



SYSTEMATIC REVIEW

Effects of vitamin D deficiency on neurobehavioural outcomes in children: a systematic review [version 1; peer review: 2 approved]

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Abstract




Introduction: Vitamin D plays an important role in brain development in experimental studies; however, the effect of vitamin D deficiency on child development remains inadequately characterized. We aimed to estimate the effects of vitamin D deficiency on neurobehavioural outcomes in children up to 18 years of age.

Methods: We searched PubMed, EMBASE, PsycINFO, Scopus, Cochrane Library, Web of Science and Open Grey for published studies up to 10th January 2020. We included all studies that assessed the effects of maternal or child vitamin D status or vitamin D supplementation on neurobehavioural outcomes in children. Study findings were synthesized qualitatively as the high level of heterogeneity in study populations and methodologies precluded a quantitative meta-analysis.

Results: Our search identified 5,633 studies, of which 32 studies with 31,445 participants from 18 countries were included in the systematic review. Of the studies identified, two were randomized controlled trials (RCTs) of vitamin D supplementation in children, while 30 were observational. One RCT (n=55) reported a beneficial effect of

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supplementation with lower doses compared to higher doses of vitamin D on motor development while the other RCT (n=70) found no beneficial effect of vitamin D supplementation on cognition in extremely preterm infants. Twelve mother-child studies (n=17,136) and five studies in children (n=1,091) reported an association between low maternal or child 25-hydroxyvitamin D levels and impaired neurobehavioural outcomes in children, while 15 mother-child studies (n=20,778) and eight studies in children (n=7,496) reported no association.

Conclusions: Although animal studies point to an effect of vitamin D deficiency on brain development, there are few studies on the effects of vitamin D deficiency on neurobehavioural outcomes in children and their findings are inconsistent. There is a need for well-conducted, adequately powered studies to further determine these effects in children.

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Keywords

Vitamin D deficiency, neurobehavioural outcomes, brain development, children, cognitive, motor, language, development



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Introduction

Impaired neurobehavioural outcomes are common among children worldwide, especially in low- and middle-income countries (LMICs)¹. Approximately 81 million children below the age of five years have impaired cognitive and socioemotional development in LMICs globally and 44% of these children live in sub-Saharan Africa². Impaired child development is associated with poor educational achievement and subsequent poverty and poor health outcomes³. Risk factors for impaired development include home environments lacking in stimulation, infections, malnutrition and micronutrient deficiencies^{3,4}. Vitamin D deficiency and/or insufficiency is an important public health concern that affects approximately one billion people globally^{5,6}. Evidence suggests that vitamin D deficiency is particularly widespread among children worldwide⁷⁻⁹.

Vitamin D may be important for brain development, especially in the early years of life when the brain is developing rapidly

and is sensitive to nutrient deficiencies¹⁰. Animal and *in vitro* studies provide consistent evidence for an important role of vitamin D in brain development. Evidence includes indication of vitamin D signalling in the brain, including presence of vitamin D receptors (VDR), metabolites and enzymes responsible for vitamin D activation and inactivation^{11,12}. Furthermore, animal studies show that vitamin D is important for neuronal differentiation and reducing apoptosis in the hippocampus, an area that is involved in language and memory^{13,14}. Vitamin D is also important for neuroprotection and anti-inflammatory effects in the brain^{15,16} (Figure 1). However, there are few epidemiological studies that have evaluated the effects of maternal or child vitamin D status on neurobehavioural outcomes.

In this systematic literature review, our objective was to estimate the effects of vitamin D deficiency and/or vitamin D supplementation on neurobehavioural outcomes in children. The outcomes of interest included cognitive and motor development, behaviour, intelligence, memory, school achievement and language.

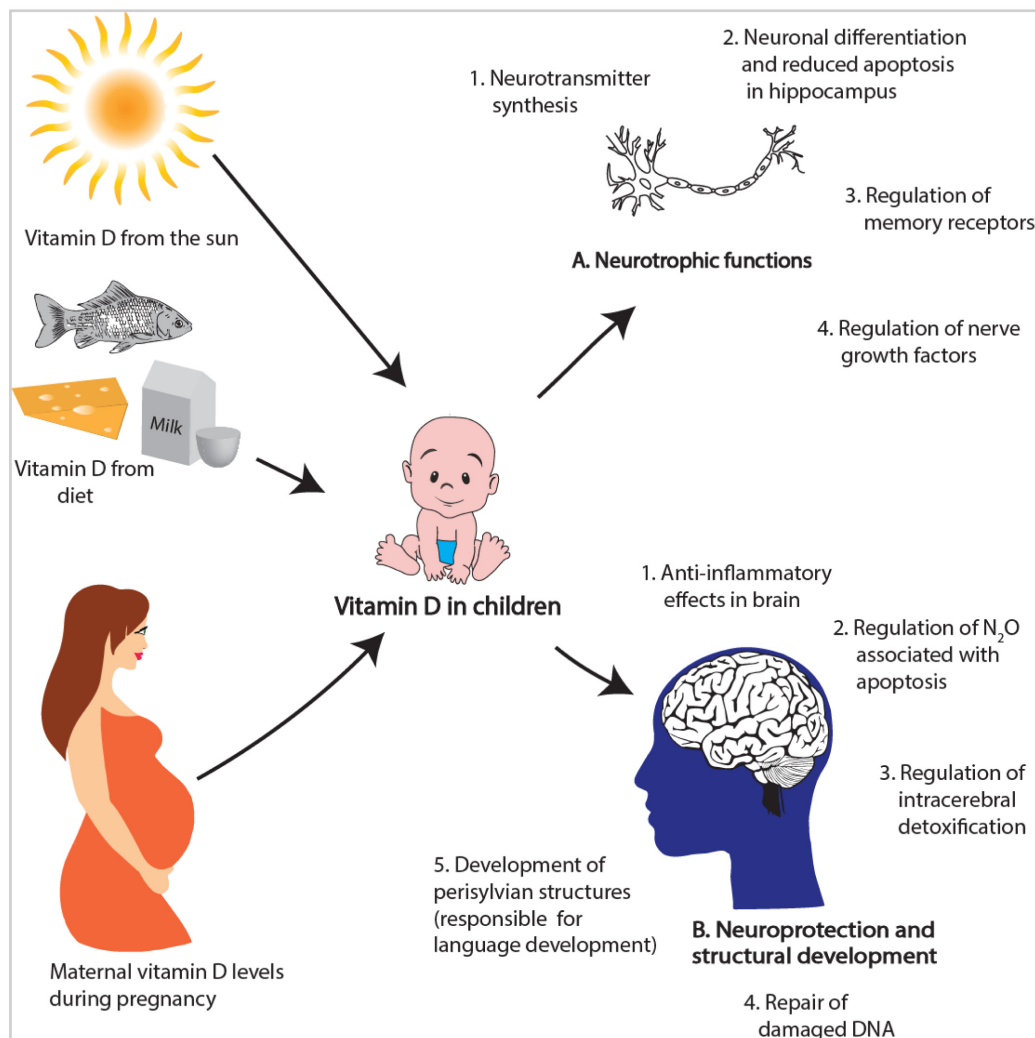


Figure 1. Factors influencing vitamin D status in children and potential roles of vitamin D in brain development¹¹⁻¹⁹.

