National Institute for Health and Care Excellence

Draft for consultation

Maternal and child Nutrition

[D] Evidence review for Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

NICE guideline number tbc

Evidence review underpinning research recommendation on optimum vitamin D dose during pregnancy for people with a BMI medically classified as overweight or obese in the NICE guideline

June 2024

Draft for consultation

These evidence reviews were developed by NICE



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ISBN:

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Optimum vitamin D dose during pregnancy

for those medically classified as

3 overweight or obese

4 Review question

- 5 What dose of vitamin D is appropriate during pregnancy for women with a BMI medically
- 6 classified as overweight or obese?

7 Introduction

Vitamin D demands increase during pregnancy and during lactation. There is a lack of scientific consensus on the amount of vitamin D intake that is deemed optimal in populations of pregnant women, regardless of additional risk factors that contribute to low vitamin D status, but current advice in the United Kingdom (UK) is to supplement with 10µg/day of vitamin D throughout pregnancy and during lactation.

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16 17 Observational studies have reported a relationship between low vitamin D status, pre-term birth, and low birth weight. There is suggestion that low vitamin D status is associated with gestational diabetes mellitus (GDM) and preeclampsia, although these relationships are not established. In the infant there is some suggestion of an association with low bone mass and poor respiratory function in childhood.

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- 20 While limited in number, observational studies have shown that pregnant women with body mass indexes (BMIs) above 29kg/m² have significantly lower vitamin D status than those with 21 a BMI ≤ 29k g/m². Vitamin D status of the neonate has been shown to be highly correlated 22 23 with maternal 25 hydroxyvitamin D (25(OH)D) concentrations. It has been suggested that 24 circulating 25(OH)D in newborns should be maintained above a minimum of 25–30 nmol/L. 25 There are no known thresholds for 25(OH)D concentrations in umbilical cord indicative of 26 improved infant health outcomes. The aim of this review is to assess what dose of vitamin D supplementation is appropriate during pregnancy for women medically classified as 27
- 28 overweight or obese.

Summary of the protocol

See Table 1 for a summary of the Population, Intervention, Comparison and Outcome (PICO) characteristics of this review.

32 Table 1: Summary of the protocol (PICO table)

Population	Women during a single or multiple pregnancy, who have a BMI medically classified as overweight or obese at booking or first scan
Intervention	 Low dose vitamin D supplementation (<10µg/400IU daily) Medium dose vitamin D supplementation (≥10µg/400IU to 20µg/800IU daily) High dose vitamin D supplementation (≥20µg/800IU daily) In combination or not with other vitamins and minerals
Comparison	 A different vitamin D supplementation dose daily (same dose category) A different vitamin D supplementation dose daily (different dose category)

	Standard care (as defined by the study)PlaceboNo intervention
Outcome	Critical Maternal serum 25(OH)D concentration Preterm birth (before 37 weeks gestational age) Low birth weight (<2500g) Important Pre-eclampsia Gestational diabetes Bone mass in infant Childhood respiratory function

- 1 25(OH)D: 25 hydroxyvitamin D; BMI: body mass index; IU: international unit
- 2 For further details see the review protocol in appendix A.

3 Methods and process

- 4 This evidence review was developed using the methods and process described in
- 5 Developing NICE guidelines: the manual. Methods specific to this review question are
- 6 described in the review protocol in appendix A and the methods document (supplementary
- 7 document 1).
- 8 Declarations of interest were recorded according to <u>NICE's conflicts of interest policy</u>.

9 Effectiveness evidence

10 Included studies

- 11 Two randomised trials were included for this review (Alhomaid 2021, Corcoy 2020).
- 12 The included studies are summarised in Table 2.
- 13 The study population included women with singleton pregnancies and mixed BMI population.
- However, only evidence on women with a BMI $\geq 25 \text{kg/m}^2$ was extracted from the studies and
- reported in this review. There was no evidence for women with multiple pregnancies.
- 16 There was evidence for 2 of the intervention groups stated in the protocol medium dose
- 17 vitamin D supplementation (≥10µg/ 400IU to 20µg/800IU daily) and high dose vitamin D
- 18 supplementation (≥20µg/800IU daily). One study compared medium dose vitamin D
- 19 supplementation of 10µg (400IU)/day with another medium dose vitamin D supplementation
- of 20µg (800IU)/day (Alhomaid 2021) and one study compared no vitamin D supplementation
- 21 to a high dose vitamin D supplementation of 40µg (1600IU)/day (Corcoy 2020). There was
- 22 no evidence for low dose vitamin D supplementation (<10µg/ 400IU daily) as defined in the
- 23 protocol, standard care or no intervention.
- 24 Two studies reported on maternal serum 25(OH)D concentration and 1 study reported on
- 25 preterm birth (before 37 weeks gestational age), low birth weight (<2500 g), pre-eclampsia
- and gestational diabetes. There was no evidence for bone mass in infant or childhood
- 27 respiratory function.
- 28 Evidence was stratified according to pre-specified BMI ranges where possible.
- 29 Pre-specified sub-group analysis (deprived socioeconomic groups, under/over 40 years of
- age, women and parents with disabilities, including learning disabilities and other physical

- and mental health conditions, women going through assisted conception, LGBTQ+ women
- and parents, children with developmental problems, geographical variation and religion and
- 3 cultural considerations) could not be conducted as there was no information within the
- 4 studies to conduct the analysis.
- 5 See the literature search strategy in appendix B and study selection flow chart in appendix C.

