Review Article

Interventions to improve vitamin D status in at-risk ethnic groups during pregnancy and early childhood: a systematic review

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Abstract

Objective: To systematically review the literature with the primary aim of identifying behavioural interventions to improve vitamin D stores in children from at-risk ethnic groups.

Design: Review based on Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines. PROSPERO registration number: CRD42017080932. Health Behaviour Model and Behaviour Change Wheel framework constructs used to underpin evaluation of interventions. Methodological quality evaluated using Cochrane Risk of Bias, Cochrane ROBINS-I and NHLBI tools.

Setting: Databases Cochrane Library, MEDLINE, EMBASE, CINAHL with secondary search of Google Scholar. No country limits set. Papers between January 1990 and February 2018, published in English included. Anticipating study heterogeneity, outcome measures not pre-specified and identified from individual full papers. Updated literature search November 2020.

Participants: Patient or population including pregnant women, newborns and children aged under 18 years, from Asian or African ethnic groups.

Results: Of 10,690 articles screened, 298 underwent full-text review, with 24 ultimately included for data extraction. All identified studies conducted a vitamin D pharmacological supplementation intervention, with two also incorporating a behavioural intervention strategy. No study explicitly defined a primary aim of evaluating a behavioural intervention, undertaken to study its effect on vitamin D supplement uptake.

Conclusions: There is a need to address the paucity of data in ethnic at-risk children on how behavioural interventions ideally developed and co-produced with the community under study, affect and help improve vitamin D uptake, within the antenatal and pregnancy phase as well as during childhood.

Keywords

Vitamin D
Ethnic minorities
Children
At risk
Optimisation
Interventions

Vitamin D deficiency is a global health problem affecting over 1 billion people worldwide(11). Vitamin D is an essential nutrient that contributes to bone health, and deficiency status increases the risk of rickets, osteomalacia and osteoporosis(2-4). Pregnant women and their breast-fed neonates as well as older children are at particular risk for vitamin D deficiency, as are Black and ethnic minority groups(1). Hypovitaminosis D(5) is common in pregnant women (up to 50%) and their breast-fed infants (up to 56%)(6) and has been reported in 64% Middle Eastern women, 58% Black women and 47% Asian women, compared to only 13% of Caucasian women(7). Short-term implications include lower neonate birth weight, length and head circumference(6,8-10) and skeletal outcomes such as muscle pain and fractures(11). Hypo-calcaemic seizures, rickets and cardiomyopathy have been reported in children under 5 years(2,12). Deficiency in childhood is associated with obesity and metabolic syndrome(13) and is a significant risk factor for dental caries(10,14,15).

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The major cause of vitamin D deficiency is low transmission or reduced penetration of solar ultra-violet B radiation, affecting the cutaneous synthesis of vitamin D. Although found naturally in some foods such as oily fish, red meat, liver and egg yolks and fortified foods (infant formula, breakfast cereals and fat spreads), less than 10% of vitamin D stores come from individual diets\(^1\)\(^2\). Often, the main source of dietary vitamin D is in the form of supplements. In women of child-bearing age, supplementation rates range between 12% and 27%\(^3\), and data indicate that 63% of women of reproductive age are vitamin D-deficient\(^4\). The Royal College of Paediatrics and Child Health and the British Paediatric Surveillance Unit recently ascertained the national incidence of nutritional rickets in children under the age of 16 years\(^5\). Black and South Asian children had a 10-fold and 5-fold greater incidence of nutritional rickets, respectively, compared with other ethnic groups under 5 years of age.

Despite current national policy and guidance to ensure supplementation in high-risk groups\(^6\), uptake is not consistent and a deeper understanding of facilitators and barriers to improve uptake is required. Behavioural interventions have shown some promise in contributing to the prevention, management and treatment of various other non-communicable diseases; obesity, diabetes, chronic pain, asthma and emotional difficulties\(^7\). In this paper, we refer to ethnic minority groups, accepting that the term is often used interchangeably with race\(^8\). Race is a social construct, usually identified based on a combination of physical, cultural and behavioural attributes, whereas ethnicity, self-identified by an individual, encompasses aspects such as nationality, culture, language and religion\(^9\). There are known structural barriers that Black, Asian and minority ethnic (BAME) groups face when accessing healthcare or being involved in clinical research\(^10\). Lack of exposure to health promotion messages and economic disadvantage are factors contributing to health inequalities. Considering these inter-related complexities, it has been recognised that interventions that are developed with the community of interest and which are context-specific and tailored may be more effective\(^11\). Our teams have a record of working with at-risk communities. With time, we have become increasingly aware of the need to understand how behavioural interventions may help at-risk communities to comply with public health advice, including that for vitamin D supplementation. Acknowledging this background, we undertook a systematic review to evaluate behavioural interventions implemented to improve or optimise vitamin D stores in children from at-risk ethnic groups.

## Methods

Our systematic review, performed based on the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines\(^12\), is registered within the PROSPERO database, CRD42017080932.

Identified interventions were planned to be evaluated using the Health Behaviour Model (HBM\(^13\)). The HBM\(^13\) attempts to explain and predict health-related behaviours across four constructs: (i) perceived susceptibility (i.e. affect with known risk of deficiency); (ii) perceived severity (i.e. affect with known consequences/health outcomes of deficiency); (iii) perceived benefits (defined here as promoters for optimal vitamin D status, i.e. sufficient sun exposure, supplementation, health information access); and (iv) perceived barriers (i.e. sufficient sun exposure, supplementation, health information access)\(^14\).

Classification of behavioural interventions was to be based on the nine categories within the Behaviour Change Wheel framework\(^15\); (1) education (e.g. increasing knowledge or understanding around benefits of supplementation/sun exposure or consequences of deficiency); (2) persuasion (e.g. reminder to supplement); (3) incentivisation (e.g. expectation of reward); (4) coercion (e.g. expectation of punishment, cost or fine); (5) training (e.g. how to take/administer supplements or drops to children); (6) restriction; (7) environmental restructuring (e.g. shared decision-making with General Practitioners (GP)/health professionals); (8) modelling (e.g. providing examples); and (9) enablement (e.g. reducing barriers).

### Search question

The PICO acronym, an established model for aiding systematic reviews, guided all elements of our research question:

1. **Patient or Population:** children under 18 years of age from an Asian or African ethnic group or race, including pregnant women and their newborns.
2. **Intervention:** to optimise vitamin D status (thresholds for optimisation as defined by research group), with primary focus on behavioural studies.
3. **Comparator:** all controls.
4. **Outcomes:** improvement in vitamin D status (no pre-specified definition).

### Data sources

The following databases were searched and verified: Cochrane Library, MEDLINE, EMBASE, CINAHL with limited secondary search on Google Scholar, in order to identify relevant studies. No country limits set. Only articles written in English published after 1990 were included. Searches were conducted in February 2018, with a limited updated search in November 2020.

### Search strategy

The search strategy included terms for ‘vitamin D’ and ‘ethnic’ and terms specifying all major subgroups. Details are as follows:

- **Term 1:** Vitamin D
  - Vitamin D * OR ricket* OR osteomalacia
- **Term 2:** Ethnic
  - Ethnic group* OR Asia* OR Africa* OR emigrant* OR immigrants
Search string:
(Vitamin D* OR ricket* OR osteomalacia) AND (ethnic group* OR Africa* OR Asia* OR emigrant* OR immigrant*)

A selective search strategy focusing on identifying behavioural interventions was used for updated review of abstracts February 2018–November 2020 (Additional File 1).

**Eligibility criteria**
We classified Asian ethnic groups as individuals of central, east, south, south-east, and western Asian origin and African as individuals of east, north, south, and western and south of the Sahara origin. Studies were included if they met the PICO inclusion criteria above, were published in English or with translation available, and were randomised controlled trials (RCT), quasi-RCT and non-RCT (before-after studies).

We excluded purely observational studies, non-English language, or no English version available, full-text not available, or studies classified as dissertations/abstracts/conference pieces/editorial letter or review. We also excluded studies with no or non-extractable ethnic/demographic or vitamin D data and those focusing on adult populations.

**Study selection and data extraction**
Two reviewers (EA and CL) shared screening of titles and abstracts. Shortlisted abstracts underwent full-text review by two researchers (PA and AT), with conflicts resolved by a third reviewer (CL). For each study of interventions, data were extracted and classified (AT and PA) and checked (EA and NT) for publication year, characteristics of study population (sample size, mean age, ethnicity, gender and age), study design, available vitamin D data, supplementation data and intervention measured. For the updated search in November 2020, in line with our primary aim, we searched only for studies with an explicitly defined behavioural intervention. Titles and abstracts were screened by two reviewers (RP and NT) and confirmed with third reviewer (EA).

**Risk of bias assessment**
Methodological quality of the studies was assessed using the Cochrane Risk of Bias-2 (231) assessment tool for RCT (nineteen studies), the Cochrane ROBINS-I (32) tool for non-randomised studies of interventions (one study) and the NHLBI (33) tools for Observational Cohort and Cross-Sectional Studies (one study) or Before-After (Pre-Post) Studies with No Control Group (three studies). The studies were quality-assessed by EA, with ratings reviewed by NT, points of uncertainty discussed and final ratings agreed with NT (n 5; 22%). Studies with a high or critical risk of bias were included but with quality ratings highlighted within the results section in order to contextualise findings.

**Results synthesis**
In view of the heterogeneity of studies identified in terms of methods, participants, vitamin D thresholds used, the interventions and outcomes, a narrative approach to synthesis was used following guidance developed by the University of York Centre for Reviews and Dissemination and the Economic and Social Research Council (34,35).