We further discuss the possible mechanisms by which vitamin D deficiency might influence child development from *in vitro* and animal studies.

Methods

We used the Preferred Reporting Items for Systematic Review and Meta-analysis (PRISMA) guidelines^{20,21} and the Centre for Reviews and Dissemination (CRD) recommendations for undertaking reviews in healthcare²². We registered the protocol on the PROSPERO database (registration number [CRD42018087619](#)).

Search strategy and eligibility criteria

We searched PubMed, EMBASE, PsycINFO, Scopus, Cochrane Library and Web of Science for relevant studies published up to 10th January 2020 without geographical or language limitations. We also searched the Open Grey database for unpublished studies. We further scanned reference lists of identified studies and previous systematic reviews for relevant studies. We applied a search strategy combining Medical Subject Heading terms for [vitamin D] AND [neurobehavioural outcomes] AND [children] and modified the search strategy as appropriate for each of the specific databases (*Extended data, file 1*)²¹.

We included studies if they (i) involved participants below 18 years of age or mother-child pairs; (ii) involved supplementation with vitamin D or measured 25(OH)D levels in children or in pregnant mothers; (iii) measured neurobehavioural outcomes in children including cognitive or motor development, intelligence quotient, attention, behaviour, school achievement or language. We excluded studies that were in older populations or only studied other outcomes apart from neurobehavioural outcomes. We also excluded reviews and case studies, commentaries, study protocols, reports and letters.

Study selection, data extraction and quality appraisal

First, we screened titles and abstracts of all identified studies against the inclusion criteria. We then reviewed full texts for potentially relevant articles to determine eligibility for inclusion. We extracted data into an Excel spreadsheet with a list of variables determined *a priori* by the authors. Variables that were extracted for the review included the first author's name and year of publication, study design, country, sample size, age at vitamin D and neurobehavioural assessment, neurobehavioural outcomes and tools used for evaluation, 25(OH)D cut-offs, and findings of the study. Study selection and data extraction was carried out by two reviewers independently (AMM and RMM) and then compared. Disagreements between reviewers were resolved through discussion. For this review, we defined severe vitamin D deficiency, vitamin D deficiency, insufficiency and sufficiency as 25(OH)D levels <25 nmol/L, 25–50 nmol/L, 50–75 nmol/L and >75 nmol/L respectively according to the Endocrine Society Practice Guidelines²³.

We used the Cochrane handbook criteria to assess the risk of bias for the randomized controlled trials (RCTs)²⁴ and the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) checklist to assess for the quality of observational studies included in the review²⁵.

Synthesis of included articles

The large degree of diversity in the study populations and methods necessitated qualitative synthesis for the studies. We grouped and discussed the findings by each neurobehavioural outcome examined and used tables to present a summary of the study characteristics and findings. For consistency, we converted all units of 25(OH)D measurements into nmol/L.

Results

Our search identified 5,633 articles and a further 18 articles were identified from scanning through reference lists of relevant articles (*Figure 2*). We removed 1,557 duplicate articles. After screening abstracts and titles, we excluded 4,003 articles that were irrelevant to our study and retained 91 papers for full review. We excluded 29 studies because the outcomes of interest were physical growth, psychiatric disorders or infections, eight because they assessed coverage of vitamin D supplementation programmes and two because the effect of vitamin D status on child development was not measured separately from the effects of other micronutrients. We excluded an additional 20 articles as they were reviews, case studies, commentaries or study protocols.

Study characteristics and outcomes

We included two vitamin D supplementation trials and 30 observational studies published between 2007 and 2019 in this review. Of the 30 observational studies, 19 evaluated the effect of maternal vitamin D status during pregnancy and 11 the effect of a child's vitamin D status on neurobehavioural outcomes in children. Of the 11 studies in children, three were cohort studies, two were case-control studies and six were cross-sectional studies; 19 studies were carried out in high-income countries and 13 in low and middle-income countries. The sample sizes varied across the studies, from 45 to 7065 children (*Figure 3*). The studies evaluated a range of neurobehavioural outcomes using a variety of assessment tools and varying definitions for vitamin D status. The characteristics and results of the included studies are shown in *Table 1*. The RCTs showed a low risk of bias (*Extended data file 2*)²¹, although the small sample sizes ($n=55$ and $n=70$) are likely to have limited their power and external validity. All observational studies adjusted for multiple potential confounding factors except one study in which it was unclear whether they adjusted for any confounders²⁶. Limitations of the observational studies included a lack of description about loss to follow-up, and many did not report the reasons for non-participation and the number of participants missing data for each variable (*Extended data, file 3*)²¹.

Cognitive development

A single small trial ($n=70$) in the USA reported that supplementation with 200 international units (IU) or 800 IU of vitamin D from birth to 28 days in extremely preterm neonates was not beneficial for cognitive development at 22 to 24 months of age²⁷.

A total of 15 studies, including 19,810 mother-child pairs evaluated the effect of vitamin D status during pregnancy on cognitive development in children. Of these, four mother-child studies ($n=6,239$) found a beneficial effect of higher maternal 25(OH)D levels on

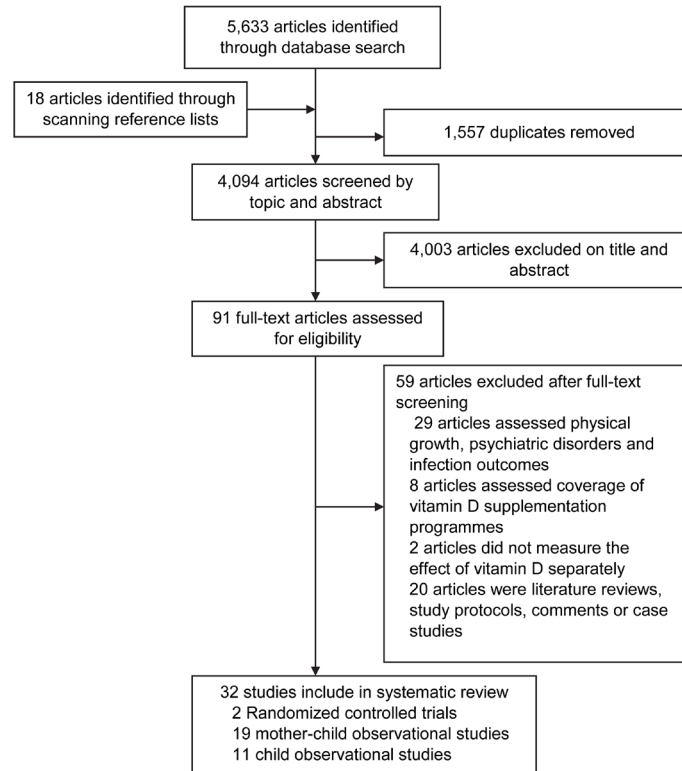


Figure 2. PRISMA flow chart showing the selection process for studies included in the review.

