30 mg/day Fe, 150 µg/day iodine, 65 µg/day Se I2: 30 mg/day Fe</td>
<td>Supplement Vitamins: 400 µg/day FA</td>
<td>27 weekends gestation to delivery</td>
<td>3, 6 and 12 months</td>
<td>BSID-II [86]</td>
<td>Positive effects were demonstrated from the multiple micronutrient supplement on child MDI scores at 12 months (p = 0.01)</td>
</tr>
<tr>
<td>Hanieh et al. [48], Vietnam, 3</td>
<td>1168</td>
<td>Supplement taken twice weekly Vitamins: I1: 1500 µg/day FA I2: 1600 µg/day vit A, 2.8 mg/day vit B1, 2.8 mg/day vit B2, 36 mg/day vit B3, 3.8 mg/day vit B6, 2.8 mg/day vit B12, 140 mg/day vit C, 10 mg/day vit D, 20 mg/day vit E, 1500 µg/day FA Minerals: I1: 60 mg/day Fe I2: 4 mg/day Cu, 60 mg/day Fe, 300 µg/day iodine, 130 µg/day Se</td>
<td>Supplement taken daily Vitamins: 400 µg/day FA Mineral: 60 mg/day Fe</td>
<td>Less than 16 weekends gestation to 3 months postnatally</td>
<td>6 months</td>
<td>BSID-III [34]</td>
<td>A positive effect was demonstrated from the twice weekly iron and folate acid supplement groups on child cognitive development scores (MD 1.89; 95% CI 0.23 to 3.56) at 6 months</td>
</tr>
<tr>
<td>Brei et al. [72], Germany, 1</td>
<td>130</td>
<td>Supplement: fish oil capsules LCPUFAs: 1020 mg/day DHA, 180 mg/day EPA Vitamins: 9 mg/day vit E Dietary counselling to lower ARA intake</td>
<td>Participants received information for eating a healthy diet during pregnancy</td>
<td>15 weekends gestation to 4 months postnatally</td>
<td>60 months</td>
<td>Child Development Inventory (German translation) [68,89], MM [90,91]</td>
<td>There were no differences between the maternal fish oil supplemented group compared to the control group on child CDI scores and MM at 4 and 5 years of age (p &gt; 0.05)</td>
</tr>
<tr>
<td>Dunstan et al. [46], Australia, 1</td>
<td>72</td>
<td>Supplement: fish oil capsules LCPUFAs: 2.2 g/day DHA, 1.1 g/day EPA Vitamins: 3–4 mg/day vit E</td>
<td>Supplement: olive oil capsules LCPUFA: 2.7 g/day olean acid Vitamins: 3–4 mg/day vit E</td>
<td>20 weekends gestation to delivery</td>
<td>18 months</td>
<td>GMDS 0-2 [92], PPVT-III [93] &amp; CBCL [94]</td>
<td>A positive effect was demonstrated from the maternal fish oil intervention on eye and hand coordination (p = 0.021) in children at 18 months</td>
</tr>
<tr>
<td>Gould et al. [47], Australia, 1</td>
<td>185</td>
<td>Intervention: marine oil capsules LCPUFAs: 0.8 g/day DHA, 0.1 g/day EPA</td>
<td>Supplement: capsule with a blend of vegetable oils</td>
<td>18–21 weekends gestation to delivery</td>
<td>27 months</td>
<td>Single and multiple object task [95,96] &amp; A-not-B task [97]</td>
<td>There were no significant differences between the intervention and control groups for child attention, working memory and inhibitory control development test scores (p &gt; 0.05)</td>
</tr>
<tr>
<td>Source, Year, Country, OECD</td>
<td>Infant Sample (n)</td>
<td>Intervention Group/s</td>
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<tr>
<td>Helland et al. [49], Norway, 1</td>
<td>262</td>
<td>Supplement: cod liver oil LCPUFAs: 75 mg/day ALA, 27.5 mg/day ARA, 1.18 g/day DHA, 0.8 g/day EPA, 160 mg/day LA; Vitamins: 117 µg/day vit A, 1 µg/day vit D, 1.4 mg/day vit E</td>
<td>Supplement: corn oil LCPUFAs: 92 mg/day ALA, 8.3 mg/day DHA, 4.7 g/day LA; Vitamins: 117 µg/day vit A, 1 µg/day vit D, 1.4 mg/day vit E</td>
<td>17–19 weekends gestation to 3 months postnatally</td>
<td>27 and 39 weekends</td>
<td>FTII-II [98]</td>
<td>There were no significant differences between the intervention and control groups for cognitive development test scores ($p &gt; 0.05$) at 27 and 39 months</td>
</tr>
<tr>
<td>Helland et al. [51], Norway, 1</td>
<td>90</td>
<td>Supplement: cod liver oil LCPUFAs: 75 mg/day ALA, 27.5 mg/day ARA, 1.18 g/day DHA, 0.8 g/day EPA, 160 mg/day LA; Vitamins: 117 µg/day vit A, 1 µg/day vit D, 1.4 mg/day vit E</td>
<td>Supplement: corn oil LCPUFAs: 92 mg/day ALA, 8.3 mg/day DHA, 4.7 g/day LA; Vitamins: 117 µg/day vit A, 1 µg/day vit D, 1.4 mg/day vit E</td>
<td>1–19 weekends gestation to 3 months postnatally</td>
<td>48 months</td>
<td>KABC-II [99]</td>
<td>A positive effect was demonstrated from the cod liver oil supplement on child mental processing composite score ($p = 0.049$) at 48 months</td>
</tr>
<tr>
<td>Helland et al. [50], Norway, 1</td>
<td>143</td>
<td>Supplement: cod liver oil LCPUFAs: 75 mg/day ALA, 27.5 mg/day ARA, 1.18 g/day DHA, 0.8 g/day EPA, 160 mg/day LA; Vitamins: 117 µg/day vit A, 1 µg/day vit D, 1.4 mg/day vit E</td>
<td>Supplement: corn oil LCPUFAs: 92 mg/day ALA, 8.3 mg/day DHA, 4.7 g/day LA; Vitamins: 117 µg/day vit A, 1 µg/day vit D, 1.4 mg/day vit E</td>
<td>1–19 weekends gestation to 3 months postnatally</td>
<td>84 months</td>
<td>KABC-II [99]</td>
<td>No significant differences between the intervention and control groups for cognitive development test scores ($p &gt; 0.05$) at 84 months was found</td>
</tr>
<tr>
<td>Hurtado et al. [68], Spain, 1</td>
<td>61</td>
<td>Supplement: fish oil drink per 100 mL</td>
<td>Supplement drink per 100 mL</td>
<td>Energy: 145 kJ</td>
<td>Macronutrients: 3.7 g prot., 6.7 g carb., 1.8 g total fat</td>
<td>LCPUFAs: 80 mg DHA, 18 mg EPA</td>
<td>Vitamins: 0.4 µg vit B12, 9 mg vit C, 0.75 µg vit D, 1.5 mg vit E, 80 µg FA Minerals: 23 µg iodide, 2.2 mg Fe</td>
</tr>
<tr>
<td>Judge et al. [59], US, 1</td>
<td>29</td>
<td>Muesli bar with fish oil. Per bar; Energy: 386 kJ Macronutrients: 18 g carb., 0.3 g fibre, 1.3 g prot., 1.7 g total fat, 300 mg DHA, 38 mg EPA</td>
<td>Muesli bar without fish oil. Per bar; Energy: 386 kJ Macronutrients: 18 g carb., 0.3 g fibre, 1.3 g prot., 1.7 g total fat</td>
<td>24 weekends gestation until delivery</td>
<td>2-step problem-solving test [100–103] &amp; FTII-II [98]</td>
<td>Positive effects were demonstrated from the fish oil intervention group on the problem-solving task ($p = 0.017$), total intentional score ($p = 0.011$) and the number of intentional solution on both cloth ($p = 0.008$) and cover steps ($p = 0.004$) at 9 months</td>
<td></td>
</tr>
</tbody>
</table>
Table 2. Cont.