**Results**

**Study selection**
Initially, 10 690 articles were identified. Title and abstract screening returned 298 potential articles. After full-text review, 274 articles were excluded, and 24 intervention studies (including 2 with behavioural components) were included. No further behavioural studies were identified from the updated literature search in November 2020 (see Fig. 1).

**Study characteristics**
The majority (11) of studies included were conducted in the USA, with two further studies each in Australia, Canada, and Mongolia. One study was undertaken in each of the following: UK, Norway, Denmark, Pakistan, India, Nigeria, and Turkey. Table 1 summarises these papers.

With reference to quality appraisal, nineteen studies were appraised using Cochrane Risk of Bias for RCT. Of these, one (36) had a low risk of bias and the remaining eighteen had some concerns. One study (37) appraised with the ROBINS-I tool, was coded as having a critical risk of bias due to confounding. Of the NHLBI appraised studies, two were rated as fair (38,39) and two as good (40,41) (see Tables 2–5).

**Behavioural interventions**
All studies included conducted a vitamin D pharmacological supplementation intervention in our target population, with two that incorporated a behavioural component to their intervention strategy (38,42) in addition to the pharmacological intervention. We found no study that explicitly defined a primary aim of evaluating a behavioural intervention undertaken with the intention to study its effect on vitamin D supplement uptake.

Madar et al. (42) (Table 2; overall risk of bias: some concerns) studied the effect of vitamin D2 drops on serum 25-hydroxy-vitamin D (25(OH)D) in infants with immigrant origin within a cluster RCT. In total, sixty-six healthy infants of Pakistani (South Asian origin), Turkish (West Asian/Middle Eastern) or Somali (East African) origin were recruited for the study from eight child health clinics in Oslo, Norway. The behavioural component involved multilingual translated brochures for mothers, with information provided to each ethnic group on vitamin D, its sources and instructions on how to administer the vitamin D drops,
made available free of charge. Aims were to evaluate whether a free supply of a 400 iu daily dose, for 6-week old infants, together with information handouts that had been translated into Urdu, Turkish or Somali languages incorporating text and simple illustrations, improved vitamin D status, assessed at 7-week follow-up in the intervention and control group. Fifty-one (78 %) infants completed the study, with serum 25(OH)D levels significantly higher in the intervention group \( v \) Control group (93·5 \( v \) 72·7 nmol/l; \( P = 0·03 \)). Amongst exclusively breast-fed infants at baseline, serum 25(OH)D levels increased by 32·3 nmol \( P = 0·035 \) in the intervention group. This study concluded that free supply of vitamin D drops, with translated information handouts, significantly improved the vitamin D status of healthy infants of immigrant background. Considering the Behaviour Change Wheel framework, this combined intervention strategy (if part of an explicitly stated behavioural intervention) could have been coded under criteria; education and training (translated information leaflets; administration advice) and enablement (free supply)\(^{30}\).