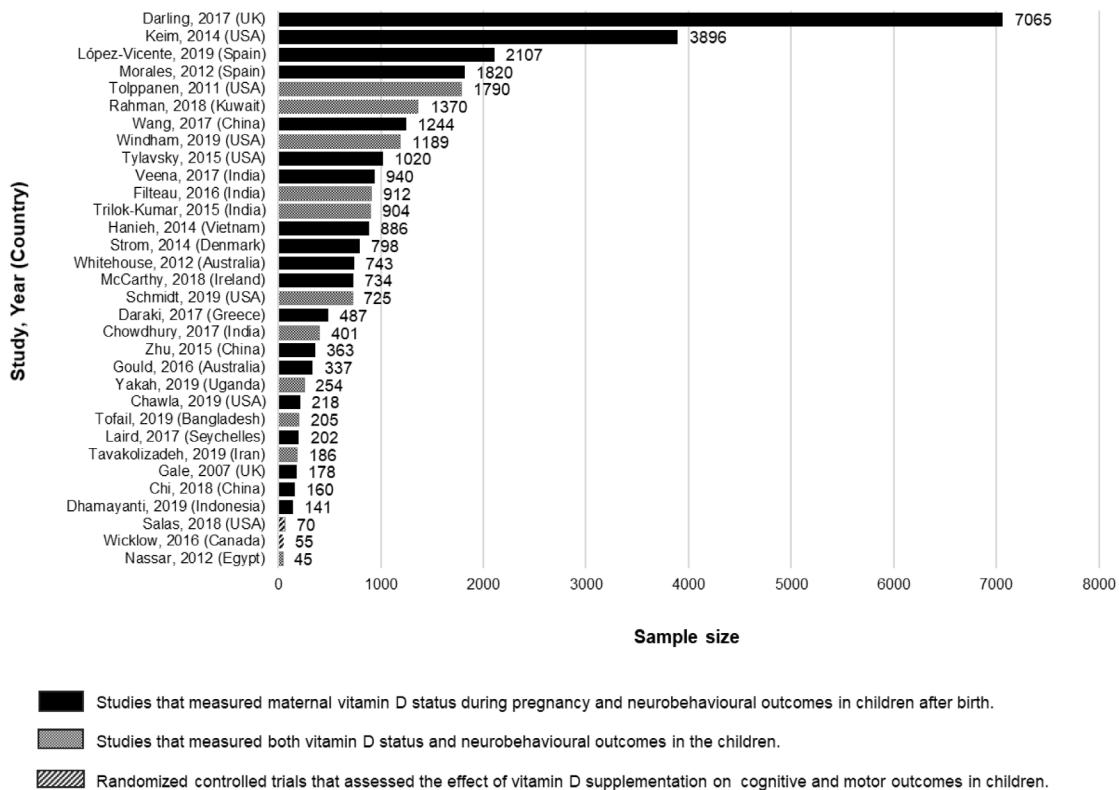


Figure 3. Sample sizes and types of studies included in the review.

Table 1. Summary of studies assessing the effect of vitamin D status on child development: characteristics and findings.

Author, year (Country)	Participants	Age at vitamin D measurement (gestational or in child)	Age at neuro-assessment in child	Study design	Domain (tool)	Definition for vitamin D status	Results
Vitamin D supplementation randomized controlled trials							
Salas, 2018 (USA) ²⁷	70 extremely preterm (<28 gestational weeks) infants	Two time-points: At birth and 28 days	22–26 months	RCT: 3 arms given 200 IU (n=22), 800 IU (n=20) or placebo (n=28) for <28 days	Cognitive (BSID III)	VDD: <30 and <50 nmol/L	Supplementation with vitamin D not beneficial for cognitive development.
Wicklow, 2016 (Canada) ²⁸	55 infants	3 and 6 months	6 months	RCT: 3 arms given 400 IU (n=19), 800 IU (n=18) or 1,200 IU (n=18) vitamin D3/day for 5.5 months	Motor (AIMS)	VDS: >75 nmol/L	Infants given 400 IU of vitamin D had higher motor scores (percentile scores (SD)=44.7 (27.7)) compared to those who received 800 IU and 1,200 IU (23.6 (13.9) and 25.6 (17.7) respectively, p <0.05).
Studies that measured maternal 25(OH)D levels during pregnancy and neurobehavioural outcomes in children after delivery							
López-Vicente, 2019 (Spain) ²⁹	2107 mother-child pairs	Mean (SD) 13.3 (1 st to 2 nd trimester)	4 time-points: 5.8, 14 and 18 years	Cohort	Social competence Total behavioural problems (CBCL, SDQ)	VDD: <50 nmol/L VDI: 50–74.9 nmol/L VDS: ≥75 nmol/L	Maternal 25(OH)D levels >75 nmol/L associated with improved social competence scores compared to levels <50 nmol/L (β =2.14, 95% CI: 0.47, 3.82). No association with total behavioural problems.
Dhamayanti, 2019 (Indonesia) ³⁰	141 mother-child pairs	11 to 14 weeks (1 st trimester)	3 time-points: 3, 6 and 12 months	Cohort	Motor Cognition (ASQ-3)	VDD: <25 nmol/L	Maternal 25(OH)D levels <25 nmol/L associated with impaired motor development at 3 and 12 months. No association with cognition.
Chawla, 2019 (USA) ³¹	218 mother-child pairs	1 st and 2 nd trimester	12 to 24 months	Cohort	Behaviour (ITSEA questionnaire)	25(OH)D levels were divided into 4 quartiles (exact values unspecified)	Lower maternal quartiles of 25(OH)D associated with less favourable behavioural scores among white and Hispanic, but not black infants.
Chi, 2018 (China) ³²	160 mother-child pairs	28 weeks (2 nd trimester)	6 months	Cohort	Mental Psychomotor (BSID-II)	VDD: <50 nmol/L VDS: >50 nmol/L	Maternal 25(OH)D <50 nmol/L associated with lower mental (OR=2.77, 95% CI: 1.44, 5.35, p=0.002) and psychomotor scores (OR=2.08, 95% CI: 1.07, 4.04, p=0.032).