<table>
<thead>
<tr>
<th>Source, Year, Country, OECD</th>
<th>Infant Sample (n)</th>
<th>Intervention Group/s</th>
<th>Control Group/s</th>
<th>Intervention Duration</th>
<th>Cognitive Testing Age</th>
<th>Cognitive Assessment Tests</th>
<th>Main Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>Makrides et al. [52], Australia, 1</td>
<td>726</td>
<td>Supplement: fish oil capsules LCPUFAs: 800 mg/day DHA, 100 mg/day EPA</td>
<td>Supplement: Capsules with vegetable oil blend</td>
<td>21 weekends gestation to delivery</td>
<td>18 months</td>
<td>BSID-III [34]</td>
<td>There were no significant differences between the groups for cognitive and language test scores ($p &gt; 0.05$)</td>
</tr>
<tr>
<td>Mulder et al. [62], Canada 4, 1</td>
<td>216</td>
<td>Supplement: algal oil capsules LCPUFAs: 400 mg/day DHA</td>
<td>Supplement: soybean and corn oil capsules</td>
<td>16 weeks gestation to delivery</td>
<td>9 and 18 months</td>
<td>CDI [82] &amp; BSID-III [34]</td>
<td>Negative effects were demonstrated from the maternal algal oil supplement for CDI language ($p &lt; 0.05$) at 14 months and 18 months and BSID-III language scores at 18 months ($p &lt; 0.05$)</td>
</tr>
<tr>
<td>Ramakrishnan et al. [74], Mexico, 2</td>
<td>797</td>
<td>Supplement: algal oil capsules LCPUFAs: 400 mg/day DHA</td>
<td>Supplement: corn and soy oil capsules</td>
<td>18–22 weeks Gestation to delivery</td>
<td>60 months</td>
<td>MSCA (Spanish translation) [104], (BASC-2) [105], and (K-CPT) [106]</td>
<td>There were no significant differences between the intervention group compared to the control group for MSCA scores ($p &gt; 0.05$), BASC-2 or scores at 5 years. of age. The intervention did significantly affect K-CPT omissions scores ($p = 0.01$) compared to the placebo group</td>
</tr>
<tr>
<td>Tofail et al. [63], Bangladesh, 4</td>
<td>249</td>
<td>Supplement: fish oil capsules LCPUFAs: 1.2 g/day DHA, 1.8 g/day EPA</td>
<td>Supplement: soy oil capsules LCPUFAs: 2.3 g/day LA</td>
<td>27 weekends gestation to delivery</td>
<td>10 months</td>
<td>BSID-II [86] &amp; Wolke’s Behaviour Rating Scale [107]</td>
<td>There were no significant differences between the intervention and control groups for cognitive or behaviour development test scores ($p &gt; 0.05$) at 10 months</td>
</tr>
<tr>
<td>Van Goor et al. [56], Netherlands, 1</td>
<td>114</td>
<td>Fish oil capsules I1: 220 mg/day DHA, 34 mg/day EPA, 15 mg/day ARA, 274 mg/day LA, 32 mg/day ALA I2: 220 mg/day DHA, 36 mg/day EPA, 220 mg/day ARA, 46 mg/day LA, 7 mg/day ALA</td>
<td>Supplement: soy bean oil capsules</td>
<td>14–20 weeks gestation to 3 months postnatally</td>
<td>18 months</td>
<td>BSID-II [86]</td>
<td>There were no significant differences between the control and intervention groups for cognitive development test scores ($p &gt; 0.05$) at 18 months</td>
</tr>
</tbody>
</table>