The second study with behavioural components centred around a nationwide prevention campaign instigated due to resurgence of vitamin D deficiency rickets in children of ethnic minority origin living in Turkey, a sunny country in West Asia/Middle East. In order to gauge the campaign’s impact which included supply of vitamin D supplements as an intervention, Mutlu et al\(^{38}\) (Table 4;
| Reference                  | Country – latitude if stated | Design                        | Population                                      | Sample size | Age (years) | Intervention – pharmacological/behavioural | Vitamin D measure (sufficiency/insufficiency/deficiency threshold used) | Primary findings                                                                                                                                                                                                 |
|---------------------------|-----------------------------|-------------------------------|-------------------------------------------------|-------------|-------------|-------------------------------------------|-------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------
| Abrams et al. (2013)[56]  | USA – sunny latitude        | 8-week double-blind RCT       | Healthy pre-pubertal; African American (12), Hispanic (23), Asian (1) and White (27) | 64 (one lost to follow-up) | 4–9 | Supplementation with 1000 iu/d for 8 weeks.                          | Serum 25(OH)D (baseline 25(OH)D concentrations: low (20 ng/ml) 50 nmol/l; middle (20–32 ng/ml) 50–80 nmol/l; high (32 ng/ml)) 80 nmol/l). | Vitamin D<sub>3</sub> supplementation with 1000 iu taken daily increased 25(OH)D levels and decreased parathyroid hormone in children, whose average vitamin D intake was below Institute of Medicine dietary recommendation; but with no significant effect of change in vitamin D<sub>3</sub> levels on Ca absorption, as after 8 weeks, with baseline values used as a covariate, no differences were seen in fractional or total Ca absorption based on supplementation group (P = 0.75 and 0.36, respectively). |
| Andersen et al. 2008[56]  | Denmark – Copenhagen area, (latitude 550N) | 1 year-double-blind RCT | Adolescent girls Pakistani origin NB: women (18.1–52.7 years) and men (17.9–63.5 years). | 21 girls | (10.1–14.7 years) | Two doses of vitamin D<sub>3</sub> (10 and 20 mcg/d). | Serum 25(OH)D | Difficult to comment on any significant difference in serum vitamin 25(OH)D<sub>3</sub> levels in the group of girls at end of study due to the low number recruited and high dropout rate. NB: Supplementation with 10 and 20 mcg vitamin D<sub>3</sub>/d increased serum 25(OH)D<sub>3</sub> levels similarly in vitamin D-deficient Pakistani women (fourfold); 10 mcg increased serum-25(OH)D<sub>3</sub> concentrations twofold and 20 mcg threefold in Pakistani men. Serum 25(OH)D<sub>3</sub> concentrations increased at 6 months and were stable thereafter. |
| Dong et al. (2010)[55]    | USA – sunny latitude        | 16-week open-label investigator-blinded RCT | Healthy African American | 49 | 14–18 | Supplementation with 2000 iu/d compared with 400 iu/d for control group. | Serum 25(OH)D (sufficiency: 25(OH)D level of at least 75 nmol/l. Insufficiency: more than 50 but less than 75 nmol/l. Deficiency: under 50 nmol/l). | Study group reached significantly higher 25(OH)D levels at 8 and 16 weeks. The 2000 iu group achieved 25(OH)D levels of 85.7 +/- 30.1 nmol/l at 16 weeks from 33.5 +/- 9.9 nmol/l at baseline; P< 0.001. Daily 2000 iu vitamin D supplementation may be effective in optimising vitamin D status and countering the progression of aortic stiffness in Black youth. Plasma vitamin 25(OH)D levels in response to 2000 iu/d supplementation negatively modulated by adiposity. |
| Reference             | Country – latitude if stated | Design                           | Population                                      | Sample size | Age (years) | Intervention – pharmacological/behavioural | Vitamin D measure (sufficiency/insufficiency/deficiency threshold used) | Primary findings                                                                 |
|-----------------------|------------------------------|----------------------------------|-------------------------------------------------|-------------|-------------|--------------------------------------------|-----------------------------------------------------------------------------|--------------------------------------------------------------------------------|
| Dougherty 2015(51)    | Philadelphia, USA            | 12-week RCT                      | African American children and young adults with SCD. | 21 with and 23 without SCD-SS                  | 5–20 years old | To assess safety and efficacy of two oral daily doses (4000 v. 7000 iu) of cholecalciferol (vitamin D₃). | Serum 25(OH)D With suboptimal vitamin D status (<32 ng/ml). | After 12-week supplementation, both vitamin D₃ doses were safe (no hypercalcaemia) and well tolerated. For both groups, deficient (<20 ng/ml) vitamin D status was eliminated only in those receiving 7000 iu/d (P < 0.05). |
| Ganmaa et al. (2008)(39) | Mongolia (East Asia) – Northern latitude | Observation/pilot feasibility study | Mongolian                                       | 46          | 9–11        | Supplementation with three 236-ml tetra pack boxes of conventional UHT-processed fortified whole milk daily (100 iu vitamin D/serving). | Serum 25(OH)D | After 1 month of drinking vitamin D fortified milk, vitamin 25(OH)D levels were improved. |
| Ganmaa 2017(57)      | Mongolia (East Asia) – Northern latitude | (i) 6-month RCT (ii) 7-week cluster-randomised classroom-based | Urban school age children                        | (i) 113 (ii) 235 | (i) 12–15 years (ii) 9–11 years | (i) 800 iu vitamin D daily (ii) 710-ml whole milk fortified with 310 iu vitamin D, daily. | Serum 25(OH)D Deficiency defined as <20 ng/ml. | Correction of baseline vitamin D deficiency in children with vitamin D 800 iu supplement daily over 6 months increased growth. |
| Green et al. (2015)(37) | Canada                        | Observation                       | Asian (28) and White (37)                        | 65          | 0.2–0.4     | Fifty-eight (89 %) breast-fed term infants had been given a vitamin D supplement, in most cases providing 10 mcg/dose. | Serum 25(OH)D | Mean 25(OH)D levels of the infants = 31 ng/ml, 95 % CI, 28–34 ng/ml. Infants who received 10 mcg vitamin D achieved significantly higher 25(OH)D concentrations than those who received less; mean 25(OH)D 9.4 ng/ml higher; P = 0.003. Maternal vitamin 25(OH)D, season, skin colour and ethnicity were not significant determinants of infant vitamin 25(OH)D levels. |
| Hanson et al. (2011)(46) | Canada                        | RCT                              | Infants receiving formula feedings during intensive care hospitalisation. White/non-White infants (African American) | 52          | Newborn; infants 32 weeks gestational age to discharge | Supplemented group received 400 iu/d vitamin D₃ in addition to formula feedings; control group received matching placebo. | Serum 25(OH)D Infants measured from cord blood, blood serum test every 7 d, and at discharge. Intact parathyroid hormone was measured at discharge. | White infants had statistically significant higher mean 25(OH)D levels at discharge compared with non-White infants (P = 0.0003). Mean supplemented group 23.1 +/- 7 ng/ml (57.66 +/- 17.47 nmol/l); Mean non-supplemented group 17.8 +/- 4.7 mg/ml (44.43 +/- 11.3 nmol/l (P = 0.007) |
| Reference     | Country – latitude if stated | Design                                      | Population                                      | Sample size | Age (years) | Intervention – pharmacological/behavioural | Vitamin D measure (sufficiency/insufficiency/deficiency threshold used) | Primary findings                                                                                                                                 |
|---------------|------------------------------|---------------------------------------------|------------------------------------------------|-------------|-------------|---------------------------------------------|-----------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------|
| Hollis et al. (2011)\(^{48}\) | USA – 32° North | Single-centre RCT                         | Women with singleton pregnancies at 12–16 weeks gestation stratified by self-defined race/ethnic group | 350         | Pregnancy/newborn | Vitamin D supplementation daily until delivery; doses 400 IU daily; 2000 IU daily; 4000 IU daily. NB: Tablet concentration verified by independent company every 6 months. | Primary outcome maternal/neonatal circulating 25(OH)D at delivery; secondary outcomes 25(OH)D > or = 80 nmol/l achieved?; 25(OH)D concentration required to achieve maximal 1,25(OH)D production. | Mean vitamin 25(OH)D levels by group at delivery and 1 month before delivery was significantly different (\(P < 0.0001\)). 4000 IU daily dose in pregnancy is safe and most effective in achieving sufficiency in all women regardless of race (\(P < 0.0001\)). |
| Hollis et al. (2015)\(^{43}\)   | Rochester, New York, USA      | 6-month RCT                                | Mother–infant pairs; Maternal race/ethnicity: Black (African American)/Hispanic/White | 334         | Newborn/breast-feeding 4–6 weeks old, with planned breast-feeding for 6 months | Maternal vitamin D\(_3\) supplementation with 6400 IU/d v. maternal and infant supplementation of 400 IU/d. | Maternal serum/neonatal serum 25(OH)D. Vitamin D deficiency < 50 nmol/l. Compared with 400 IU vitamin D\(_3\)/d, 6400 IU/d was safe; with significant increase in maternal vitamin 25(OH)D from baseline (\(P < 0.0001\)). | Vitamin D deficiency in breast-fed infants affected by race. African American mothers and infants had substantially lower circulating 25(OH)D levels; several minority infants had severe vitamin D deficiency (2.5 nmol/l 25(OH)D) after 1 month of breast-feeding. |
| Hossain et al. 2014\(^{47}\)      | Pakistan, South Asia          | Single-centre open-label RCT                | Pakistani pregnant women and neonates          | 193         | Pregnancy/newborn, within 48 h of birth | 4000 IU vitamin D daily from week 20 to delivery v. routine care (200 mg ferrous sulphate and 600 mg Ca). | Serum 25(OH)D Cord blood or neonatal serum levels. Maternal serum at baseline and delivery. | Maternal vitamin D supplementation improved maternal and neonatal vitamin D status. Maternal 25(OH)D levels increased in treated group 18.3 +/- 11 ng/dl v. 8.8 +/- 11.94 ng/dl; \(P = 0.001\).  Routine care group 25(OH)D levels 6.9 +/- 7.0 ng/dl v. 6.35 +/- 3.97 ng/dl; \(P = 0.6\). There was a positive correlation between maternal and neonatal 25(OH) vitamin D levels (\(r = 0.83; P = 0.001\)). |
| Lewis et al. (2013)\(^{54}\)       | USA – latitudes 34°N and 40°N, | RCT – multisite triple-masked trial        | Black and White                                | 323         | 9–13         | Children received 1 of 5 oral vitamin D\(_3\) doses (0, 400, 1000, 2000 and 4000 IU/d) and were sampled over 12 weeks. | Serum 25(OH)D Serum 1,25(OH)\(_2\)D | Mean baseline 25(OH)D for entire sample 70 nmol/l. Increases in 25(OH)D levels depended on dose taken, with 12-week changes ranging from ~10 nmol/l for placebo to 76 nmol/l for 4000 IU. Larger vitamin 25(OH)D gains observed for Whites compared with Black children at highest dose (\(P < 0.01\)). |
| Reference          | Country – latitude if stated | Design          | Population                              | Sample size | Age (years) | Intervention – pharmacological/ behavioural | Vitamin D measure (sufficiency/ insufficiency/deficiency threshold used) | Primary findings                                                                                                                                                                                                 |
|--------------------|-----------------------------|-----------------|-----------------------------------------|-------------|-------------|---------------------------------------------|--------------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Madar et al. (2009)(42) | Norway                      | RCT             | Pakistani, Turkish and Somali           | 51          | 7-week-old | 7-week supplementation of 400 iu of vitamin D2 and tailor-made information about vitamin D and its sources. | Serum 25(OH)D                                                           | Vitamin D supplementation provided free of charge together with translated information brochures significantly increased vitamin serum 25(OH)D concentrations when compared to a control group receiving usual care. |
| Mondal et al. (2014)(44)  | India                       | RCT             | Indian children with nutritional rickets. | 71          | 6 months to 5 years | Single intramuscular dose of 600 000 iu of vitamin D or 10 weeks of 60 000 iu of vitamin D taken weekly for 10 weeks. 12-week follow-up. | Serum 25(OH)D                                                           | No difference found in efficacy of the two regimens on comparing diagnostic parameters for rickets. Both treatments were safe and effective. No difference in mean serum 25(OH)D levels in both groups at baseline and 12 weeks (P = 0.887); Increase in serum 25(OH)D in both groups statistically significant; P < 0.001 in both groups at 12 weeks v. baseline. No significant difference between groups on comparing proportion of children with biochemical features suggestive of rickets at any point of the study (P > 0.05), except at 4 week follow-up, there were more cases of hypocalcaemia in oral group (P = 0.039). |
| Mutlu 2011 (38)        | Turkey (Middle East/West Asia) | Observational cohort cross-sectional study to evaluate efficacy of nationwide prevention programme | Turkish infants, recalled as part of national screening for congenital hypothyroidism; Information regarding age at start of supplementation, the dosage and compliance obtained from mothers with face-to-face interview. | 85          | 16.5 +/- 20.7 (3 – 120) d | Babies had been provided with free vitamin D (cholecalciferol) solution and recommended to receive 400 iu (3 drops) daily. | Serum 25(OH)D Mean 25-(OH)D level was 42.5 ± 25.8 (median: 38.3) ng/ml. | Taking 400 iu/d of vitamin D seems adequate to prevent vitamin D deficiency. |
| Reference | Country – latitude if stated | Design | Population | Sample size | Age (years) | Intervention – pharmacological/behavioural | Vitamin D measure (sufficiency/insufficiency/deficiency threshold used). | Primary findings |
|-----------|-----------------------------|--------|------------|-------------|-------------|-------------------------------------------|-------------------------------------------------|----------------|
| Rajakumar et al. (2005)(41) | Philadelphia, USA | Non-randomised pre-post comparison | African American; at-risk vitamin D sample | Total 41: 27 males and 14 females | 6–10 | 4-week 400 iu vitamin D supplementation. | Serum 25(OH)D 1,25(OH)2D (i) 12–20 ng/ml essential for bone health in children. (ii) Deficiency: less than 10 ng/ml. Insufficiency: 10–20 ng/ml based on definition for adults. | Vitamin D insufficiency in approximately half of participants at baseline. Although with supplementation vitamin D levels increased significantly*, vitamin D insufficiency persisted in 18% of experimental group. *Significant incremental increase in 25(OH)D levels found only among vitamin D insufficient children (15·4 +/- 3·2 ng/ml to 23 +/- 6·4, P < 0·001 v. 32·1 +/- 8·1 ng/ml to 31·8 +/- 5·8 ng/ml, P = 0·79). |
| Rajakumar et al. (2008)(40) | Philadelphia, USA | Non-randomised pre-post comparison | African American | Total 41: 21 obese and 20 non-obese | 6–10 | 4-week 400 iu/d vitamin D supplementation. | Serum 25(OH)D Deficiency: less than 20 ng/ml. Insufficiency: 21–29 ng/ml. | Vitamin D deficiency common among obese and non-obese pre-adolescent African American children. Vitamin D 400 iu daily dose for 1 month is not enough to raise blood levels to 30 ng/ml. Vitamin D deficiency in 12/21 (57%) obese v. 8/20 (40%) non-obese at baseline (P = 0·35); persisted in 5/21 (24%) obese v. 2/18 (11%) non-obese (P = 0·42) after treatment. |
| Rajakumar et al. (2015)(49) | Philadelphia, USA | RCT | African American (84) and White (73) | 157 | 8–14 | 6 months supplementation with vitamin D3 1000 iu/d | Serum 25(OH)D | Mean circulating levels of vitamin D were higher in supplemented group v. placebo group. Effect of vitamin D3 supplementation varied by race and was found to be more effective and significant in Black children*. *Levels at 2 months 26·4 +/- 8·1 v. 18·9 +/- 8·1 ng/ml; P < 0·001 and at 6 months 26·7 +/- 7·6 v. 22·4 +/- 7·3; P = 0·003. Lower baseline 25(OH)D levels associated with greater change at 2 months overall (r = −0·49; P < 0·001) but not in White children (r = −0·005; P = 1·0). |
| Reference | Country – latitude if stated | Design | Population | Sample size | Age (years) | Intervention – pharmacological/behavioural | Vitamin D measure (sufficiency/insufficiency/deficiency threshold used) | Primary findings |
|-----------|-----------------------------|--------|------------|-------------|-------------|------------------------------------------|------------------------------------------------|-----------------|
| Rajakumar et al. (2016) | Philadelphia, USA | RCT | African American (52) and White (44) | 96 | 8–14 | Supplemented for 6 months with 1000 iu/d vitamin D₃ | Serum 25(OH)D | At baseline, Black children had lower mean 25(OH)D levels ($P < 0.0001$) and higher Ca levels ($P = 0.001$) than White children. With intervention, the within-group change $P$-value was <0.0001 for treated v. 0.041 for placebo group. Vitamin D intake required in order to maintain 25(OH)D levels at 20 ng/ml in 97.5 % of Black and White children, adjusting for race and pubertal status, was three times higher than (current USA) recommended daily allowance of 600 iu/d. |
| Rodda et al. (2015) | Australia, latitude 38 degrees south | RCT | Vitamin D-deficient pregnant women and their newborns. Dark skin or veiled: 84 % in treated group v. 97 % control group. | 78 | Pregnancy; newborn Treatment 12–16 weeks' gestation to delivery when maternal and neonate vitamin D levels tested | Supplementation with 2000–4000 iu/d until 28-week gestation | Serum 25(OH)D | Recruited women with singleton pregnancies with serum 25(OH)D levels less than 75 nmol/l, defined as deficiency or insufficiency, at first antenatal appointment at 12–16 weeks gestation. Umbilical cord serum 25(OH)D concentration at delivery was higher in neonates of the treatment group mothers compared with neonates of control group mothers (81 nmol/l (range 70–91) v. 42 nmol/l (range 34–50), CI 95 %; $P < 0.0001$). Strong positive correlation between maternal and umbilical serum 25(OH)D levels at delivery; spearman rank correlation coefficient 0.88; $P < 0.0001$. |
| Sacheck et al. (2017) | USA | RCT | Black, Hispanic, White and Asian. | 604 | 8–15 | Supplemented with vitamin D₃ at 600, 1000 or 2000 iu/d for 6 months | Serum 25(OH)D | Serum vitamin 25(OH)D levels increased over 6 months in all three dose groups. The 2000 iu/d group achieved a higher vitamin D concentration than the other two dose groups (33.1 v. 26.3 and 27.5 ng/ml; $P < 0.001$). |
| Reference       | Country – latitude if stated | Design          | Population                          | Sample size | Age (years)            | Intervention – pharmacological/behavioural | Vitamin D measure (sufficiency/insufficiency/deficiency threshold used). | Primary findings                                                                                                                                 |
|-----------------|------------------------------|-----------------|-------------------------------------|-------------|------------------------|---------------------------------------------|---------------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------|
| Talib et al. (2016)⁵⁵ | USA, New York latitude 40·8 ° NE | RCT             | Adolescents with vitamin D deficiency Hispanic, Black, White and Asian | 122         | 13–20 years; mean age 16·6 +/− 2·2 | Supplemented for 8 weeks with either 50 000 iu/week, 5000 iu/d or 1000 iu/d. | Serum 25(OH)D                                                                      | Adolescents require 8 weeks of high-dose cholecalciferol (at least 5000 iu/d) to correct vitamin D deficiency. Obese adolescents have poorer response to treatment (13·7 +/− 10·7 v. 21·9 +/− 16·9 ng/ml; P < 0·001). Mean change in 25(OH)D levels post-treatment: 24·9 +/− 15·1 v. 21·0 +/− 15·2–50 000 iu group v. 6·2 +/− 6·5 ng/ml–1000 iu respectively; P < 0·001. |
| Thacher et al. (1999)⁵⁸ | Nigeria, Africa             | RCT             | Nigerian children with rickets or bone deformities. | 109         | 1–14 years             | Supplemented for 24 weeks with either vitamin D 600 000 iu intramuscularly, calcium 1000 mg/d or a combination of vitamin D and calcium. | Serum 25(OH)D                                                                      | Calcium supplementation with or without vitamin D was more effective than supplementation with vitamin D alone in healing active rickets (P < 0·001). Daily dietary calcium intake was low in both children with rickets and control group children (median 203 mg and 196 mg, respectively; P = 0·64). |
| Yu et al. 2009⁷    | London, UK                  | Prospective RCT, | Indian Asian, Middle Eastern, Black and Caucasian | 180         | Study arms 27 weeks’ gestation to delivery | Study arms: single-dose 200 000 iu/800 iu daily/no treatment | Mothers – serum 25(OH)D. Babies – cord 25(OH)D. Thresholds: sufficient > 50 nmol/l; insufficient 25–50 nmol/l; deficiency < 25 nmol/l. | Final maternal 25(OH)D levels supplemented v. no treatment; daily dose median 42 (31–76) v. 27 (27–39) nmol/l; P < 0·0001. Cord levels significantly higher with supplementation; daily dose median 25 (18–34) v. 17 (14–22) nmol/l in no-treatment group; P = 0·001. Single or daily dose improved serum vitamin D levels significantly. Only small percentage of women (30 %) and babies (8 %) who were given supplements were vitamin D-sufficient. |
### Table 2: Cochrane risk of bias for randomised controlled trials (RCT)