Author, year (Country)	Participants	Age at vitamin D measurement (gestational or in child)	Age at neuro-assessment in child	Study design	Domain (tool)	Definition for vitamin D status	Results
McCarthy, 2018 (Ireland) ³³	734 mother-child pairs	Two time-points: 15 weeks (2 nd trimester) and cord blood	5 years	Cohort	IQ Behaviour (KBIT-2, CBCL)	VDD: <30 nmol/L VDI: 30–50 nmol/L VDS: ≥ 50 nmol/L	No association with IQ or behaviour.
Darling, 2017 (UK) ³⁴	7065 mother-child pairs	Median 29.6 (12.7–33.3) weeks (1 st to 3 rd trimester)	Motor, socio-development at 6 to 42 months Behaviour at 7 years IQ at 8 years Reading ability at 9 years	Cohort	Motor Socio-development and communication Behaviour IQ Reading ability (SDQ, WISC)	VDD: <50 nmol/L VDS: >50 nmol/L	Maternal 25(OH)D <50 nmol/L associated with lower gross-motor (OR=1.20, 95% CI: 1.03, 1.40, p=0.02), fine-motor (OR=1.23, 95% CI: 1.05, 1.44, p=0.01) and social development (OR=1.20, 95% CI: 1.01, 1.41, p=0.04). No association with IQ or reading ability.
Veena, 2017 (India) ³⁵	940 mother-child pairs	28 to 32 weeks (3 rd trimester)	9–14 years	Cohort	Cognitive Memory Language (KABC-II, WISC)	VDD: <50 nmol/L VDS: >50 nmol/L	No association with cognitive, memory or language development.
Wang, 2017 (China) ³⁶	1244 mother-child pairs	Cord blood	2 years	Cohort	Motor Cognitive Behaviour (ASQ)	VDD: <50 nmol/L VDI: 50–74.9 nmol/L VDS: ≥75 nmol/L	No association with motor, cognitive or behavioural development.
Laird, 2017 (Seychelles) ³⁷	202 mother-child pairs	Maternal blood samples collected 1 day after delivery	5 years	Cohort	Cognitive Behaviour Language (FT, PLS, WSAT, CBCL, KBIT)	VDS: ≥50 nmol/L VDI: 30–49.9 nmol/L VDD: <30 nmol/L	No association with cognitive, language or behavioural development.
Daraki, 2017 (Greece) ³⁸	487 mother-child pairs	13±2.4 weeks (1 st trimester)	4 years	Cohort	Cognitive Motor Behaviour (MSCA, SDQ)	Tertiles: <38.4 nmol/l; 38.4–50.7 nmol/l; >50.7 nmol/l.	Maternal 25(OH)D levels >50.7 nmol/l associated with higher behavioural scores compared to levels <38.4 nmol/l. No association with cognitive or motor development.

Author, year (Country)	Participants	Age at vitamin D measurement (gestational or in child)	Age at neuro-assessment in child	Study design	Domain (tool)	Definition for vitamin D status	Results
Gould, 2016 (Australia) ³⁹	337 mother-child pairs	Cord blood	Cognitive, motor, language, behaviour and socio-emotional at 18 months Language at 4 years	Cohort	Cognitive Motor Behaviour Language (BSID-III, DAS II, CELF-P2)	VDD: <25 nmol/L VDI: 25–50 nmol/L VDS: >50 nmol/L	Each 10 nmol/L increase in cord blood 25(OH)D associated with an increase in mean language scores of 0.60 points at 18 months (95% CI: 0.04, 1.17, p=0.04) and 0.68 points at 4 years (95% CI: 0.07, 1.29, p=0.03). No association with cognitive, motor, behaviour or socio-emotional development.
Zhu, 2015 (China) ⁴⁰	363 mother-child pairs	Cord blood	16–18 months	Cohort	Mental Psychomotor (BSID-CR)	Quintiles: 5.56–20.8, 20.9–30.9, 31.0–39.8, 39.9–51.0, 51.3–111 nmol/L	Cord blood 25(OH)D levels of 5.56–20.8 nmol/L associated with a reduction in mental and psychomotor scores (-7.60, 95% CI: -12.4, -2.82; p=0.002 and -8.04, 95% CI: -12.9, -3.11; p=0.001 points respectively) compared to the reference category (levels 39.9–51.0 nmol/L). Levels of 51.3 to 111 nmol/L associated with a reduction in psychomotor scores -12.3 (95% CI: -17.9, -6.67; p<0.001) points) compared to the reference category.
Tylavsky, 2015 (USA) ⁴¹	1020 mother-child pairs	2 nd trimester	2 years	Cohort	Cognitive Language (BSID III)	Categories: <49.9, 50.00–74.9, ≥75 nmol/L	Each 25 nmol/L increase in maternal 25(OH)D was associated with an increase in scaled receptive language scores of 0.24 points (p=0.017). No association with scaled cognitive or expressive language scores.
Hanieh, 2014 (Vietnam) ⁴²	886 mother-child pairs	32 weeks (3 rd trimester)	6 months	Cohort	Cognitive Motor Behaviour Language (BSID III)	VDD: <37.5nmol/L VDI: ≥37.5–75 nmol/L VDS: ≥75 nmol/L	Maternal 25(OH)D <37.5 nmol/L associated with a reduction in language scores (mean difference: -3.48; 95% CI: -5.52, -1.44). No association with cognitive, motor, or behaviour development.
Keim, 2014 (USA) ⁴³	3896 mother-child pairs	Two time-points: ≤26 weeks (2 nd trimester) and cord blood	Cognitive and motor at 8 months IQ and behaviour at 4 and 7 years School achievement at 7 years	Cohort	IQ Motor Behaviour School achievement (BSID, SBIS, WISC, WRAT)	Categories: <25, 25–50, 50–75, ≥75 nmol/L	Each 5 nmol/L increase in both maternal and cord blood 25(OH)D associated with a slight increase in IQ scores of 0.10 points (95% CI: 0.00, 0.19) at 7 years of age. No association with motor, school achievement or behaviour development.

Author, year (Country)	Participants	Age at vitamin D measurement (gestational or in child)	Age at neuro-assessment in child	Study design	Domain (tool)	Definition for vitamin D status	Results
Strøm, 2014 (Denmark) ⁴⁴	798 mother-child pairs	30 weeks (2 nd trimester)	15–16 years	Cohort	School achievement (Exam scores)	VDD: <50 nmol/L VDS: ≥50–75 nmol/L Optimal levels: ≥75–125 nmol/L	No association with school achievement.
Morales, 2012 (Spain) ⁴⁵	1820 mother-child pairs	11.6% assessed in 1 st trimester, 88% in 2 nd trimester and <1% in 3 rd trimester	11–23 months	Cohort	Mental Psychomotor (BSID)	VDD: <50 nmol/L VDS: >75 nmol/L	Each 25 nmol/L increase in maternal 25(OH)D associated with 0.79 (95% CI: 0.14, 1.45) and 0.88 (95% CI: 0.22, 1.54) points increase in mental and psychomotor scores, respectively.
Whitehouse, 2012 (Australia) ⁴⁶	743 mother-child pairs	18 weeks (2 nd trimester)	2 years	Cohort	Language Behaviour (CBCL, PPVT-R)	VDD: <46 nmol/L	Maternal 25(OH)D levels <46 nmol/L associated with impaired language development compared to levels >72 nmol/L (OR=1.97, 95% CI: 1.00, 3.92, p<0.05). No association with behavioural development.
Gale, 2007 (UK) ⁴⁷	178 mother-child pairs	28–42 weeks (3 rd trimester)	9 years	Cohort	Cognitive (WASI)	VDS: >50 nmol/L VDI: 27.5–50 nmol/L VDD: <27.5 nmol/L	No association with cognitive development.
Studies that measured 25(OH)D levels and neurobehavioural outcomes in children							
Tofail, 2019 (Bangladesh) ⁴⁸	205 children	7 months	6–8 months	Cross-sectional	Cognitive Motor Language Behaviour (BSID III, parental reports)	VDS≥50 nmol/L VDD<50 nmol/L	Children with VDD had lower behavioural scores (activity: $\beta \pm SE$ 1.64±0.78, 95% CI: 0.10, 3.18, p=0.037) and soothability (2.02±0.70, 0.64, 3.41, p=0.004) compared to the VDS group. No association with cognitive, motor or language development.
Yakah, 2019 (Uganda) ⁴⁹	254 children (3 groups: perinatally HIV-infected (PHIV), exposed-uninfected (HEU), or unexposed-uninfected (HUU))	6–10 years	3 time-points: 6–10, 6.5–10.5 and 7–11 years	Cohort	Behaviour BASC-3	VDD: <50 nmol/L VDI: 50–62 nmol/L VDS: >62 nmol/L	VDD associated with poor behavioural development in perinatally HIV exposed but not in uninfected and HIV unexposed uninfected.
Windham, 2019 (USA) ⁵⁰	1189 children (753 cases with developmental disability and 436 controls from the general population)	New-born stored blood spots	4.5–9 years	Case-control	IQ (Intellectual disability)	VDS: ≥ 75nmol/L VDI: 50–74nmol/L VDD: <50 nmol/L	No association with IQ.