**LCPUFA and Folic Acid Intervention vs. Control**

<p>| Catena et al. [66], Germany, Spain &amp; Hungary, 1 | 136 | Dairy drink per sachet: I1: 500 mg/day DHA, 150 mg/day EPA I2: 400 µg/day EPA I3: 500 mg/day DHA, 150 mg/day EPA, 400 µg/day FA | Dairy drink with vitamins and minerals in accordance to European dietary guidelines for pregnancy | Less than 20 weekends gestation to delivery | 102 months | Attention Network Test (child version) [108,109] | Children born to mothers supplemented with folic acid alone solved the response conflict more quickly than the control group and the fish oil and folic acid group ($p &lt; 0.05$) |</p>
<table>
<thead>
<tr>
<th>Source, Year, Country, OECD</th>
<th>Infant Sample (n)</th>
<th>Intervention Group/s</th>
<th>Control Group/s</th>
<th>Intervention Duration</th>
<th>Cognitive Testing Age</th>
<th>Cognitive Assessment Tests</th>
<th>Main Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>Multiple Micronutrient Intervention vs. Control</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Christian et al. [71], Nepal, 4</td>
<td>676</td>
<td>Supplement: Vitamin: 11: 1000 µg/day vit A, 400 µg/day FA I2: 1000 µg/day vit A, 400 µg/day FA I3: 1000 µg/day vit A, 400 µg/day FA I4: 1000 µg/day vit A, 400 µg/day FA, 1.6 mg/day vit B1, 1.8 mg/day vit B2, 20 mg/day vit B3, 2.2 µg/day vit B6, 2.6 µg/day vit B12, 100 mg/day vit C, 10 µg/day vit D, 10 mg/day vit E, 65 µg/day vit K, Minerals: I2: 60 mg/day Fe I3: 60 mg/day Fe, 30 mg/day Zn I4: 2 mg/day Cu, 60 mg/day Fe, 100 mg/day Mg, 30 mg/day Zn</td>
<td></td>
<td></td>
<td>Pregnancy to 3 months postnatally</td>
<td>84–108 months</td>
<td>UNIT [110], Go/no-go [111], the Stroop test [112], backward digit span [113], MABC [114], finger-tapping test [115]</td>
</tr>
<tr>
<td>Christian et al. [67], Bangladeshi, 4</td>
<td>734</td>
<td>Supplement: Vitamin: 11: 770 µg/day vit A, 1.4 mg/day vit B1, 1.4 mg/day vit B2, 18 mg/day vit B3, 1.9 mg/day vit B6, 2.6 µg/day vit B12, 85 mg/day vit C, 5 µg/day vit D, 15 mg/day vit E, 600 µg/day FA, Minerals: 1000 µg/day Cu, 220 µg/day iodine, 27 mg/day Fe, 60 µg Se, 12 mg/day Zn</td>
<td></td>
<td></td>
<td>Pregnancy to 3 months postnatally</td>
<td>24 months</td>
<td>BSID-III [34]</td>
</tr>
<tr>
<td>Joos et al. [116], Taiwan, 2</td>
<td>198</td>
<td>Supplement: 1680 kJ, 50 g/day carb., 0.55 g/day fibre, 20 g/day prot., 13.3 g/day total fat, Vitamins: 750 µg/day vit A, 0.8 mg/day vit B1, 0.9 mg/day vit B2, 10 mg/day vit B3, 0.08 mg/day vit B6, 1 mg/day vit B12, 37.5 mg/day vit C, 5 µg/day vit D, 3.35 mg/day vit E, Minerals: 0.5 mg/day Ca, 0.5 mg/day Cu, 6 mg/day Fe, 1 mg/day Mn, 0.9 mg/day K, 0.2 mg/day Na, 0.4 mg/day</td>
<td>No supplement</td>
<td>Conception to lactation of 2nd child</td>
<td>8 months</td>
<td>BSID-I [117]</td>
<td>A positive effect was demonstrated from the multiple micronutrient supplement on PDI (mean ± SD: 3.80 ± 1.90 compared to 3.31 ± 1.71) at 8 months</td>
</tr>
</tbody>
</table>
Table 2. Cont.

<table>
<thead>
<tr>
<th>Source, Year, Country, OECD</th>
<th>Infant Sample (n)</th>
<th>Intervention Group/s</th>
<th>Control Group/s</th>
<th>Intervention Duration</th>
<th>Cognitive Testing Age</th>
<th>Cognitive Assessment Tests</th>
<th>Main Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>Prado et al. [53], Indonesia, 3</td>
<td>487</td>
<td>Supplement</td>
<td>Vitamins: 800 µg/day vit A, 1.4 mg/day vit B1, 1.4 mg/day vit B2, 18 mg/day vit B3, 1.9 mg/day vit B6, 2.6 mg/day vit B12, 70 mg/day vit C, 5 µg/day vit D, 100 mg/day vit E, 400 µg/day FA Minerals: 2 µg/day Cu, 30 mg/day Fe, 150 µg/day iodine, 65 µg/day Se</td>
<td>Supplement</td>
<td>Vitamins: 400 µg/day FA Minerals: 30 mg/day Fe</td>
<td>Pregnancy to 3 months postnatally</td>
<td>42 months</td>
</tr>
<tr>
<td>Tofail et al. [64], Bangladesh, 4</td>
<td>2853</td>
<td>Food &amp; supplements</td>
<td>Energy: 2512 kJ, Vitamins: I1: 400 µg folate I2: 1.4 mg/day vit B1, 1.4 mg/day vit B2, 18 mg/day vit B3, 2.6 mg/day vit B6, 1.9 mg/day vit B12, 70 mg/day vit C, 5 µg/day vit D, 10 mg/day vit E, 400 µg/day FA Minerals: I1: 30 mg/day Fe I2: 2 mg/day Cu, 60 mg/day Fe, 150 µg/day iodine, 65 µg/day Se 15 mg/day Zn</td>
<td>Food &amp; supplements</td>
<td>Energy: 2512 kJ, Vitamins: I1: 400 µg folate I2: 1.4 mg/day vit B1, 1.4 mg/day vit B2, 18 mg/day vit B3, 2.6 mg/day vit B6, 1.9 mg/day vit B12, 70 mg/day vit C, 5 µg/day vit D, 10 mg/day vit E, 400 µg/day FA Minerals: I1: 30 mg/day Fe I2: 2 mg/day Cu, 60 mg/day Fe, 150 µg/day iodine, 65 µg/day Se 15 mg/day Zn</td>
<td>17 weekends gestation to delivery</td>
<td>7 months</td>
</tr>
<tr>
<td>Vuori et al. [76], Colombia, 2</td>
<td>244</td>
<td>Food &amp; supplements:</td>
<td>Energy: 3595 kJ Prot: 38.4 g/day Vitamin: 1807 µg/day vit A Mineral: 18 mg/day Fe</td>
<td>No supplement</td>
<td>27 weekends gestation and continued during lactation</td>
<td>15 days</td>
<td>Checkboard activity [76]</td>
</tr>
</tbody>
</table>
Table 2. Cont.