6 Excluded studies

- 7 Studies not included in this review are listed, and reasons for their exclusion are provided in
- 8 appendix K.

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Summary of included studies

Summaries of the studies that were included in this review are presented in Table 2.

11 Table 2: Summary of included studies.

Otrala			^	0
•				Outcomes
Study Alhomaid 2021 RCT Northern Ireland	Population N = 240 pregnant women Age in years [Mean (SD)]: Intervention (20μg/d) = 29.5 (5.5) Control (10μg/d) = 29.7 (5.1) Gestational age in weeks [Mean (SD)]: Intervention (20μg/d) = 12.8 (1.4) Control (10μg/d) = 13.0 (1.4) BMI in kg/m² [Mean (SD)]: Intervention (20μg/d) = 27.8 (5.4) Control (10μg/d) = 27.8 (5.4) Control (10μg/d) = 28.1 (5.7) Singleton	Intervention 20µg (800IU) vitamin D supplement from 12 weeks gestational age until birth 1 multivitamin tablet containing 10µg (400IU) vitamin D and 1 tablet containing 10µg (400IU) vitamin D	Comparison 10µg (400IU) vitamin D supplement from 12 weeks gestational age until birth 1 multivitamin tablet containing 10µg (400IU) vitamin D and 1 tablet containing 0µg vitamin D (placebo)	• Maternal serum 25(OH)D concentration at 28- and 36- weeks gestation – data was presented separately for overweight and obese women
	pregnancies only	49 (4999)		
Corcoy 2020 Cluster RCT	N = 154 pregnant women	40μg (1600IU)/day vitamin D3 supplement from ~	Placebo from ~ 15 weeks gestational age	 Maternal serum 25(OH)D concentration at
		15 weeks	until birth	24-28 weeks,

Study	Population	Intervention	Comparison	Outcomes
7 European countries • United Kingdom • Ireland • Australia • Poland • Italy • Spain • Belgium	Age in years [Mean (SD)]: Intervention (40μg/d) = 32.3 (5.2) Control (Placebo) = 32.8 (5.4) Gestational age in years [Mean (SD)]: Intervention (40μg/d) = 15.0 (2.9) Control (Placebo) = 15.4 (2.5) BMI in kg/m² [Mean (SD)]: Intervention (40μg/d) = 34.2 (4.3) Control (Placebo) = 34.2 (4.2) Singleton pregnancies only	gestational age until birth 4 tablets each containing 10µg (400IU) of vitamin D3	4 tablets identical to the intervention tablets	35-37 weeks and at delivery Preterm birth at delivery Low birth weight at delivery (<2500g) Preeclampsia at delivery Gestational diabetes mellitus at 24-28 weeks and 35-37 weeks

- 1 25(OH)D: 25 hydroxyvitamin D; BMI: body mass index; IU: international unit; RCT: randomised controlled trial
- 2 See the full evidence tables in appendix D. No meta-analysis was conducted (and so there
- 3 are no forest plots in appendix E).

4 Summary of the evidence

- 5 See appendix F for full GRADE tables.
- 6 The below paragraphs summarise the evidence for 2 comparisons: medium dose vitamin D
- 7 supplementation (≥10μg/ 400IU to 20μg/800IU daily) versus a different vitamin D
- 8 supplementation dose daily (same dose category) and high dose vitamin D supplementation
- 9 (≥20μg/800IU daily) versus placebo.
- 10 Comparison 1: Comparison between 20µg/day and 10µg/day vitamin D
- 11 supplementation in pregnant women medically classified as overweight or obese (BMI
- 12 ≥25.0 kg/m²) singleton pregnancies only
- One study was included in this comparison.
- 14 Overall 20µg (800IU)/day of vitamin D supplementation showed inconsistent findings for
- maternal serum 25(OH)D concentration when compared to 10µg (400IU)/day in overweight
- or obese women by gestational age. In women medically classified as overweight, the
- 17 evidence showed a possible important benefit at 36 weeks gestational age, and no important
- difference at 28 weeks gestational age. No important difference or no evidence of an
- important difference was found in women medically classified as obese at the same time

- 1 points. No other relevant outcomes were reported for this comparison group. The quality of
- 2 the evidence in this comparison was moderate.
- 3 Comparison 2: Comparison between 40µg/day vitamin D supplementation and no
- 4 supplementation in pregnant women medically classified as overweight or obese (BMI
- 5 ≥ 29.0kg/m²) singleton pregnancies only
- 6 One study was included in this comparison.
- 7 Overall 40µg (1600 IU)/day of vitamin D supplementation in pregnant women medically
- 8 classified as overweight or obese showed an important benefit over placebo in terms of
- 9 maternal serum 25(OH)D concentration at 24 to 28 and 35 to 37 weeks gestational age, and
- 10 at delivery. There was no evidence of important difference or no important difference for
- other outcomes such as preterm birth, low birth weight (<2.5kg), preeclampsia and
- 12 gestational diabetes. The quality of the evidence in this comparison was very low to
- 13 moderate.