Author, year (Country)	Participants	Age at vitamin D measurement (gestational or in child)	Age at neuro-assessment in child	Study design	Domain (tool)	Definition for vitamin D status	Results
Schmidt, 2019 (USA) ⁵¹	725 children (491 cases with developmental disability and 234 controls from the general population)	New-born stored blood spots	24–60 months	Case-control	Cognitive (MSEL, VABS)	VDS: ≥ 75 nmol/L VDI: 50–75 nmol/L VDD: < 50 nmol/L	No association with cognitive development.
Tavakolizadeh, 2019 (Iran) ³⁶	186 children	12 months	12 months	Cross-sectional	Gross motor (Assessed by paediatrician based on achievement of pre-selected milestones)	VDS: > 75 nmol/L VDI: 50–74 nmol/L VDD: < 50 nmol/L	VDD associated with impaired gross motor ability compared to VDS (OR=3.9, 95% CI: 1.9, 8.4, $p < 0.001$).
Rahman, 2018 (Kuwait) ⁵²	1,370 adolescents	11–16 years	11–16 years	Cross-sectional	Cognitive (RSPM), School achievement (Exam scores)	VDS: ≥ 75 nmol/L VDI: 50–75 nmol/L VDD: < 50 nmol/L Severe VDD: < 25 nmol/L	No association with cognitive development or school achievement.
Chowdhury, 2017 (India) ⁵³	401 pre-school children	6–30 months	12–36 months	Cohort	Cognitive Motor Personal-social (ASQ-3)	VDD: < 25 nmol/L	No association with cognitive, motor or personal social development.
Fliteau, 2016 (India) ⁵⁴	912 school children	5 years	5 years	Cross-sectional	Gross motor (ASQ-3)	VDD: < 25 nmol/L Borderline: 25–50 nmol/L VDS: > 50 nmol/L	No association with gross motor development.
Trilok-Kumar, 2015 (India) ⁵⁵	904 children	2 time-points: 6 months, 3–6 years	3–6 years	Cohort (follow up of a vitamin D supplementation trial in infancy)	Gross motor (ASQ-2)	VDD: < 50 nmol/L	No association with gross motor development.
Nassar, 2012 (Egypt) ⁵⁶	45 school children	School age	School age	Cross-sectional	Cognitive (WISC, BVRT)	VDD: < 27.5 nmol/L VDI: 27.5–37.5 nmol/L VDS: 37.5–77.5 nmol/L	VDS associated with improved cognitive and school achievement compared to VDD.
Tolppanen, 2011 (USA) ⁵⁷	1790 adolescents	12–16.9 years	12–16.9 years	Cross-sectional	Cognitive (WISC-R, WRAT)	Quartiles: 8.7–49.2, 49.4–65.6, 65.9–82.1, 82.4–205.2 nmol/L	No association with cognitive development.

AIMS, Alberta Infant Motor Scale; ASQ-3, Ages and Stages Questionnaire-3; BASC-3, Behavioural Assessment System for Children; BSID, Bayley Scales of Infant Development-China Revision; BSID-III, Bayley Scales of Infant Development-version 3; BVRT, Benton's Visual Retention Test; CBCL, Child Behaviour Checklist; CELF-P2, Clinical Evaluation of Language Fundamentals Preschool; Second Edition; DAS-II, Differential Ability Scales Second Edition; FT, Finger tapping; PLS, Preschool Language Scale-Revised Edition; MSCA, McCarthy Scales of Children's Abilities; WISC, Woodcock-Johnson Scholastic Achievement Test; KBIT, Kaufman Brief Intelligence Test; RSPM, Raven's Standard Progressive Matrices; PPVT-R, Peabody Picture Vocabulary Test-Revised; SBIS, Stanford-Binet Intelligence Scale; WASI, Wechsler Abbreviated Scale of Intelligence; WISC, Wechsler Intelligence Scale for Children; WISC-R, Wechsler Intelligence Scale for Children-revised; WRAT, Wide Range Achievement Test; SDQ, Strengths and Difficulties Questionnaire; KABC-II, Kaufman Assessment Battery for Children-2nd edition; MSEL, Mullen Scales of Early Learning; VABS, Vineland Adaptive Behaviour Scales. IQ, intelligence quotient; IU, international units; VDD, vitamin D deficiency; VDI, vitamin D insufficiency; VDS, vitamin D sufficiency; β , Beta coefficient; SD, standard deviation; SE, standard error.

cognitive development and 11 studies ($n=13,234$) reported no association. In a large study of 3,896 American mother-child pairs, higher maternal 25(OH)D levels were associated with higher intelligence quotient (IQ) scores in children at 7 years of age⁴³. Likewise, in a study of 1,820 Spanish mother-child pairs, maternal 25(OH)D levels >75 nmol/L were associated with higher cognitive scores in children aged 11–23 months compared to maternal 25(OH)D levels <50 nmol/L⁴⁵. Another study of 363 Chinese mother-child pairs reported that cord blood 25(OH)D levels between 5.56 and 20.8 nmol/L were associated with lower cognitive scores in 16–18 month-old children than levels between 39.9 and 51 nmol/L⁴⁰. Similarly, a study of 160 Chinese mother-child pairs found that maternal 25(OH)D levels <50 nmol/L were associated with low cognitive scores in six-month-old children³².

However, a large study of 7,065 mother-child pairs in the UK reported no association between maternal 25(OH)D levels and cognition in eight-year-old children³⁴. Other studies in 1,020 American and 1,244 Chinese mother-child pairs also found no association between maternal 25(OH)D levels during pregnancy and cognition in children aged two years or below^{36,41}. Two mother-child studies in India ($n=940$) and Vietnam ($n=886$) also found no association between maternal 25(OH)D levels during pregnancy and cognition in children aged six months to 14 years^{35,42}. Similarly, in six other studies of 2,079 mother-child pairs, maternal 25(OH)D levels were not associated with cognitive development after birth^{30,33,37–39,47}.