<table>
<thead>
<tr>
<th>Source, Year, Country, OECD</th>
<th>Infant Sample (n)</th>
<th>Intervention Group/s</th>
<th>Control Group/s</th>
<th>Intervention Duration</th>
<th>Cognitive Testing Age</th>
<th>Cognitive Assessment Tests</th>
<th>Main Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>Waber et al. [125], Colombia, 2</td>
<td>304</td>
<td>Food, supplements &amp; nutrition education 3595 kJ energy, 38.4 g prot., 25.8 g total fat. Vitamins: 1807 µg vit A, 5.4 mg vit B1, 6.6 mg vit B2, 31.6 mg vit B3, 104.6 mg vit C. Mineral: 1146 mg Ca</td>
<td>No supplement</td>
<td>27 weekends gestation to 6 months postnatally</td>
<td>4, 6, 12, 18, 24 and 36 months</td>
<td>GMDS 0–2 [92] &amp; Escalona &amp; Corman’s Albert Einstein Scales of Sensorimotor Development [126]</td>
<td>Positive effects were demonstrated from food supplements on subtests including personal social (p = 0.006), speech and language (p = 0.019), eye and hand coordination (p = 0.008), as well as general quotient (p = 0.003)</td>
</tr>
<tr>
<td>Multiple Micronutrient or Vitamin A Intervention vs. Control</td>
<td></td>
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<tr>
<td>McGrath et al. [61], Tanzania, 4</td>
<td>327</td>
<td>Supplement: Vitamins: I1: 1500 µg/day vit A, 30 mg/day beta-carotene, 20 mg/day vit B1, 20 mg/day vit B2, 100 mg/day vit B3, 25 mg/day vit B6, 50 mg/day vit B12, 500 mg/day vit C, 30 mg/day vit E, 800 µg/day FA I2: 1500 µg/day vit A, 30 mg/day beta-carotene, 20 mg/day</td>
<td>Supplement: Vitamins: C1: 20 mg/day vit B1, 20 mg/day vit B2, 100 mg/day vit B3, 25 mg/day vit B6, 50 mg/day vit B12, 500 mg/day vit C, 30 mg/day vit E, 800 µg/day FA C2: No supplements</td>
<td>12–27 weekends gestation to 18 months in postnatal period</td>
<td>6, 12 and 18 months</td>
<td>BSID-II [86]</td>
<td>Positive effects were demonstrated from the multiple micronutrient supplement by increasing PDI score by 2.6 points (95% CI: 0.1–5.1) over 6 to 18 months Vitamin A supplementation was also associated with 2.8 point increase (95% CI: 0.4–5.2) in PDI score at 6 months</td>
</tr>
<tr>
<td>Vitamin A, Folic Acid and Iron Intervention vs. Control</td>
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<tr>
<td>Schmidt et al. [69], Indonesia, 4</td>
<td>188</td>
<td>Supplement: Vitamins: 4800 µg/day vit A, 500 µg/day FA. Mineral: 120 mg/day Fe</td>
<td>Supplement: Vitamin: 500 µg/day FA. Mineral: 120 mg/day Fe</td>
<td>16–20 weekends gestation to delivery</td>
<td>6 and 12 months</td>
<td>BSID-I [117]</td>
<td>There were no significant effects of vitamin A supplementation on MDI and PDI scores at 6 or 12 months of age (p &gt; 0.05). There was no significant difference between weekly compared to daily iron supplementation on child MDI and PDI scores (p &gt; 0.05)</td>
</tr>
<tr>
<td>Vitamin B12 Intervention vs. Control</td>
<td></td>
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</tr>
<tr>
<td>Srinivasan et al. [75], India, 3</td>
<td>178</td>
<td>Supplement: Vitamin: 50 µg/day vit B12</td>
<td>No supplement</td>
<td>Less than 14 weekends gestation to 6 weekends postnatally</td>
<td>9 months</td>
<td>BSID-III [34]</td>
<td>There were no significant effects of maternal B12 supplementation on child MDI and PDI scores at 9 months (p &gt; 0.05)</td>
</tr>
<tr>
<td>Zinc Intervention vs. Control</td>
<td></td>
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</tr>
<tr>
<td>Hamadani et al. [58], Bangladesh, 4</td>
<td>168</td>
<td>Supplement: Vitamin: 30 mg/day Zn</td>
<td>No supplement</td>
<td>Pregnancy to delivery</td>
<td>13 months</td>
<td>BSID-II [86] &amp; Wolke’s Behaviour Rating Scale [107]</td>
<td>Negative effects were demonstrated from maternal zinc supplementation on child MDI (p = 0.04) and psychomotor development index (PDI) (p = 0.04) at 13 months</td>
</tr>
</tbody>
</table>
Table 2. Cont.

<table>
<thead>
<tr>
<th>Source, Year, Country, OECD</th>
<th>Infant Sample (n)</th>
<th>Intervention Group/s</th>
<th>Control Group/s</th>
<th>Intervention Duration</th>
<th>Cognitive Testing Age</th>
<th>Cognitive Assessment Tests</th>
<th>Main Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tamura et al. [55], US, 1</td>
<td>347</td>
<td>Supplement:</td>
<td>No supplement</td>
<td>19 weekends gestation to delivery</td>
<td>60 months</td>
<td>DAS [127], VSM [128], ASM [128], Knox Cube [129], PDMS-2 [130] &amp; Grooved Pegboard [131]</td>
<td>There were no significant differences between the intervention and control groups for cognitive, memory or motor development test scores (p &gt; 0.05) at 60 months</td>
</tr>
</tbody>
</table>

Zinc, Folic Acid and Iron vs. Control

| Caulfield et al. [57], Peru, 2 | 184               | Supplement:          | Supplement:     | 10–14 weekends gestation to delivery | 54 months             | WPPSI-III [113], bear story [132,133], the counting game [57], draw a person [134], the friendship interview [135,136], Vineland’s II [97] & PBQ [137] | There were no significant differences between the intervention and control groups for child cognitive, social or behavioural development test scores (p > 0.05) at 54 months |

Supplement: Vitamins: 250 µg/day FA Minerals: 25 mg/day Zn, 60 mg/day Fe

Supplement: Vitamins: 250 µg/day FA Minerals: 60 mg/day Fe

3.7. Outcome 1: Behaviour

Ten RCTs [46,48,52,53,57,64,65,70,72,74] measured the effect of a nutritional intervention on behaviour composite scores. Multiple outcomes were combined to create a composite score in Brei et al. [72] (2 outcomes), Makrides et al. [52] (2 outcomes), Ramakrishnan et al. [74] (4 outcomes), Zhou et al. [65] (5 outcomes), Zhou et al. [70] (2 outcomes), Caulfield et al. [57] (5 outcomes), and Dunstan et al. [46] (4 outcomes). There was no significant effect of LCPUFA supplementation (n = 4 trials) [46,52,72,74], multiple micronutrient supplementation (n = 1 trial) [53], iron supplementation (n = 1 trial) [65], iodine supplementation (n = 1 trial) [70], zinc supplementation (n = 1 trial) [57] or a food-based intervention (n = 1 trial) [64] (Table 3) (Table 3 and Figure 2). Hanieh et al. [48] investigated the effect of nutrient supplements of varying dosage on cognition, no significant effect on infant behaviour composite score was reported. From the statistical tests of Egger (p = 0.72), there was no evidence of publication bias (Supplementary Figure S2). From the sensitivity analysis, the country-income of the studies did not significantly (p > 0.05) affect child behaviour outcomes in studies that provided LCPUFA supplementation (Supplementary Figure S3).