| Study                          | Risk of bias arising from the randomisation process | Risk of bias due to deviations from the intended interventions (effect of assignment to intervention) | Missing outcome data | Risk of bias in measurement of the outcome | Risk of bias in selection of the reported result | Overall risk of bias |
|-------------------------------|----------------------------------------------------|-------------------------------------------------------------------------------------------------------|----------------------|---------------------------------------------|-----------------------------------------------|---------------------|
| Abrams et al. (2013)          | Low                                                | Low                                                                                                   | Low                  | Low                                         | Low                                           | Low                 |
| Anderson et al. 2006          | Some concerns                                     | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Dong et al. (2010)            | Low                                                | Some concerns                                                                                         | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Dougherty 2015                | Some concerns                                     | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Gannmaa 2017                 | Some concerns                                     | Low                                                                                                   | Low                  | Low                                         | Low                                           | Low                 |
| Hanson et al. (2011)          | Low                                                | Some concerns                                                                                         | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Hollis et al. (2011)          | Some concerns                                     | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Hollis et al. (2015)          | Low                                                | Some concerns                                                                                         | Low                  | Low                                         | Low                                           | Low                 |
| Hossain N et al. 2014         | Low                                                | Low                                                                                                   | Low                  | Low                                         | Low                                           | Low                 |
| Lewis et al. (2013)           | Low                                                | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Madar et al. (2009)           | Some concerns                                     | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Mondal et al. (2014)          | Some concerns                                     | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Rajakumar et al. (2015)       | Low                                                | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Rajakumar et al. (2016)       | Low                                                | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Rodda et al. (2015)           | Low                                                | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Sacheck et al. (2017)         | Low                                                | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Talib et al. (2016)           | Low                                                | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Thacher et al. (1999)         | Some concerns                                     | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |
| Yu et al. 2009                | Some concerns                                     | Low                                                                                                   | Some concerns        | Low                                         | Low                                           | Some concerns       |

### Table 3: Cochrane ROBINS-I

| Study                          | Bias due to confounding (low/moderate/serious/critical/no info (NI)) | Bias in selection of participants into the study (low/moderate/serious/critical/NI) | Bias in classification of interventions (low/moderate/serious/critical/NI) | Bias due to deviations from intended interventions (low/moderate/serious/critical/NI) | Bias due to missing data (low/moderate/serious/critical/NI) | Bias in measurement of outcomes (low/moderate/serious/critical/NI) | Bias in selection of the reported result (low/moderate/serious/critical/NI) | Overall bias |
|-------------------------------|--------------------------------------------------------------------|---------------------------------------------------------------------------------------|--------------------------------------------------------------------------|---------------------------------------------------------------------------------|-----------------------------------------------------------------|------------------------------------------------------------------|---------------------------------------------------------------|---------------|
| Green et al. (2015)           | Critical                                                           | Moderate                                                                              | Low                                                                      | Moderate                                                                       | No Info                                                        | Moderate                                                        | Moderate                                                       | Critical       |
### Table 4 Quality assessment tool for observational cohort and cross-sectional studies

| Was the research question or objective in this paper clearly stated? | Was the study population clearly specified and defined? | Was the participation rate of eligible persons at least 50%? | Was a sample size justification, power description, or variance and effect estimates provided? | For the analyses in this paper, were the exposure(s) of interest measured prior to the outcome(s) being measured? | Was the time frame sufficient so that one could reasonably expect to see an association between exposure and outcome if it existed? | For exposures that can vary in amount or level, did the study examine different levels of the exposure as related to the outcome (e.g. categories of exposure, or exposure measured as continuous variable)? | Were the exposure measures (independent variables) clearly defined, valid, reliable, and implemented consistently across all study participants? | Was the exposure(s) assessed more than once over time? | Were the outcome measures (dependent variables) clearly defined, valid, reliable, and implemented consistently across all study participants? | Were the outcome assessors blinded to the exposure status of participants? | Was loss to follow-up after baseline 20% or less? | Were key potential confounding variables measured and adjusted statistically for their impact on the relationship between exposure(s) and outcome(s)? | Quality rating (good, fair, poor) |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| Mutlu 2011(38) | Yes | Yes | NR | Yes | No | No | Yes | Yes | No | Yes | No | NR | NA | No | Fair |