Seven observational studies measured both 25(OH)D levels and cognitive development in children. One small cross-sectional study ($n=45$) in Egyptian school children reported that those with 25(OH)D levels of 37.5–77.5 nmol/L had improved cognitive function compared to those with 25(OH)D levels <27.5 nmol/L⁵⁶. However, a cohort study in 401 Indian children found no association between vitamin D status at six to 30 months of age with cognitive development at 12 to 36 months of age⁵³. Additionally, two case-control studies of 1,244 children with neurodevelopmental disabilities and 670 controls with normal development in the USA reported no association between vitamin D status and cognitive development at two to nine years of age^{50,51}. Likewise, a cross-sectional study of 1,790 American adolescents aged 12 to 16.9 years did not find an association with cognitive development⁵⁷. Similarly, two cross-sectional studies, one of 1,370 Kuwaiti adolescents aged 11 to 16 years old and another of 205 Bangladeshi children aged six to eight months reported no association between vitamin D status and cognitive development^{48,52}.

Motor development

One small vitamin D supplementation trial in 55 Canadian children aged three to six months evaluated the effect of vitamin D in doses of 400 international units (IU), 800 IU and 1200 IU on gross motor skills. The study did not include a placebo arm. They reported that supplementation with 400 IU vitamin D was beneficial for gross motor development in infants compared to supplementation with 800 or 1200 IU of vitamin D²⁸.

A total of 10 studies including 11,503 mother-child pairs examined the association between motor development and vitamin D status. In five mother-child studies ($n=9,549$), high maternal 25(OH)D levels were associated with improved motor development; however, five studies ($n=6,850$) found no association. A large study of 7,065 mother-child pairs in the UK reported a trend of improved gross and fine motor scores in 18–30 month-old children with increasing maternal 25(OH)D levels³⁴. Another study of 1,820 Spanish mother-child pairs, similarly showed that maternal 25(OH)D levels >75 nmol/L were associated with higher motor scores in infancy compared to levels <50 nmol/L⁴⁵. A study of 160 Chinese mother-child pairs showed that vitamin D deficient (25(OH)D levels <50 nmol/L) mothers were more likely to have children with lower psychomotor development scores at six months of age compared to mothers without deficiency³². Another study of 141 Indonesian mother-child pairs reported that maternal 25(OH)D levels <25 nmol/L were associated with impaired motor development in children at three and 12 months of age³⁰. Interestingly, a study of 363 Chinese mother-child pairs reported an inverted u-shaped trend of association with lower motor scores in toddlers that had cord blood 25(OH)D levels between 5.56 and 20.8 nmol/L and between 51.3 and 111 nmol/L compared to the reference category (levels 31.0 to 39.8 nmol/L)⁴⁰. However, two large mother-child studies in the USA ($n=3,896$) and China ($n=1,244$) and smaller studies in Vietnam ($n=886$), Greece ($n=487$) and Australia ($n=337$) reported no association between 25(OH)D levels during pregnancy and motor development in children aged between six months to four years^{36,38,39,42,43}.

Five studies assessed both 25(OH)D levels and motor development in children. A cross-sectional study among 186 Iranian children reported that 25(OH)D levels >75 nmol/L were associated with improved gross motor ability compared to 25(OH)D levels <50 nmol/L at one year of age²⁶. However, a cohort study in 401 Indian children reported no association between vitamin D status measured at six to 30 months and motor development at 12 to 36 months⁵³. Another cohort study in 904 Indian children found that vitamin D status measured at six months and at three to six years was not associated with gross motor development at three to six years of age⁵⁵. Likewise, two cross-sectional studies, one in 912 five-year-old Indian children and another in 205 Bangladeshi infants aged six to eight months reported no association with motor development^{48,54}.

Language development

The association between maternal 25(OH)D levels and language development was evaluated in six studies including a total of 4,128 mother-child pairs. Four studies, including a total of 2,986 mother-child pairs, reported a positive association between high maternal 25(OH)D levels and improved language development, while two other studies including 1,142 mother-child pairs reported no association. A study of 1,020 American mother-child pairs reported that maternal 25(OH)D levels ≥ 75 nmol/L were positively associated with receptive language but not expressive language in two-year-old children⁴¹. Likewise, a study of 886 Vietnamese mother-child pairs reported that

six-month-old infants born to women with 25(OH)D levels <37.5 nmol/L had lower language scores compared to those with 25(OH)D levels ≥ 75 nmol/L⁴². Similarly, in a study involving 743 mother-child pairs, children of mothers with 25(OH)D levels <46 nmol/L were almost two times more likely to have language impairment compared to those of women with 25(OH)D levels >72 nmol/L at two years of age⁴⁶. In another study of 337 Australian mother-child pairs, higher cord blood 25(OH)D levels were associated with an increase in mean language scores at 18 months and four years³⁹. However, two studies of 1,142 mother-child pairs reported no association between vitamin D status and language development in children aged five to 14 years^{35,37}. A single cross-sectional study in 205 Bangladeshi infants aged six to eight months reported a positive association between 25(OH)D levels >50 nmol/L in children and language development compared to levels <50 nmol/L⁴⁸.

Behavioural development

The association between maternal 25(OH)D levels and behavioural development was evaluated in 11 mother-child studies. Four mother-child studies (n=7,770) reported an association between high 25(OH)D levels during pregnancy and improved behavioural development in children; however, eight studies (n=10,149) found no association. A large study of 7,065 mother-child pairs in the UK reported an increasing trend of social development scores with increasing maternal 25(OH)D levels in children at 42 months of age³⁴. Another study of 487 mother-child pairs in Greece reported that maternal 25(OH)D levels >50.7 nmol/L were associated with better behavioural scores compared to 25(OH)D levels <38.4 nmol/L³⁸. Likewise, a study of 218 mother-child pairs in the USA reported that low maternal 25(OH)D levels (unspecified values) were associated with impaired behavioural development among white and Hispanic, but not black, infants at one to two years of age³¹. A study of 2,107 mother-child pairs reported a positive association between maternal 25(OH)D levels >75 nmol/L and improved social competence scores compared to levels <50 nmol/L; however, they found little association between maternal 25(OH)D levels and total behavioural problems²⁹. Two large mother-child studies in the USA (n=3,896) and China (n=1,244) found no association between maternal 25(OH)D levels and behaviour in children aged two to 18 years^{36,43}. Two other studies of 1,080 mother-child pairs in Australia reported no association between maternal 25(OH)D levels and behaviour in 18–24 month-old children^{39,46}. Likewise, three mother-child studies in Vietnam (n=886), Ireland (n=734) and Seychelles (n=202) reported no association between maternal 25(OH)D levels and behaviour in children aged six months to five years^{33,37,42}.

Three studies measured both 25(OH)D levels and behavioural development in children. A cohort study of 254 Ugandan children found an association between 25(OH)D levels <50 nmol/L at enrollment, at six to 10 years of age, and poor behavioural outcomes measured at enrollment, and at six and 12 months after enrollment, among children who were perinatally exposed to HIV but were uninfected and unexposed HIV negative children⁴⁹. A cross-sectional study of 401 Indian

children found that children with 25(OH)D levels <25 nmol/L had lower scores on the personal-social sub-scale at six to 36 months⁵³, and a cross-sectional study in 205 Bangladeshi children aged six to eight months reported an association between 25(OH)D levels <50 nmol/L and impaired parent-reported behaviour⁴⁸.