![Forest plot for child behaviour outcomes.](image-url)

**Figure 2.** Forest plot for child behaviour outcomes. The overall effect size was estimated by standardised mean difference (SMD). LCPUFA, long chain polyunsaturated fatty acids. D+L, random-effects estimate (Der Simonian and Laird method) and I-V, fixed-effects estimate (inverse variance method).
**Table 3.** Meta-analyses for cognitive outcomes in evaluation of nutritional interventions during pregnancy.

<table>
<thead>
<tr>
<th>Dietary Nutrient</th>
<th>Cognitive Outcome</th>
<th>Study Authors</th>
<th>Country (OECD)</th>
<th>Studies (n)</th>
<th>Children (n)</th>
<th>SMD (95% CI) †</th>
<th>p-Value</th>
<th>I² ‡</th>
</tr>
</thead>
<tbody>
<tr>
<td>LCPUFA</td>
<td>Attention</td>
<td>Gould et al. [47] Ramakrishnan et al. [74]</td>
<td>Australia (1) Mexico (2)</td>
<td>2</td>
<td>955</td>
<td>−0.07 (−0.17 to 0.03)</td>
<td>0.19</td>
<td>0%</td>
</tr>
<tr>
<td></td>
<td>Behaviour</td>
<td>Dunstan et al. [46] Makrides et al. [52] Brei et al. [72] Ramakrishnan et al. [74]</td>
<td>Australia (1) Australia (1) Germany (1) Mexico (2)</td>
<td>4</td>
<td>1725</td>
<td>−0.05 (−0.12 to 0.03)</td>
<td>0.25</td>
<td>0%</td>
</tr>
<tr>
<td>MMN</td>
<td>Motor skills</td>
<td>Li et al. [60] McGrath et al. [61] Prado et al. [53] Chang et al. [73] Christian [71] Christian [67]</td>
<td>China (2) Tanzania (4) Indonesia (3) China (3) Nepal (4) Bangladesh (4)</td>
<td>6</td>
<td>3572</td>
<td>0.02 (−0.04 to 0.17)</td>
<td>0.55</td>
<td>0%</td>
</tr>
<tr>
<td>LCPUFA</td>
<td>Motor skills</td>
<td>Dunstan et al. [46] Makrides et al. [52] Mulder et al. [62] Tofail et al. [63] Van Goor et al. [56] Brei et al. [72] Ramakrishnan et al. [74]</td>
<td>Australia (1) Canada (1) Bangladesh (4) Netherlands (1) Germany (1) Mexico (2)</td>
<td>7</td>
<td>2265</td>
<td>0.06 (−0.03 to 0.15)</td>
<td>0.22</td>
<td>8.9%</td>
</tr>
<tr>
<td>Zinc</td>
<td>Motor skills</td>
<td>Caulfield et al. [57] Hamadani et al. [58] Tamura et al. [55]</td>
<td>Peru (2) Bangladesh (4) United States (1)</td>
<td>3</td>
<td>985</td>
<td>−0.10 (−0.38 to 0.17)</td>
<td>0.49</td>
<td>72.5%</td>
</tr>
<tr>
<td>LCPUFA</td>
<td>Fluid intelligence</td>
<td>Brei et al. [72] Dunstan et al. [46] Ramakrishnan et al. [74]</td>
<td>Germany (1) Mexico (2)</td>
<td>3</td>
<td>999</td>
<td>0.05 (−0.08 to 0.18)</td>
<td>0.45</td>
<td>10.1%</td>
</tr>
<tr>
<td>MMN</td>
<td>Fluid intelligence</td>
<td>Christian et al. [71] Prado et al. [53]</td>
<td>Nepal (4) Indonesia (3)</td>
<td>2</td>
<td>755</td>
<td>0.07 (−0.20 to 0.33)</td>
<td>0.63</td>
<td>78.2%</td>
</tr>
<tr>
<td>Zinc</td>
<td>Fluid intelligence</td>
<td>Caulfield et al. [57] Tamura et al. [55]</td>
<td>Peru (2) United States (1)</td>
<td>2</td>
<td>539</td>
<td>−0.10 (−0.25 to 0.06)</td>
<td>0.23</td>
<td>0%</td>
</tr>
<tr>
<td>MMN</td>
<td>Global cognition</td>
<td>Joos et al. [116] Li et al. [60] McGrath et al. [61] Waber et al. [125] Chang et al. [73] Christian et al. [67]</td>
<td>Taiwan (2) China (2) Tanzania (4) Colombia (2) China (3) Bangladesh (4)</td>
<td>6</td>
<td>3126</td>
<td>0.09 (−0.02 to 0.19)</td>
<td>0.11</td>
<td>57.2%</td>
</tr>
</tbody>
</table>
Table 3. Cont.