### Table 5 Quality assessment tool for before-after (pre-post) studies with no control group

| Was the study question or objective clearly stated? | Were eligibility/selection criteria for the study population pre-specified and clearly described? | Were the participants in the study representative of those who would be eligible for the test/service/intervention in the general or clinical population of interest? | Were all eligible participants that met the pre-specified entry criteria enrolled? | Was the sample size sufficiently large to provide confidence in the findings? | Was the test/service/intervention clearly described and delivered consistently across the study population? | Were the outcome measures pre-specified, clearly defined, valid, reliable, and assessed consistently across all study participants? | Were the people assessing the outcomes blinded to the participants’ exposures/interventions? | Was the loss to follow-up after baseline 20% or less? | Were those lost to follow-up accounted for in the analysis? | Did the statistical methods examine changes in outcome measures from before to after the intervention? | Were statistical tests done that provided P-values for the pre-to-post changes? | Were outcome measures of interest taken multiple times before the intervention and multiple times after the intervention (i.e. did they use an interrupted time series design)? | If the intervention was conducted at a group level (e.g. a whole hospital, a community, etc.), did the statistical analysis take into account the use of individual-level data to determine effects at the group level? | Quality rating (good, fair, poor) |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| Gannmaa et al. (2008)(39) | Yes | No | No | Yes | Yes | Yes | Yes | NR | Yes | Yes | No | Yes | No | Yes | Fair |
| Rajakumar et al. (2008)(40) | Yes | Yes | Yes | NR | Yes | Yes | Yes | NR | Yes | No | Yes | No | Yes | No | Good |
| Rajakumar et al. (2005)(41) | Yes | Yes | Yes | NR | Yes | Yes | Yes | No | Yes | Yes | No | Yes | No | Yes | Good |
overall risk of bias: Fair) report from evaluation undertaken with a small sample of 85 healthy infants (45 girls and 40 boys; individual ethnic origin data not provided). The over-arching aims for the campaign, which also incorporated a curriculum to train healthcare workers, were to encourage the entire population, and especially pregnant and nursing women and infants, to have adequate sunlight exposure. Distribution of vitamin D supplements to every newborn at no cost to families was made through Turkey’s network of primary care units and maternal health centres. The study had limitations; no control group, high risk of bias from confounding and no statistical test was performed (Table 4). Evaluation was by way of face-to-face interviews with the mothers, advised to administer three drops (400 iu) daily. Overall, seventy-six (89 %) infants received the vitamin D dose recommended. However, mothers for twenty-six (31 %) infants stated that they did not administer the dose on a daily basis. The research team concluded that for prevention of vitamin D deficiency in infants, a daily dose of 400 iu was sufficient. Considering the Behaviour Change Wheel framework, the intervention strategy utilised could be coded under criteria of education and enablement(30). The authors(38) noted that the major obstacles for use of vitamin D supplements in Turkey included limited public awareness, access to healthcare and supplement costs.

Pharmacological interventions
All studies included in this review incorporated some form of pharmacological intervention, the results of which are summarised in the following sections. Ten studies(7,37,38,42-48) were undertaken during the pregnancy/newborn phase with ethnic minority origin as a main risk factor. One study specifically targeted a cohort with known ‘at-risk status’, that is, nutritional rickets(44).

Pregnancy/newborn group
Green et al.'s observational study assessed sixty-five Canadian breast-fed infants (Asian/White; aged 0·2–0·4 weeks), noting that supplementation with 10 mcg vitamin D helped infants achieve significantly higher 25(OH)D concentrations compared with those who received less (mean 25(OH)D 9·4 ng/ml higher; P = 0·003). Maternal vitamin 25(OH)D levels, season, skin colour and ethnicity were not significant determinants of infant vitamin 25(OH)D levels. Hanson et al.'s study(40) also undertaken in Canada evaluated newborn infants receiving formula feeds during intensive care hospitalisation. Classified as White or non-White African American origin, the study arms included supplementation with 400 iu vitamin D or matching placebo with formula feeds. White infants achieved significantly higher mean 25(OH)D levels by time of discharge (P = 0·0003).

Hollis et al. reported findings from two studies(43,48). The 2011 US cohort(48) were women with singleton pregnancies, given varying doses of vitamin D (400/2000/4000 iu daily) from 12–16 weeks of gestation to delivery. The 4000 iu daily dose was safe and the most effective in achieving sufficiency in all women regardless of race (Caucasian; African American; Hispanic; P < 0·0001). The 2015 US study(43) included 334 mother–infant pairs with newborns of 4–6 weeks old, recruited as mothers planned to breast-feed for 6 months. Compared to a 400 iu daily dose, the higher 6400 iu daily dose showed a significant increase in maternal vitamin D levels (P < 0·0001) and was safe. Vitamin D deficiency in breast-fed infants was affected by race, with African American mothers and infants having substantially lower circulating 25(OH)D levels.

Rodda et al.’s Australian study(45) recruited 78 ethnic minority pregnant women and newborns, identified on the basis of dark skin or veiling (84 % treated and 97 % control group). They were supplemented with 2000–4000 iu daily from 12 to 16 weeks’ gestation to delivery. Umbilical cord serum 25(OH)D levels at delivery were higher in neonates of mothers in the treated group (81 nmol/l v. 42 nmol/l; P < 0·0001).

Yu et al.’s prospective UK study(47) recruited 180 women, of Indian Asian, Middle Eastern, Black and Caucasian origin, treated from 27 weeks’ gestation to delivery. Study arms compared a single 200 000 iu dose injection, an 800 iu daily dose and no treatment. Treated mothers had significantly improved vitamin D levels (42 v. 27 nmol/l; P < 0·0001), but only a small percentage (women 30 %; babies 8 %) given supplements achieved sufficient levels (50 nmol/l or over).

The intervention in Hossain et al.’s single-centre open-label RCT(47) in Pakistan (South Asia) was 4000 iu vitamin D taken daily from gestational week 20 to delivery. Positive correlation was found between maternal and neonatal 25(OH)D levels (r = 0·83; P = 0·001), with improved status with supplementation (maternal 25(OH)D increased from mean 8·82 ng/dl to 18·3 ng/dl; P = 0·001). Mondal et al.(44) studied seventy-one Indian children (South Asia) with nutritional rickets, aged between 6 months and 5 years, given either a single intramuscular dose of 600 000 iu vitamin D or 10 weeks of a 60 000 iu dose taken weekly. Both were safe and effective with no difference found in efficacy on comparing diagnostic parameters for rickets (P > 0·05).

The Madar et al.(42) and Mutlu et al.(38) studies discussed above included a behavioural component in addition to the pharmacological intervention.

Children
Mondal et al.(44) study on children aged between 0·5 and 5 years is discussed above. Of fourteen further papers in children (aged 1–20 years), ten were undertaken in the USA and included African American(40,41,49–52), Hispanic, Asian and White(50), or Black and White(53–55) children. None reported including any type of behavioural intervention.

Lewis et al.(50) assessed five daily oral vitamin D doses (0, 400, 1000, 2000 and 4000 iu) in Black and White children aged 9–13 years. Increases in vitamin 25(OH)D levels
depend on dose taken; 12-week changes ranged between −10 and 76 nmol/l (placebo v. 4000 iu, respectively). In the highest dose group, larger vitamin D gains were seen in White compared with Black children (P < 0.01).

Rajakumar et al. reported vitamin D insufficiency at baseline[41] in about half of their study group (6–10 years old, African American children). Using the 2005 definition for adults, deficiency status was defined as less than 10 ng/ml and insufficiency as between 10 and 20 ng/ml[41]. Supplementation with 400 iu daily for 4 weeks helped achieve a significant increase in levels (see Table 1), but insufficiency status still persisted in 18% of the experimental group[41]. In a separate study, Rajakumar et al. also noted[40] that vitamin D deficiency is common amongst obese and non-obese preadolescent African American children, finding that 400 iu daily dose was not enough to raise vitamin D levels to 30 ng/ml (Table 1; deficiency defined as less than 20 ng/ml; insufficiency 21–29 ng/ml). In further work[49,52], Rajakumar et al. studied African American and White children aged 8–14 years supplemented with vitamin D3 1000 iu/d for 6 months. They reported that vitamin D3 supplementation varied with race and was more effective and significant in Black children[49] (Table 1). In addition, the intake required to maintain concentrations at 20 ng/ml in 97.5% of Black and White children, adjusting for race and pubertal status, was three times higher than the US recommended allowance of 600 iu/d[52]. Sachek et al.[53]) randomised control trial recruited 604 children (Black, Hispanic, White and Asian), aged 8–15 years. They found that serum vitamin D levels increased over 6 months with all dose ranges (600, 1000 or 2000 iu daily) with the 2000 iu daily dose achieving a higher vitamin D concentration compared with the two lower doses (33.1 v. 26.3 and 27.5 ng/ml; P < 0.001). Talib et al.[55] found that adolescents (Hispanic, Black, White and Asian; 13–20 years old) required 8 weeks of high-dose colecalciferol, of at least 5000 iu/d in order to correct deficiency status, noting also that obese adolescents had poorer response to treatment (13.7 +/− 10.7 v. 21.9 +/− 16.9 ng/ml; P < 0.001).

Abrams et al.[50] found no significant effect of an increase in vitamin D3, with decrease in parathyroid hormone levels, on calcium absorption, which was one of their study’s primary outcomes of interest. The pharmacological intervention used was supplementation with 1000 iu daily vitamin D for 8 weeks in African American, Hispanic, Asian, and White children aged between 4 and 9 years. Dong et al.[50] studied supplementation in forty-nine healthy African American children, aged 14–18 years, with 2000 iu/d v. 400 iu/d in the control group. They reported the 2000 iu daily dose as probably more effective (see Table 1) in optimising vitamin D status and counteracting progression of aortic stiffness in Black youth.

Doherty et al.[51] target group were African American children and young adults (5–20 years old) with or without sickle cell disease (SCD-SS). The study aim was to assess safety and efficacy of two oral vitamin D3 daily doses (4000 iu and 7000 iu). They noted that with 12 weeks of supplementation, both doses were safe and well tolerated but deficiency status (<20 ng/ml) was eliminated only in those receiving 7000 iu/d (P < 0.05).