School achievement

Two mother-child studies and two cross-sectional studies evaluated the effect of vitamin D status on school achievement. The two mother-child studies, including a total of 4,694 mother-child pairs, observed no association between maternal vitamin D status and children's educational achievement^{43,44}. The two cross-sectional studies reported conflicting findings. A study of 45 Egyptian children reported that 25(OH)D levels between 37.5 and 77.5 nmol/L were associated with improved school achievement compared to 25(OH)D levels <27.5 nmol/L⁵⁶, while a study of 1,370 Kuwaiti adolescents reported no association between vitamin D status and school achievement⁵².

Discussion

In this systematic review, we found inconclusive evidence for the effects of maternal or child vitamin D status on neurobehavioural outcomes in children below 18 years of age. A small RCT (n=55) reported that infants given lower vitamin D doses had improved motor development compared to those given higher vitamin D doses suggesting that higher doses of vitamin D might impair motor development. Another RCT of 70 extremely preterm neonates reported no beneficial effect of vitamin D supplementation on cognitive development in infancy. Similarly, evidence from observational studies was conflicting. Twelve studies of 17,136 mother-child pairs and five studies in of 1,091 children reported an association between low maternal or child 25(OH)D and impaired neurobehavioural outcomes in children. However, 15 studies of 20,778 mother-child pairs and eight studies of 7,496 children found no association between vitamin D status and neurobehavioural outcomes.

We found conflicting evidence for the effects of vitamin D status on cognitive development. One small RCT reported no beneficial effect of vitamin D supplementation in very preterm neonates on cognition at two years of age, but this study was limited by the small sample size and the short period of vitamin D supplementation (28 days). Overall, we found little evidence for a detrimental effect of low maternal or child 25(OH)D levels on cognitive development. Lack of association might be explained by the adequate baseline levels of 25(OH)D observed in some of the study populations. Of the 17 observational studies that reported no association, 11 observed maternal or child mean or median 25(OH)D levels >50 nmol/L, while four of the five studies that reported an association observed sub-optimal 25(OH)D levels (mean or median levels <50 nmol/L). Also, it is possible that some of the neurodevelopmental assessment tools were not sensitive enough to detect subtle effects of vitamin D on cognition. For example, Tylavsky *et al.* suggest that the Bailey Scales of Infant

Development used in their study could have failed to detect small changes in recognition memory, symbolic play, acuity, or information processing speed⁴¹. Although there is little evidence in humans, animal and *in vitro* studies suggest that vitamin D influences cognitive development^{58–60}. *In vitro* studies show that vitamin D receptors and cytochrome enzymes responsible for vitamin D metabolism are present in the cortex and other parts of the brain involved in cognition¹¹. Additionally, vitamin D in its active form (1,25(OH)₂D) influences cytokine production which may affect neurotransmission and synaptic plasticity, in turn altering learning processes and cognition as observed in rats⁶¹.

Similarly, findings for an effect of vitamin D status on motor development were mixed. More mother-child studies indicated adverse effects for lower maternal 25(OH)D levels on motor development compared to cross-sectional studies in children. Only one cross-sectional study reported an association between lower 25(OH)D levels in children and motor development but these findings were unadjusted for potential confounders²⁶. Three of the four studies that reported no association between child vitamin D status and motor function used the Ages and Stages Questionnaire-3 to assess gross motor development^{53–55}. This tool was designed to diagnose children with developmental abnormalities and may not be sensitive enough to detect small changes in motor function⁶². How might maternal vitamin D status affect motor development? Maternal vitamin D deficiency has been associated with congenital rickets with apparent foetal bone disease and impaired bone quality postnatally⁶³. One study reported an inverted u-shaped association so that both lower (5.56 to 20.8 nmol/L) and higher (51.3 to 111 nmol/L) maternal 25(OH)D levels were associated with impaired motor development in children⁴⁰. This finding is supported by a small vitamin D supplementation trial in 55 infants, which found that a lower dose of vitamin D supplementation might be more beneficial for gross motor function compared to higher doses of vitamin D²⁸. However, the findings of this trial may be limited by the lack of a control arm that did not receive vitamin D supplementation. It is possible that excess levels of vitamin D may result in toxic and pro-inflammatory effects which could lead to impaired motor development^{64,65} and that an optimal 25(OH)D level may be required. Vitamin D status might also influence motor development through its role in neurotransmission through the dopaminergic system. Data from animal studies show that vitamin D deficiency is associated with alterations in the synthesis and turnover of dopamine, norepinephrine and dihydroxyphenylacetic acid in rats^{17,66,67} and ventral mid-brain dopaminergic neurons have been shown to play a crucial role in the regulation of motor processes⁶⁶.

We found evidence for associations between low vitamin D status and impaired language development in four out of six mother-child studies. In one study, impairment of language development was only observed when 25(OH)D levels fell below 37.5 nmol/L in late pregnancy, suggesting that language impairment is a feature of advanced vitamin D deficiency rather than insufficiency⁴². However, two mother-child studies (n=1,142)

reported no association between maternal vitamin D status and language scores in adolescence, but these studies did not account for the vitamin D status of the adolescents, and it is possible that detrimental effects may have been evident earlier in life, but not in adolescence^{35,37}. A possible mechanism for the observed association could be that *in utero* exposure to low vitamin D concentrations during the second and third trimesters may affect the development of the perisylvian structures, which are responsible for language development in children as reported in structural neuroimaging studies^{18,68}. Additionally, maternal vitamin D deficiency has been shown to alter the brain structure and genes that are involved in neuronal survival and speech and language development (brain-derived neurotrophic factor (*Bdnf*)) and forkhead box protein P2 (*Foxp2*) genes, respectively) in mice¹⁷.

We found mixed evidence for the effect of vitamin D deficiency on behaviour in children. Eight of 11 mother-child studies reported no association between maternal vitamin D status and behaviour. Of these studies, four relied on parent-reported behaviours only, which might be prone to bias, and only three relied on psychologist assessments or well-validated tools in evaluating behaviour. Additionally, it might be difficult to estimate the effect of maternal vitamin D status on behavioural outcomes in childhood since a wide variety of maternal, environmental and genetic risk factors may cause confounding. Animal and *in vitro* studies suggest that vitamin D status might influence behaviour possibly via the anti-inflammatory effects of vitamin D, which protect against maternal immune activation, which may affect foetal brain development and behavioural manifestations⁶⁹. Additionally, vitamin D deficiency is associated with alterations in the synthesis of serotonin, a neurotransmitter that is important for behavioural outcomes⁷⁰.

There may be many explanations for our findings of inconsistent evidence for the effects of hypovitaminosis D on child development. There were substantial differences in study design between studies with some assessing maternal and some children's vitamin D status in relation to child development. Studies also had different inclusion criteria and definitions for vitamin D status and were also adjusted for different confounding variables. Secondly, vitamin D status was measured at varying gestational ages in mothers and different ages in the children. Several studies that measured maternal 25(OH)D in the 2nd trimester, rather than the 3rd trimester, reported an association between low maternal 25(OH)D and low developmental scores in the children^{32,41,43,45,46}, which might suggest a critical window for the impact of low 25(OH)D levels earlier in pregnancy when there is rapid brain development⁷¹. In addition, studies suggest that there might be optimal levels of vitamin D required for development with higher levels also being associated with impaired development^{28,40}. Thirdly, the timing of neurobehavioural assessment varied markedly between studies. For example, several studies that evaluated the effect of maternal 25(OH)D levels on neurobehavioural outcomes in adolescence reported no association. Putative effects might be more likely to be detected in earlier years of life and may also be

influenced by 25(OH)D levels throughout childhood. Studies that measured 25(OH)D levels during pregnancy and child development did not account for the vitamin D status of children at the time of neuro assessment, which may also have influenced neurobehavioural outcomes. The application of findings from experimental studies to human beings may also be limited. For instance, animal experiments can create conditions of extremely low 25(OH)D levels which may not normally be found in humans.