<table>
<thead>
<tr>
<th>Dietary Nutrient</th>
<th>Cognitive Outcome</th>
<th>Study Authors</th>
<th>Country (OECD)</th>
<th>Studies (n)</th>
<th>Children (n)</th>
<th>SMD (95% CI) †</th>
<th>p-Value</th>
<th>I² ‡</th>
</tr>
</thead>
<tbody>
<tr>
<td>LCPUFA</td>
<td>Global cognition</td>
<td>Dunstan et al. [46]</td>
<td>Australia (1)</td>
<td>10</td>
<td>2632</td>
<td>0.03 (−0.07 to 0.13)</td>
<td>0.55</td>
<td>21.3%</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Helland et al. [49]</td>
<td>Norway (1)</td>
<td></td>
<td></td>
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<tr>
<td></td>
<td></td>
<td>Helland et al. [51]</td>
<td>Norway (1)</td>
<td></td>
<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Judge et al. [59]</td>
<td>United States (1)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Makrides et al. [52], Mulder et al. [62], Tofail et al. [63], Van Goor et al. [56], Brei et al. [72], Ramakrishnan et al. [74]</td>
<td>Australia (1), Canada (1), Bangladesh (4), Netherlands (1), Germany (1), Mexico (2)</td>
<td>10</td>
<td>2632</td>
<td>0.03 (−0.07 to 0.13)</td>
<td>0.55</td>
<td>21.3%</td>
</tr>
<tr>
<td>Zinc</td>
<td>Crystallised intelligence</td>
<td>Caulfield et al. [57], Tamura et al. [55]</td>
<td>Peru (2), United States (1)</td>
<td>2</td>
<td>539</td>
<td>−0.04 (−0.20 to 0.12)</td>
<td>0.61</td>
<td>0%</td>
</tr>
<tr>
<td>LCPUFA</td>
<td>Crystallised intelligence</td>
<td>Dunstan et al. [46], Makrides et al. [52], Mulder et al. [62], Brie et al. [72], Ramakrishnan et al. [74]</td>
<td>Australia (1), Canada (1), Germany (1), Mexico (2)</td>
<td>5</td>
<td>1941</td>
<td>0.25 (−0.04 to 0.53)</td>
<td>0.09</td>
<td>87.8%</td>
</tr>
<tr>
<td>MMN</td>
<td>Crystallised intelligence</td>
<td>Christian et al. [67], Prado et al. [53]</td>
<td>Bangladesh (4), Indonesia (3)</td>
<td>2</td>
<td>1207</td>
<td>0.01 (−0.11 to 0.12)</td>
<td>0.91</td>
<td>0%</td>
</tr>
</tbody>
</table>

CI: confidence interval; LCPUFA: long chain polyunsaturated fatty acids; MMN: multiple micronutrient; SMD: standardised mean difference; OECD: the organisation for economic co-operation and development criteria; 1 = high income country, 2 = higher middle income country, 3 = lower middle income country & 4 = low income country; † the main measure of effect was SMD (also known as Coehens d). The SMD was determined by taking the difference between the mean of the intervention group compared to the control group, and dividing the pooled standard deviation for the outcome across the whole trial. A random effects model using the method DerSimonian & Laird [43] was applied to the data; ‡ the I² statistic is the percentage of observed total variation across studies that is due to heterogeneity rather than chance. It is calculated using the following formula: $I^2 = 100\% \times (Q – df)/Q$, where Q is Cochran’s heterogeneity and df is the degrees of freedom [44].
3.8. Outcome 2: Memory

Four RCTs [45,55,65,74] measured the effect of a nutritional intervention on memory composite score. There was no significant effect of iron supplementation (n = 1 trial) [55], zinc supplementation (n = 1 trial) [65], choline supplementation (n = 1 trial) [45] or LCPUFA supplementation (n = 1 trial) [74] on memory composite scores.

3.9. Outcome 3: Motor Skills

Eighteen RCTs [46,48,52,53,55–58,60–64,67,71–74] measured the effect of a nutritional intervention on the motor skills score. Multiple outcomes were combined to create a composite score in Brei et al. [72] (2 outcomes), Dunstan et al. [46] (2 outcomes), Mulder et al. [62] (2 outcomes), and Christian et al. [71] (2 outcomes). Li et al. [60] Chang et al. [73], and Christian et al. [71] used two interventions and one control group, the sample populations were not independent, therefore the effect sizes were combined. McGrath et al. [61] also used multiple independent intervention and control groups, therefore the effect sizes were kept separate. There was no significant effect of multiple micronutrient interventions (n = 6 trials) [53,60,61,67,71,73], LCPUFA supplementation (n = 7 trials) [46,52,56,62,63,72,74], zinc supplementation (n = 3 trials) [55,57,58] or food-based intervention (n = 1 trial) [64] on motor skills composite scores (Table 3 and Figure 3). Hanieh et al. [48] investigated the effect of nutrient supplementation of varying dosage on cognition, no significant effect on infant motor skills composite score was reported. The heterogeneity among the zinc intervention trials was significant (p = 0.03). From the statistical tests of Egger (p = 0.54), there was no evidence of publication bias (Supplementary Figure S4). The heterogeneity among the zinc intervention trials was significant (p = 0.03). From the sensitivity analysis, the country-income of the studies did not significantly (p > 0.05) affect child motor skills (Supplementary Figure S5).

Figure 3. Forest plot for child motor skills outcomes. The overall effect size was estimated by standardised mean difference (SMD). LCPUFA, long chain polyunsaturated fatty acids; MMN, multiple micronutrient. D+L, random-effects estimate (Der Simonian and Laird method) and I-V, fixed-effects estimate (inverse variance method).
### 3.10. Outcome 4: Fluid Intelligence

Eight RCTs [46,53,55,65,71,72,74] measured the effect of a nutritional intervention on the fluid intelligence composite score. Multiple outcomes were combined to create a composite score in Brei et al. [72] (2 outcomes), and Ramakrishnan et al. [74] (2 outcomes). There was no significant effect of LCPUFA supplementation (n = 3 trials) [46,72,74], multiple micronutrient supplementation (n = 2 trials) [53,67], zinc supplementation (n = 2 trials) [55,57] or iron supplementation (n = 1 trial) [65] on fluid intelligence composite scores (Table 3 and Figure 4). From the statistical tests of Egger (p = 0.39), there was no evidence of publication bias (Supplementary Figure S6). From the sensitivity analysis, the country-income of the studies did not significantly (p > 0.05) affect child fluid intelligence (Supplementary Figure S7).

![Forest plot for child fluid intelligence outcomes.](image)

**Figure 4.** Forest plot for child fluid intelligence outcomes. The overall effect size was estimated by standardised mean difference (SMD). LCPUFA, long chain polyunsaturated fatty acids; MMN, multiple micronutrient. D+L, random-effects estimate (Der Simonian and Laird method) and I-V, fixed-effects estimate (inverse variance method).

### 3.11. Outcome 5: Global Cognition

Twenty RCTs [46,48,49,51,52,56,57,59–63,65,67,69,72–74,116,125] measured the effect of a nutritional intervention on global cognition. The effect size of multiple time points were combined in Chang et al. [73], Helland et al. [49], Helland et al. [51] and Helland et al. [50]. Li et al. [60] Van Goor et al. [56] Chang et al. [73] and Waber et al. [125] used multiple interventions and one control group, the sample populations were not independent, therefore the effect sizes were combined. McGrath et al. [61] also used multiple independent intervention and control groups, therefore the effect sizes were kept separate. There was no significant effect of multiple micronutrient interventions (n = 6 trials) [60,61,67,73,116,125] LCPUFA interventions (n = 10 trials) [46,49,51,52,56,59,62,63,72,74] iron supplementation (n = 1 trial) [65] or vitamin A supplementation (n = 1 trial) [69] on global cognition composite scores (Table 3 and Figure 5). Hanieh et al. [48] investigated the effect of nutrient supplementation of varying dosage on cognition, iron and folic acid supplements given twice weekly was shown to significantly effect global cognition composite scores in infants at six months. The heterogeneity among the multiple micronutrient intervention trials was significant (p = 0.04). From the statistical tests of Egger (p = 0.82), there was no evidence of publication bias...
(Supplementary Figure S8). From the sensitivity analysis, the country-income of the studies did not significantly ($p > 0.05$) affect child global cognition outcomes (Supplementary Figure S9).