Andersen et al.[56] recruited adolescent girls of Pakistani (South Asia) origin living in Denmark, aged 10–15 years, to study two vitamin D doses (10, 20 mcg/d). It is difficult to comment on any significant difference in 25(OH)D levels in the group after 1 year, as numbers recruited were small (n = 21). Gannmaa et al.[59,57] found that drinking ultra-high-temperature (UHT) processed milk fortified with about 100 iu vitamin D per serving improved levels in Mongolian (East Asia) children aged 9–11 years, and that correction of baseline deficiency in Mongolian children aged 12–15 years with 800 iu daily over 6 months increased growth[57]. Thatcher et al.[58] studied Nigerian (West Africa) children aged between 1 and 14 years with nutritional rickets, finding that calcium supplementation with or without vitamin D was more effective than vitamin D alone in healing active rickets (P < 0.001). (Tables 1, 6 and 7).

Research published after completion of the initial review

The updated literature review on MEDLINE, EMBASE, CINAHL and Cochrane Library databases, including secondary search of Google Scholar, undertaken for February 2018–November 2020, focused purely on identifying behavioural intervention studies. No publication with an explicitly defined behavioural intervention implemented to study vitamin D supplementation outcomes was identified (see Fig. 1). Protocols for two new systematic reviews have been published[60,67], but there is no indication that any behavioural interventions per se will be considered.

Discussion

This systematic review included twenty-four publications for final evaluation. We had planned to analyse behavioural interventions using HBM constructs[29] and the Behaviour Change Wheel framework criteria[30]. We have utilised HBM constructs for assessment in a previous patient and public involvement study[29].

Despite our a priori aim, we found no studies that explicitly defined and studied the effects of a behavioural intervention. Therefore, we completed a narrative evaluation of the two studies that undertook a pharmacological intervention and incorporated a behavioural component, which included translated information brochures, some support from health professionals and free supply of vitamin D supplements[38,42]. These studies did not evaluate which individual component helped improve vitamin D status in the infants, for example, whether it was the translated information brochure provided to the mothers or the free supplement supply. The combined strategies broadly meet criteria of education, training and enablement from the total of nine that underpin the Behaviour Change Wheel model[20].
### Table 6 Pharmacological vitamin D doses used in the trials included in systematic review and intervention outcome

| Reference               | Age (years) | Vitamin D supplement doses given and frequency | Intervention outcome positive/negative/other | Other study outcomes                                                                 |
|-------------------------|-------------|-----------------------------------------------|-------------------------------------------|-------------------------------------------------------------------------------------|
| Abrams *et al.* (2013)  | 4–9         | 1000 iu daily for 8 weeks                     | Positive                                  | No significant effects of change in vitamin D3 level on calcium absorption           |
| Andersen *et al.* 2008 | 10–15       | 10 mcg or 20 mcg/d dose over 1 year           | Positive                                  | Sample included women (18–53 years) and men 18–64 years.                           |
| Dong *et al.* (2010)   | 14–18       | 2000 iu v. 400 iu in control group over 16 weeks | Positive                                  | 2000 iu daily maybe effective in optimising vitamin D status and counteracting progression of aortic stiffness in Black youth. |
| Dougherty 2015          | 5–20        | 4000 iu/d v. 7000 iu/d over 12 weeks          | Positive                                  | Plasma 25(OH)D concentrations negatively modulated by adiposity                    |
| Green *et al.* (2015)  | Newborn     | 400 iu daily in addition to formula feed      | Positive                                  | In both groups of patients with sickle cell disease (SCD-SS), deficiency status eliminated only in those on 7000 iu/d dose. |
| Hollis *et al.* (2015)  | Pregnant women and newborns | Pregnant women on 400 iu vitamin D with additional 0 iu, 1600 iu or 3600 iu until delivery | Positive                                  | Maternal vitamin 25(OH)D levels, season, skin colour and ethnicity were not significant determinants of infant vitamin 25(OH)D levels. |
| Hollis *et al.* (2015)  | Newborn 4–6 weeks with breast-feeding planned for 6 months | 6400 iu vitamin D/d v. 400 iu/d          | Positive                                  | White infants had statistically significant higher mean 25(OH)D levels at discharge than non-White infants. |
| Hossain *et al.* 2014   | Pregnant women and newborns | 4000 iu/d from week 20 to delivery          | Positive                                  | 4000 iu/d for pregnant women safe and most effective in achieving sufficiency in all neonates as well regardless of race 6400 iu vitamin D dose daily was safe. Vitamin D deficiency in breast-fed infants affected by race. There was a positive correlation between maternal and neonatal 25(OH)D levels. Larger gains in vitamin D levels in White v. Black population at the highest dose. Translated information brochures and free supply of vitamin D supplement. No difference found in efficacy of both regimens when comparing diagnostic parameters of rickets at 12 weeks. Vitamin D insufficiency status persisted in 18% study group of African American children. Dose inadequate to raise blood levels to sufficiency status in both obese and non-obese African American children. Effect of supplementation varied by race and was more effective and significant in Black children. |
In general, there was a great deal of variability in the approaches of the studies. Several studies were undertaken in healthy populations (36, 42, 50, 54), including populations where the aim was to evaluate a large-scale national prevention campaign (38) or study the effect of fortification (39, 57). Other studies were undertaken in patients with medical conditions that precluded achievement of vitamin D sufficiency status (Table 7). Examples include a sickle cell anaemia – SS patient cohort (51), infants in intensive care (46), treatment for rickets (44, 58) or studying treatment in obesity (40, 53). Variation in study outcomes is to be expected due to heterogeneity within the studies, which used different thresholds for insufficiency and deficiency status when measuring serum vitamin D levels (7, 36, 40, 41, 50, 57).

Hanson et al. (46), Rodda et al. (45) and Yu et al. (7) undertook cord blood sampling, whereas Hollis et al. (43) evaluated maternal and infant urine and blood samples. The majority of the studies aimed to assess whether a vitamin D 400 iu daily dose was sufficient, using various surrogate endpoints to rationalise and justify their conclusions, with study of higher doses in studies that included patients with research group-defined deficiency/insufficiency status (Tables 1 and 7).

Cochrane reviews have shown that supplements taken during pregnancy could reduce the risks for pre-eclampsia, gestational diabetes and low birth weight (9, 59). The 2018 Food Standards Agency’s commissioned research on use of food supplements (60) found that consumers with higher levels of education or currently working were more likely to take supplements. The most common reason cited for taking food supplements was an aspiration for a healthy lifestyle. However, consumers who may be at risk may not be adequately informed about food supplements, to include vitamin D. Michie and Tayarachakul (61) suggest that any ‘information and advisory’ health professional role for pregnant women should include the recommendation of 400 iu vitamin D daily; with communication, especially with vulnerable mothers, and these should include mothers of darker skinned ethnic populations, undertaken and supported by midwives or community pharmacists (61). High-risk groups may need individualised advice and information on the need for vitamin D supplements. In addition, it may be important to address perceived medicalisation of vitamin D supplements, whereby women feel they only need to take a supplement on the recommendation or prescription from their GP (29). Currently in the UK, patients and the public are advised to purchase vitamin D, unless eligible for the Healthy Start Scheme (19). Even if prescribed per se, there are important issues that need to be addressed. Wan et al. (62) demonstrated an increase in prescribing of vitamin D in the UK, but their 2008–2016 database study found inconsistency between supply regimens prescribed, an absence of pre-supplementation (range 29–56 % annually), and a trend for increased prescribing of higher pharmacological treatment doses rather than maintenance doses. Global and national data (1–9, 11, 12, 16–18, 20, 59–61, 63) show there are still ongoing issues relating to vitamin D deficiency status and supplementation. More importantly,
studies have not adequately addressed the best approach to improve uptake in high-risk groups. This area requires further work to identify the most effective behavioural interventions, which should ideally be co-developed working with the high-risk community itself.

We used ethnicity as an inclusion criterion in order to consider at-risk groups of Asian or African origin. Both culture and religion can impact on uptake of supplements, but no study in this review discussed this complex relationship and effect on uptake, compliance or adherence with vitamin D supplementation. We noted variations in study design to support improved uptake of pharmacological treatment and these observations could be used as supporting information for future behavioural interventions. These modifications