We identified two previous systematic reviews and one meta-analysis of the effects of maternal vitamin D status on neurobehavioural outcomes in children. Consistent with our review, a systematic review of maternal micronutrient status (vitamin D, B12, iron and folate) and carbohydrate, protein and fat intake during pregnancy and cognitive development in childhood found limited and conflicting evidence for the effect of vitamin D status on cognitive development⁷². However, of 38 eligible studies, this review included only three mother-child studies on vitamin D status and cognitive development, also included in our review^{45–47}. Another systematic review of vitamin D status during pregnancy and offspring neurocognitive and psychological outcomes also found contradictory evidence for the effect of vitamin D status on neurobehavioural outcomes and neuropsychological disorders in children⁷³. This review included only 10 human studies, four of which evaluated neuropsychological disorders and six neurobehavioural outcomes, but cognition, motor, language and behavioural outcomes were not evaluated⁷³. Another systematic review and meta-analysis of the neurodevelopmental effects of prenatal vitamin D reported an association between increased maternal 25(OH)D levels and improved cognition and reduced risk of autism and attention deficit and hyperactivity disorder (ADHD)⁷⁴. Unlike our review, they did not include studies that measured vitamin D status of children. Additionally, the findings of the meta-analysis may be limited by the different definitions for vitamin D status used across the included studies⁷⁴.

Strengths and limitations

A key strength of our review is that it was very comprehensive, since we searched seven databases without geographic location, date or language limitations. We included randomized controlled trials and observational studies assessing the effects of maternal and child vitamin D status on neurobehavioural outcomes. Also, we included children up to the age of 18 years, hence capturing child development through to adulthood. There were several limitations in this review. We found substantial heterogeneity in study populations and study methods including timing of vitamin D measurements, definitions for vitamin D status and tools used to measure neurobehavioural outcomes which made comparison of the studies difficult and precluded a quantitative meta-analysis of the studies. There were only two small vitamin D supplementation trials and few studies measured 25(OH)D levels in children. We also found a relatively small number of studies from low-income countries. Evidence for the effects of vitamin D deficiency on neurobehavioural outcomes from studies in high-income countries might not be generalizable to low-income settings, where impaired child development may be more common, and confounding effects might also differ between these settings.

Conclusion

Evidence as to whether prenatal or childhood vitamin D status affects neurobehavioural outcomes in children is limited and inconsistent and further investigation is warranted. For instance, standardized cut-offs for assessing vitamin D deficiency, as well as well-validated and standardized methods of assessing neurobehavioural outcomes would be useful to allow comparison among studies globally. Well-powered randomized controlled trials are required to determine whether vitamin D is beneficial for brain development in children and for optimal dosage and timing of vitamin D supplementation.

Data availability

Underlying data

All data underlying the results are available as part of the article and no additional source data are required.

Extended data

Figshare: Effects of vitamin D deficiency on neurobehavioural outcomes in children: a systematic review-supplementary files. <https://doi.org/10.6084/m9.figshare.11799063.v7>²¹.

This project contains the following extended data:

- Extended datafile 1: Search terms.
- Extended datafile 2. Risk of bias in included randomized controlled trials based on the Cochrane handbook criteria.
- Extended datafile 3. STROBE checklist assessing the quality of observational studies included in the review.

Reporting guidelines

Figshare: PRISMA checklist for 'Effects of vitamin D deficiency on neurobehavioural outcomes in children: a systematic review'. <https://doi.org/10.6084/m9.figshare.11799063.v7>²¹.

Data are available under the terms of the [Creative Commons Attribution 4.0 International license](#) (CC-BY 4.0).

Author information

AMM, SHA and AA conceived the idea of the review and developed the protocol. AMM conducted the literature search, selected studies and extracted relevant information with the guidance of SHA, RMM and AA. AMM, SHA and AA synthesized the data and wrote the first draft of the manuscript. AMM, SHA, AA, RMM, ELW, TNW and AME revised successive drafts of the paper and approved the final version.

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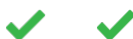
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Mairead Kiely 

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This is a comprehensive SLR of vitamin D status and neurological development in children.

The review includes both trials, of which there were only 2 (both very small), mother-infant prospective cohort studies and cross-sectional studies. Implementation of the search strategy & review are clearly and transparently reported.

The results are logically described and well-written. This is a highly heterogenous dataset describing complex outcomes and the authors treated the results objectively.

A couple of points:

- The Salas paper on preterm infants should not be included as preterm infants are a different and high-risk subgroup from a developmental perspective.
- Although the authors correctly point out the deficits in the trial data, I thought that the discussion could have appraised the evidence basis more from the prospective cohorts from a vitamin D perspective; for example were gold standard methods of 25(OH)D analysis used? Were appropriate methods used for cord blood sampling and analysis?
- Could some sensitivity analysis have been conducted on the basis of 25(OH)D concentrations reported given the widely variant definitions of "vitamin D deficiency"?
- Were the developmental assessments led by clinical psychologists within a MDT?
- Title: Currently suggestive rather than descriptive - Suggest amendment to: "Inconclusive associations between vitamin D status during pregnancy or at birth and neuro-behavioural outcomes in children: a systematic review."

Are the rationale for, and objectives of, the Systematic Review clearly stated?

Yes

Are sufficient details of the methods and analysis provided to allow replication by others?

Yes

Is the statistical analysis and its interpretation appropriate?

Yes

Are the conclusions drawn adequately supported by the results presented in the review?

Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Nutrition; Vitamin D; Maternal & Child nutrition and development.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Reviewer Report 23 March 2020

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Andrew S. Day

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This MS focuses on the impact of vitamin D upon child development, using a systematic approach.

Overall comments:

- The MS is well written and easy to follow. The objectives are clear and the conclusions well presented.
- The authors have used we/our throughout rather than third person or indirect language.

Specific comments:

1. The first paragraphs of the Methods and Results sections don't have subheadings - these should be added.
2. The authors used very broad inclusion criteria, for instance including children to age 18 years. A more limited focus, for instance on infancy or the preschool years, may have provided more emphasis on key early aspects of development.
3. Further, vitamin D may be hard to consider as an isolated effect (separate to other micro or macronutrient factors). Was this considered?

Are the rationale for, and objectives of, the Systematic Review clearly stated?

Yes

Are sufficient details of the methods and analysis provided to allow replication by others?

Yes

Is the statistical analysis and its interpretation appropriate?

I cannot comment. A qualified statistician is required.

Are the conclusions drawn adequately supported by the results presented in the review?

Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Paediatric gastroenterology, with focus on IBD and the impact of vitamin D

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.