**Figure 5.** Forest plot for child global cognition outcomes. The overall effect size was estimated by standardised mean difference (SMD). LCPUFA, long chain polyunsaturated fatty acids; MMN, multiple micronutrient. D+L, random-effects estimate (Der Simonian and Laird method) and I-V, fixed-effects estimate (inverse variance method).

### 3.12. Outcome 6: Crystallised Intelligence

Eleven RCTs [46,48,52,53,55,57,62,65,67,72,74] measured the effect of a nutritional intervention on crystallised intelligence composite score. Multiple outcomes were combined to create a composite score in Brei et al. [72] (2 outcomes), Caulfield et al. [57] (3 outcomes), Dunstan et al. [46] (4 outcomes) and Mulder et al. [62] (7 outcomes). There was no significant effect of LCPUFA supplementation ($n = 5$ trials) [46,52,62,72,74], multiple micronutrient supplementation [53,67] ($n = 2$ trials), zinc supplementation ($n = 2$ trials) [55,57] or iron supplementation ($n = 1$ trial) [65] on crystallised intelligence composite scores (Table 3 and Figure 6). Hanieh et al. [48] investigated the effect of nutrient supplementation of varying dosage on cognition, no significant effect on child crystallised intelligence composite score was reported. The heterogeneity among the LCPUFA intervention trials was significant ($p = 0.00$). From the statistical tests of Egger ($p = 0.05$), there was no evidence of publication bias, however there is asymmetry in the funnel plot (Figure 7). From the sensitivity analysis, the country-income of the studies did not significantly ($p > 0.05$) affect child crystallised intelligence (Supplementary Figure S10).
**Figure 6.** Forest plot for child crystallised intelligence outcomes. The overall effect size was estimated by standardised mean difference (SMD). LCPUFA, long chain polyunsaturated fatty acids; MMN, multiple micronutrient. D+L, random-effects estimate (Der Simonian and Laird method) and I-V, fixed-effects estimate (inverse variance method).

**Figure 7.** Funnel plot for child crystallized intelligence outcomes with 95% confidence limits. meanSE, mean standard error.
3.13. Outcome 7: Visual Processing

Three RCTs [45,53,55] measured the effect of a nutritional intervention on visual processing. There was no significant effect of choline supplementation \((n = 1 \text{ trial})\) [45], multiple micronutrient supplementation \((n = 1 \text{ trial})\) [53] or zinc supplementation \((n = 1 \text{ trial})\) [55] on visual processing composite scores.

3.14. Outcome 8: Attention

Four RCTs [47,66,74,76] measured the effect of a nutritional intervention on attention. There was no significant effect of multiple or LCPUFA supplementation \((n = 2 \text{ trials})\) [47,74] on attention. Vuori et al. [76] reported that multiple micronutrient supplementation during pregnancy did positively affect attention in infants at 15 days postnatally.

3.15. Outcome 9: Problem Solving

Two RCTs [59,64] measured the effect of a nutritional intervention on the problem solving composite score. There was no significant effect of multiple micronutrient intervention \((n = 1 \text{ trial})\) [64] or LCPUFA intervention \((n = 1 \text{ trial})\) [59] on problem composite scores.

4. Discussion

This systematic review and meta-analysis indicates that nutritional interventions during pregnancy do not significantly impact on child cognitive outcomes. There is some evidence to suggest that LCPUFA supplementation may marginally improve child crystallised intelligence, however the effect was not statistically significant and study heterogeneity among the trials was significant. Although a wide scope of literature was assessed and a rigorous selection methodology was applied, the data included in this systematic review were affected by significant methodological limitations. From the included studies, 79% were exclusively nutrient supplementation trials, the effect of food-based interventions or nutrition counselling alone during pregnancy on child cognition was not well explored. Many studies were small, and attrition rates were high due to the length of follow-up required. Child cognition was a secondary outcome of interest in many studies, even with meta-analysis our findings are likely to be underpowered. This systematic was unable to perform a meta-analysis in specific age ranges due to the lack of data available for children older than 35 months, especially in low and middle income countries. Therefore, it is important that future trials include a longer child follow-up, to explore any possible delayed effects of maternal nutritional interventions on child cognition.

4.1. Quality of Included Studies

Many included studies were considered to be of high quality. While, the Quality Checklist Criteria [36] is designed to assess the methodological quality and reporting within an article, it does not assess the appropriateness of the methodology in addressing the research question. This is a concern considering that many of included RCTs are likely to be at risk of type 2 error, due to the use of secondary data analysis and not powered to detect differences in cognitive outcome as a primary outcome variable. In addition, many post-hoc studies excluded participants after randomisation and in some cases multiple variable adjustments were required to address the significant characteristic differences between the groups, suggesting the assumptions of randomisation had been breached. Most post-hoc studies did not perform intention-to-treat analysis to evaluate the randomisation of the study groups. This approach is likely to introduce bias and systematic error that will reduce the power of the study. Therefore, it is important that future evidence is derived from adequately powered primary studies to reduce the risk of a type 2 error and systematic bias in post-hoc studies and determine whether there are any effects of nutritional interventions during pregnancy on child cognition.
4.2. Measuring Child Cognition

There are many methodological challenges when measuring child cognition using psychological assessment tests. The cognitive performance of infants and children will be affected by their mood, motivation, amount of sleep and personal effort [138]. Environmental factors, including who conducts the cognition tests, who else is present during testing and where the tests are held can also influence a child’s performance ability [138]. In addition, most cognition tests were devised in developed countries, using stimuli relevant to their culture, therefore this stimuli is likely to be unfamiliar or unacceptable to children from developing countries.