Table 7  Vitamin D intervention study in healthy v. at-risk population as defined by research group

| References | Author defined study group | Ethnic group study | Classification of vitamin D status in study |
|------------|--------------------------|------------------|-------------------------------------------|
| Dong et al. (2010)[50] | Healthy group: Healthy | Yes | Sufficiency: 75 nmol/l |
| Abrams et al. (2013)[36] | Healthy | Yes | Insufficiency: more than 50 but less than 75 nmol/l |
| Thacher et al. (1999)[58] Rajakumar et al. (2005)[41] | Vitamin D at-risk group: Rickets or bone deformities At-risk vitamin D sample | Nigerian children, in Nigeria Yes | Deficiency: under 50 nmol/l |
| Rajakumar et al. (2008)[40] | Obese African American | Yes | High 32 ng/ml; 50–80 nmol/l |
| Mondal et al. (2014)[44] Dougherty 2015[51] | Nutritional rickets Sickle cell disease Study of two doses of vitamin D | Indian children, in India Yes | Middle 2–32 ng/ml; 50–80 nmol/l |
| Talib et al. (2016)[55] | Adolescents with vitamin D deficiency Pregnancy/neonate: Pregnancy | Yes | Low 20 ng/ml; 50 nmol/l |
| Yu et al. 2009[7] | | | (i) 12–20 ng/ml essential for bone health in children |
| Madar et al. (2009)[42] Hollis et al. (2011)[48] | Neonate Pregnancy/newborn | Yes | Insufficiency: 10–20 ng/ml Deficiency: < 10 ng/ml based on definition for adults |
| Hanson et al. (2011)[46] Mutlu 2011[38] | Infants/formula fed Infants | Yes | Suboptimal vitamin D status (<32 ng/ml) |
| Hossain et al. 2014[47] | Pregnancy/neonate | Yes | Deficiency 20 g/ml |
| Hollis et al. (2015)[43] | Mother–infant pairs and breast-feeding | Yes | Sufficient > 50 nmol/l |
| Green et al. (2015)[37] Rodda et al. (2015)[45] | Breast-fed infants Pregnancy/newborn vitamin D-deficient | Yes | Insufficiency: 25–50 nmol/l |
| Andersen et al. 2008[56] | Study of two doses of vitamin D | Yes | Deficiency: < 25 nmol/l |
| Gannmaa et al. (2008)[39] Lewis et al. (2013)[34] | Fortification Study of five oral doses in children | Mongolian children, in Mongolia Yes | Insufficiency as ≥50 to <80 nmol/l (≥20 to <32 ng/ml) |
| Rajakumar et al. (2015)[49] | Study of 1000 iu daily dose for 6 months | Yes | Deficiency <50 nmol/l (20 ng/ml) |
| Rajakumar et al. (2016)[52] | Study of 1000 iu daily dose for 6 months | Yes | |
| Gannmaa 2017[57] | Correction of baseline deficiency; fortification | Mongolian children, in Mongolia Yes | |
| Sacheck et al. (2017)[53] | Study of three doses of vitamin D | Yes | |
included use of chewable tablets (53), liquid (37,51) or flavoured formulations (36,53), drops for infants (37,38,251) and halal-certified products (45,56). Some of these modifications would be applicable in the case of all children, whereas some are important when considering ethnicity and culturally acceptable treatment.

The Palacios et al. (39) updated Cochrane ‘interventions’ systematic review aimed to evaluate the effect of vitamin D supplementation. With a focus on pregnancy only and not on at-risk ethnic minority groups per se, they assessed the evidence base for three distinct patient cohorts; vitamin D only (22 trials; 3725 pregnant women), calcium and vitamin D supplementation (9 trials; 1916 pregnant women) and vitamin D and other micronutrients (1 study; 1300 pregnant women). The studies considered pharmacological interventions evaluating risk or harm; no behavioural interventions were described. As in our review, they observed that supplementation increased serum 25(OH) D concentrations during pregnancy but with large heterogeneity in the results, possibly related to differences in vitamin D doses and methods used to assess outcomes. Another Cochrane review (39) considered variations in vitamin D supplement regimens during pregnancy, focusing on pregnancy and neonatal outcomes, but did not specifically consider at-risk ethnic minority groups and no behavioural interventions were reported.

Although we identified few publications in this systematic review, there is work that shows that milk fortification (35,53) appears to be acceptable to mothers, and chewable formulations (53) in addition to drops or liquid products (34,35,39,47) appear tolerable in children. This could have important implications for public policy on mass fortification. Variations in the available forms of supplements should be highlighted, especially to those groups less likely to present to health services.

Structural barriers will be important to overcome to ensure messaging reaches at-risk groups, including health literacy and low socio-economic status, especially where the targeted community comprises first-generation immigrants or residents. Other facets that deserve due consideration include individual communication preferences, comprehension and concerns, and research teams will need to seriously consider these with use of co-production methodologies. The NIHR INVOLVE guidance (24) defines the approach used by researchers, practitioners and the public working together on a co-produced research project as one of joint ownership, with sharing of power and responsibility from the start to the end of the project. Working together helps promote understanding by including all perspectives and skills. With close community engagement, there is an expectation that implementation of the co-designed intervention will result in better outcomes.

The two behavioural type strategies that we identified used elements of education, training and enablement as possible supportive behaviour change techniques, with the associated pharmacological intervention demonstrating achievement of significantly higher vitamin D levels. As these were mainly medical intervention trials, it is to be expected that the focus would be on participant adherence rather than on changing behaviour related to ongoing supplementation. However, bearing in mind racial/ethnic disparities, using a behavioural strategy may be important. We accepted that the search term ‘interventions’ would identify both pharmacological- and behavioural-type interventions, and we evaluated on the premise that there can be learning from any behavioural component incorporated within pharmacological trials as well. Future research groups could consider incorporating explicitly defined behavioural interventions, underpinned by HBM constructs and Behaviour Change Wheel categories, as a formal aspect of their intervention study. This may help address structural barriers and could be implemented, for example, during routine antenatal visits, for widespread population benefits. It is important to note the results of a 2020 study of 125 UK children with nutritional rickets, which found that over three-quarters (77.6%) of these children were not taking vitamin D supplements (43). Accepting that there is still a need for more data on children who did not have nutritional rickets, it is evident that current recommendations from Department of Health (1,3,19) are not reaching high-risk groups. Our aim to study behavioural interventions aiming to improve both initiation and maintenance of supplementation in pregnancy and beyond has been informed by this observation. The findings of our study, namely a lack of research into behavioural interventions and the forms of intervention that would be most effective in engaging at-risk groups, lead us to recommend this as a research priority.

Limitations of this review

Mutlu et al.’s study (38) describes a national campaign implemented in Turkey, aiming to encourage the entire population, including pregnant and nursing women and infants, to have adequate sunlight exposure. Although the primary source of vitamin D is sun exposure, the ability of the body to create and maintain sufficient levels is affected by many variables. These include socio-economic/socio-demographic status, geographical/environmental, and cultural and religious, lifestyle and dietary as well as genetic factors. Knoss et al. (68) studied people of different ethnic origins with similar practices, noting that skin pigmentation was a significant risk factor for vitamin D deficiency, irrespective of ultra-violet light exposure. A Swedish, primarily a food-based intervention study (69), did not report ethnic origin but recruited children living in the northern and southern latitudes of the country. With children stratified by skin colour using Fitzpatrick’s definition, the results showed that those with darker skin required higher vitamin D intake or supplementation. We focused on ethnicity to help identify behavioural interventions implemented, if any, but note that there may be other relevant work on sun exposure-related behavioural interventions in children. We used ethnicity as a primary search term to help identify high-risk groups, but studies reported their
selected populations based on ethnic origin or alternatively used race for classification. This adds to the heterogeneity of data available. No studies considered inter-related complexities including cultural or religious beliefs, distance from country of origin, or first- or second-generation immigrant status, and this needs due acknowledgement. The primary focus for all studies was outcome of a pharmacological intervention, rather than behavioural interventions, with different thresholds used to define vitamin D deficiency and extrapolation of findings should be undertaken cautiously. We only included studies published in English, and studies where full text was available, meaning some relevant studies may have been omitted.

Conclusions

In summary, all the studies included in our systematic review, which focused on high-risk ethnic groups aged under 18 years, evaluated the effect of a pharmacological vitamin D intervention. There were no studies identified that included direct evaluation of an explicitly defined behavioural intervention. There is a need for additional rigorous high-quality and larger RCT to evaluate the effects of vitamin D supplementation in pregnancy. Equally, there is a need for research into clearly defined behavioural interventions targeted to individual ethnic groups, ideally designed with use of co-production methodologies, in order to help our understanding of how behaviour change can affect vitamin D use in antenatal care, pregnancy and childhood.

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Supplementary material

For supplementary material accompanying this paper visit https://doi.org/10.1017/S1368980021000756

References

1. National Institute for Health and Care Excellence (NICE) (2016) Vitamin D: supplement use in specific population groups. https://www.nice.org.uk/guidance/ph56/chapter/5-Context (accessed May 2020).
2. Julies P, Lynn RM, Pall K et al. (2020) Nutritional rickets under 16 years: UK surveillance results. Arch Dis Child 105, 587–592.
3. Scientific Advisory Committee on Nutrition (2016) Vitamin D and health report. https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/537616/SACN_Vitamin_D_and_Health_report.pdf (accessed May 2020).
4. National Osteoporosis Society (2016) Vitamin D and Bone Health: A Practical Clinical Guideline for Patient Management. https://nos.org.uk/media/2073/vitamin-d-and-bone-health-adults.pdf (accessed May 2020).
5. Podd D (2015) Hypovitaminosis D: a common deficiency with pervasive consequences. J Am Acad PAs 28, 20–26.
6. Mulligan ML, Felton SK, Rick AE et al. (2010) Implications of vitamin D deficiency in pregnancy and lactation. Am J Obstet Gynecol 202, 429.e1–429.e9.
7. Yu CKH, Sykes L, Sethi M et al. (2009) Vitamin D deficiency and supplementation during pregnancy. Clin Endocrinol (Oxf) 70, 685–690.
8. De-Regil LM, Palacios C, Lombardo LK et al. (2016) Vitamin D supplementation for women during pregnancy. Cochrane Database Syst Rev, CD008873. doi: 10.1002/14651858. CD008873.pub3.
9. Palacios C, Kostikov LK & Peña-Rosas JP (2019) Vitamin D supplementation for women during pregnancy. Cochrane Database Syst Rev Issue 7, CD008873. doi: 10.1002/14651858.CD008873.pub4.
10. Theodoratou E, Tzoulaki I, Zgaga L et al. (2014) Vitamin D and multiple health outcomes: umbrella review of systematic reviews and meta-analyses of observational studies and randomised trials. BMJ 348, g2035.
11. Callaghan AL, Moy RJD, Booth IW et al. (2006) Incidence of symptomatic vitamin D deficiency. Arch Dis Child 91, 606–607.
12. Blair M (2016) Pregnant women need vitamin D for neonatal health. BMJ 355, i6685.
13. Haimi M & Kremer R (2017) Vitamin D deficiency/insufficiency from childhood to adulthood: insights from a sunny country. World J Clin Pediatr 6, 1–9.
14. Chhonkar A, Gupta A & Arya V (2018) Comparison of vitamin D level of children with severe early childhood caries and children with no caries. Int J Clin Pediatr Dent 11, 199–204.
15. Almoudi MM, Hussein AS, Abu Hassan MI et al. (2019) Dental caries and vitamin D status in children in Asia. Pediatr Int 61, 327–338.

16. Public Health England (2017) Results from Years 1–4 (combined) of the Rolling Programme (2008/2009–2011/12). https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/594560/NDNS_Y1_to_4_UK_report_executive_summary_revised_February_2017.pdf (accessed May 2020).

17. Gardner DB (2012) Vitamin D Supplementation among Women of Childbearing Age: Prevalence and Disparities. Thesis, University of Washington. https://depts.washington.edu/mchprog/docs/abstracts/2012/GardnerAbstract.pdf (accessed October 2020).

18. Gomes F, Bourassa MW, Adu-Afarwuah S et al. (2020) Setting research priorities on multiple micronutrient supplementation in pregnancy. Ann NY Acad Sci 1465, 76–88.

19. Davies SC, Jewell T, McBride M et al. (2012) Vitamin D – Advice on Supplementation for At Risk Groups. London: Department of Health.

20. National Institutes of Health Office of Dietary Supplements (2020) Vitamin D Fact Sheet for Health Professionals. https://ods.od.nih.gov/factsheets/VitaminD-HealthProfessional/ (accessed October 2020).

21. Brignden A, Parslow RM, Linney C et al. (2019) How are behavioural interventions delivered to children (5–11 years old): a systematic mapping review. BMJ Pediatr Open 3, e000543.

22. Johnson MRD, Bhopal RS, Ingleby JD et al. (2019) A glossary for the first world congress on migration, ethnicity, race and health. Public Health 172, 85–88.

23. Ford ME & Kelly PA (2005) Conceptualizing and categorizing race and ethnicity in health services research. Health Serv Res 40, 1658–1675.

24. INVOLVE (2019) Guidance on co-producing a research review. https://www.involve.org.uk/wp-content/uploads/2019/04/Coprod_Guidance_Feb19.pdf (accessed May 2020).

25. George S, Duran N & Norris K (2014) A systematic review of barriers and facilitators to minority research participation among African Americans, Latinos, Asian Americans, and Pacific Islanders. Am J Public Health 104, e16–e31.

26. Szczepura A (2005) Access to health care for ethnic minority populations. Postgrad Med J 81, 141.

27. Moher D, Liberati A, Tetzlaff J et al. (2009) Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. BMJ 339, b2535.

28. Rosenstock IM (1974) The health belief model and behavioural interventions delivered to children (5–8 years-old): a systematic mapping review. BMJ Pediatr Open 3, e1675–1679.

29. Green TJ, Li W, Barr SI et al. (2015) Vitamin D supplementation is associated with higher serum 25OHD in Asian and White infants living in Vancouver, Canada. Matern Child Nutr 11, 253–259.

30. Muthu GY, Kusal Y, Oszu E et al. (2011) Prevention of Vitamin D deficiency in infancy: daily 400 IU vitamin D is sufficient. Int J Pediatr Endocrinol 2011, 4.

31. Stanhope LE, Dietz WH & Service TD (2008) Obesity and vitamin D status in young children: results of an open-label, randomized controlled trial. Pediatrics 136, 625–634.

32. Mondal K, Seth A, Marwaha RK et al. (2014) A randomized controlled trial on safety and efficacy of single intramuscular versus staggered oral dose of 600 000 IU Vitamin D in treatment of nutritional rickets. J Trop Pediatr 60, 205–210.

33. Redda CP, Benson JE, Vincent AJ et al. (2015) Maternal vitamin D supplementation during pregnancy prevents vitamin D deficiency in the newborn: an open-label randomized controlled trial. Clin Endocrinol (Oxf) 83, 363–368.

34. Hanson C, Armass L, Suppes E et al. (2011) Vitamin D status and associations in newborn formula-fed infants during initial hospitalization. Am J Diet Assoc 111, 1836–1843.

35. Hossain N, Kanani FH, Ramirez S et al. (2014) Obstetric and neonatal outcomes of maternal vitamin D supplementation: results of an open-label, randomized controlled trial of antenatal vitamin D supplementation in Pakistani women. J Clin Endocrinol Metab 99, 2448–2455.

36. Hollis BW, Johnson D, Hulsey TC et al. (2011) Vitamin D supplementation during pregnancy: double-blind, randomized clinical trial of safety and effectiveness. J Bone Miner Res 26, 2341–2357.

37. Rajakumar K, Moore CG, Yahes J et al. (2015) Effect of Vitamin D3 supplementation in black and in white children: a randomized, placebo-controlled trial. J Clin Endocrinol Metab 100, 3183–3192.

38. Dong Y, Stollman-Jorgensen IS, Pollock NK et al. (2010) A 16-week randomized clinical trial of 2000 international units daily vitamin D3 supplementation in black youth: 25-hydroxyvitamin D, adiposity, and arterial stiffness. J Clin Endocrinol Metab 95, 4584–4591.

39. Dougherty KA, Bertolaso C, Schall JH et al. (2015) Safety and efficacy of high-dose daily vitamin D3 supplementation in children and young adults with sickle cell disease. J Pediatr Hematol Oncol 37, e308–e315.
52. Rajakumar K, Moore CG, Yabes J et al. (2016) Estimations of dietary vitamin D requirements in black and white children. Pediatr Res 80, 14–20.

53. Sachine JM, Van Rompay MI, Chomitz VR et al. (2017) Impact of three doses of Vitamin D3 on serum 25(OH)D deficiency and insufficiency in at-risk schoolchildren. J Clin Endocrinol Metab 102, 4496–4505.

54. Lewis RD, Laing EM, Hill Gallant KM et al. (2013) A randomized trial of vitamin D3 supplementation in children: dose-response effects on vitamin D metabolites and calcium absorption. J Clin Endocrinol Metab 98, 4816–4825.

55. Talib HJ, Ponnapakkam T, Gensure R et al. (2016) Treatment of vitamin D deficiency in predominantly hispanic and black adolescents: a randomized clinical trial. J Pediatr 170, 266.e1–272.e1.

56. Andersen R, Molgaard C, Skovgaard LT et al. (2008) Effect of vitamin D supplementation on bone and vitamin D status among Pakistani immigrants in Denmark: a randomised double-blinded placebo-controlled intervention study. Br J Nutr 100, 197–207.

57. Gannaa D, Stuart JJ, Sumberzul N et al. (2017) Vitamin D supplementation and growth in urban Mongol school children: results from two randomized clinical trials. PloS One 12, e0175237–e0175237.

58. Thacher TD, Fischer PR, Pettifor JM et al. (1999) A comparison of calcium, vitamin D, or both for nutritional rickets in Nigerian children. N Engl J Med 341, 563–568.

59. Palacios C, Trak-Fellermeier MA, Martinez RX et al. (2019) Regimens of vitamin D supplementation for women during pregnancy. Cochrane Database Syst Rev 10, CD013446.

60. Food Standards Agency (2018) Food Supplements Consumer Research Final Report 2018. https://www.food.gov.uk/sites/default/files/media/document/food-supplements-consumer-research.pdf (accessed October 2020).

61. Michie C & Tayarachakul S (2019) We should not write off vitamin D supplementation — especially in pregnancy. Pharm J 303, 7932. doi: 10.1211/PJ.2019.20207549.

62. Wan M, Horsfall LJ, Basatemedur E et al. (2019) Vitamin D prescribing in children in UK primary care practices: a population-based cohort study. BMJ Open 9, e031870.

63. Chibuzor MT, Graham-Kalio D, Osajiy JO et al. (2020) Vitamin D, calcium or a combination of vitamin D and calcium for the treatment of nutritional rickets in children. Cochrane Database Syst Rev 4, CD012581.

64. NICE (2009) Medicines adherence: involving patients in decisions about prescribed medicines and supporting adherence. Clinical guideline (CG76). https://www.nice.org.uk/guidance/cg76 (accessed October 2020).

65. NICE (2019) 2019 surveillance of medicines adherence (NICE guideline CG76) and medicines optimisation (NICE guideline NG5). Surveillance report. https://www.nice.org.uk/guidance/cg76/resources/2019-surveillance-of-medicines-adherence-nice-guideline-cg76-and-medicines-optimisation-nice-guideline-ng5-pdf-8714208880069 (accessed October 2020).

66. Yousef S, Elliot J, Manuel D et al. (2019) Study protocol: worldwide comparison of vitamin D status of immigrants from different ethnic origins and native-born populations – a systematic review and meta-analysis. Syst Rev 8, 211.

67. Best CM, Xu J, Patchen B et al. (2019). Vitamin D supplementation in pregnant or breastfeeding women or young children for preventing asthma. Cochrane Database Syst Rev 8, CD013396.

68. Knoss R, Halsey L & Reeves S (2012) Ethnic dress, vitamin D intake, and calcaneal bone health in young women in the United Kingdom. J Clin Densitom 15(2), 250–254. doi: 10.1016/j.jocd.2011.09.005.

69. Ohlund I, Lind T, Hernell O et al. (2017) Increased vitamin D intake differentiated according to skin color is needed to meet requirements in young Swedish children during winter: a double blind randomized clinical trial. Am J Clin Nutr 106, 105–112.

70. Biolab Medical Unit (2010) Vitamin D. https://www.biolab.co.uk/docs/vitd.pdf (accessed May 2020).