Many studies in this review used global development assessments (e.g., Bayley’s Scales of Infant Development) to determine the children’s overall cognitive ability. However, broad measures of cognitive function may only detect highly significant nutritional effects, especially if the study is underpowered [138]. Specific cognitive skills (e.g., attention, memory) that contribute to overall function are regulated by distinct neural mechanisms, that may be differentially affected by nutritional manipulations [139]. Therefore, Cognitive Assessment Tests that analyse specific skills, may be more sensitive to detect subtle effects of nutritional interventions on cognition. For this reason, future trials may be improved by the use of assessment tests that measure specific cognitive skills, as well as global measures of cognition. The choice of specific cognition test should be based on the hypothesised neural mechanisms that are expected to be altered by nutrients.

To determine cognitive function later in childhood and adolescence, future studies should consider linking and analysing school performance data. Pilot data should be used to determine the appropriateness of the sample size and ensure that the study is adequately powered. This approach also has the advantage of the outcome assessors being blinded to the randomisation.

4.3. Nutrition Intervention Design

Due to methodological and ethical challenges, most trials analysed the effects of nutrient supplementation during pregnancy on child cognition rather than using food-based interventions. This may reflect the perception that obtaining adherence to food-based interventions is a challenge and therefore nutrient supplementation is a more feasible strategy/study design. However, performing a meta-analysis for the supplementation trials proved to be difficult because many studies used a supplemented control group rather than an unsupplemented placebo group. Consequently, only the effect of the additional nutrient supplemented in the intervention group compared to the control group could be accessed. In Australia, evidence-based guidelines recommend the use of folic acid and iodine supplementation during pregnancy, therefore it is unethical to use an unsupplemented control group within this population. Li et al. [60] and Caulfield et al. [57] attempted to address this dilemma by providing the same dosage of folic acid across all study groups while women in the intervention groups received a multiple micronutrient supplement (including folic acid) or folic acid with iron. In addition, 88% of the studies conducted in high-income countries provided a single nutrient supplement to women during pregnancy. Since the effect of a single nutrient supplement on child cognition is likely to be modest, especially in a population that is well-nourished, this is likely to explain why a significant summary effect was not found. While, 88% of middle and low-income country studies, provided multiple micronutrient supplementation in a population that is likely to be undernourished, but no significant effect on child cognition was found. However, since child cognition assessment test were validated in a western population, this may have resulted in summary effect that was underestimated.

By convention, many randomised control trials analyse the effect of nutrition on cognition by distilling complex dietary patterns into a single agent using a nutrient supplement. A single nutrient supplement is However, the bioavailability and efficacy of nutrient supplements can differ compared to nutrients contained in food [140]. Nutrient bioavailability is affected by the specific nutrient, absorptive capacity of intestinal mucosa, physiological states (e.g., pregnancy) and nutrient-nutrient interactions [140]. For example, the bioavailability of inorganic iron is enhanced by
an interaction with vitamin C and increasing the bioavailability of iron can decrease magnesium and calcium absorption [140]. Huang et al. [141] reported that the bioavailability of natural vitamin was approximately 50 percent greater than synthetic vitamin E (α-tocopherol) in adults. In addition, synthetic vitamin E supplementation was also associated with a significant reduction in the bioavailability of other forms of vitamin E [141]. This trial highlights the limitations of analysing the effects of nutrition via a single agent without considering the biochemical pathways in which nutrients interact that is likely to be important for cognitive development [142–144]. Given the limited evidence available, there is a clear need for trials that are designed from a sound biologically plausible basis.

4.4. Public Health Implications

Based on the meta-analyses presented, there is a lack of robust evidence on which to base recommendations for maternal diet or for the use of nutrient supplements as a strategy to enhance cognition in children. Given that the synergistic effect of micronutrients in metabolic pathways is important for enhancing cognitive development of children [145], it is important that future trials focus on food-based interventions that aim to improve the overall diet quality of pregnant women. In developing countries, pregnant women are more likely to experience protein-energy malnutrition, as well as multiple micronutrient deficiencies in comparison to high-income countries [146]. Therefore, further evidence is required to determine whether protein and energy as well as micronutrients interventions are effective for improving child cognition in the long-term in developing countries.

5. Conclusions

In conclusion, this systematic review suggests that the infant and child cognitive outcomes were not significantly improved by nutritional interventions (dietary counselling, education, nutrition supplementation, fortified foods and/or whole foods). However, the possible association between LCPUFA supplementation and child crystallised intelligence warrants further investigation in high quality RCTs powered to detect difference in cognitive variables as the primary outcome. Further evidence from primary research is required to test whether maternal nutritional interventions during pregnancy, in particular food-based interventions, are beneficial for improving child cognition. This review emphasises the need for well-designed pilot studies to inform adequately powered RCTs and data linkage studies. This evidence is important because gains in child cognition are likely to have extended reach to other areas of public health and carry economic benefit.

Supplementary Materials: The following are available online at www.mdpi.com/2072-6643/9/11/1265/s1, Table S1: Search strategy used for systematic review; Table S2: Computing a combined effect across outcomes; Table S3: Summary of quality assessment; Table S4: Summary of the nutritional interventions for included studies; Table S5: Cognitive Assessment Tests; Figure S1: Review protocol; Figure S2: Funnel plot: behavior; Figure S3: Sensitivity analysis: behavior; Figure S4: Funnel plot: motor skills; Figure S5: Sensitivity analysis: motor skills; Figure S6: Funnel plot: fluid intelligence; Figure S7: Sensitivity analysis: fluid intelligence; Figure S8: Funnel plot: global cognition; Figure S9: Sensitivity analysis: global cognition; Figure S10: Sensitivity analysis: crystallised intelligence.

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Author Contributions: R.M.T., A.J.H., A.B., C.E.C., R.S. were involved in the research design. R.M.T. and S.F. conducted the research including literature search, quality assessment and data extraction. A.J.H. assisted with publication eligibility and assessing methodological quality. R.M.T., A.B. and T.-J.E. performed the statistical analysis. R.M.T. drafted the paper and was primary responsible for the final content. All authors provided constructive feedback and approved the final manuscript.

Conflicts of Interest: The authors declare no conflict of interest.
Abbreviations

CHC  Cattell–Horn–Carroll
CI  confidence interval
DHA  docosahexaenoic acid
HIV-1  human immunodeficiency virus type 1
LCPUFA  long chain polyunsaturated fatty acids
OECD  Organisation for Economic Co-operation and Development
PRISMA  Preferred Reporting Items for Systematic Review Meta-Analyses
PRISMA randomised controlled trials
SMD  standardised mean difference

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