14 Economic evidence

15 Included studies

- 16 No economic studies were identified which were applicable to this review question. See the
- 17 literature search strategy in appendix B and economic study selection flow chart in appendix
- 18 G.

19 Excluded studies

20 No economic studies were reviewed at full text and excluded from this review.

21 Economic model

- No economic modelling was undertaken for this review because the committee agreed that
- other topics were higher priorities for economic evaluation.

24 The committee's discussion and interpretation of the evidence

25 The outcomes that matter most

- 26 Maternal serum 25(OH)D concentration, preterm birth (before 37 weeks gestational age) and
- low birth weight (<2500g) were prioritised as critical outcomes by the committee. The main
- aim of this review was to determine the optimal dose of vitamin D during pregnancy for
- women medically classified as overweight and obese. Maternal serum 25(OH)D was
- prioritised as a critical outcome because it is a key measure of the vitamin D concentration in
- the blood, which will help to determine what dosage will achieve optimal blood vitamin D
- 32 levels. Preterm birth and low birth weight were prioritised as critical outcomes because these
- outcomes are crucial to the health of the neonate and are impacted by maternal vitamin D
- 34 levels.
- 35 Pre-eclampsia, gestational diabetes, bone mass in infants and childhood respiratory function
- 36 were prioritised as important outcomes as they are associated with vitamin D levels and may
- 37 have long-term effects on the mother and the infant. In addition, the committee recognised
- that preeclampsia and gestational diabetes have been prioritised as outcomes in published
- 39 literature. Bone mass in infants was agreed as an important outcome because bone
- 40 formation is directly associated with vitamin D and the fetus obtains vitamin D from the
- 41 mother. The committee acknowledged that there is evidence that shows the relationship
- 42 between vitamin D supplementation and reduced risk of severe asthma attacks but agreed

- that studies' follow up may not be long enough to capture this. Therefore, they agreed to
- 2 include any childhood respiratory function as an important outcome.
- 3 Evidence was available for all outcomes except for bone mass in infants and childhood
- 4 respiratory function.

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The quality of the evidence

- 6 The quality of the evidence was assessed using GRADE and ranged from moderate to very
- 7 low, with most of the evidence being of moderate quality. The main issues with the quality
- 8 were due to methodological risk of bias due to issues with randomisation and missing
- 9 outcome data, and bias arising from seriously imprecise findings. Studies were assessed for
- 10 quality using the Cochrane Risk of Bias (Rob) 2.0 tool for randomised studies and the
- 11 Cochrane RoB 2.0 Cluster randomised trials tool for cluster randomised studies.

12 Benefits and harms

- 13 The committee discussed the evidence from the review and agreed that whilst there was
- some evidence suggesting a benefit for vitamin D for some outcomes, given the low quality
- and uncertainty in evidence, the committee decided not to make any recommendations.
- 16 Evidence for 20μg (800IU) versus 10μg (400IU) of vitamin D supplementation showed that
- there was minimal increase in maternal serum 25(OH)D concentration for women medically
- 18 classified as overweight and no difference in women medically classified as obese. Therefore
- due to insufficient evidence and inconsistency in the evidence, the committee agreed not to
- 20 make any recommendations for this specific dosage.
- 21 The evidence for 40µg (1600IU) vitamin D supplementation showed that there was increased
- 22 maternal serum concentration levels at various timepoints during pregnancy and at birth but
- there was no difference for any other outcomes when compared to placebo. Therefore due to
- lack of sufficient evidence, the committee did not make any recommendation for this vitamin
- 25 D dose.

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- The committee from their knowledge and experience discussed that there was increased
- 27 insufficiency of vitamin D (vitamin D sufficiency in the UK is set at 25mmol) in pregnant
- women who are medically classified as overweight or obese. However, they agreed that the
- 29 evidence reviewed was not sufficient to make recommendations on optimum dose of vitamin
- D for pregnant women who are medically classified as overweight or obese. The committee
- 31 agreed that further research should be undertaken in this area to determine the optimum
- dose of vitamin D during pregnancy for women with a BMI medically classified as overweight
- or obese. They therefore developed a research recommendation to inform future guidance
- 34 (see appendix K for full details of the research recommendation).

Cost effectiveness and resource use

- 36 No economic evidence was identified for this review. As the committee made no
- 37 recommendations relevant to this review question, there are no resource implications.

Other factors the committee took into account

- For this review question, the population in the evidence was women and no evidence was
- identified or reviewed for trans men or non-binary people. The protocol and literature
- searches were not designed to specifically look for evidence on trans men or non-binary
- 42 people but they were also not excluded. However, there is a small chance evidence on them
- 43 may not have been captured, if such evidence exists. In discussing the evidence, the
- committee considered whether the recommendations could apply to a broader population,
- 45 and used gender inclusive language to promote equity, respect and effective communication
- with everyone. Healthcare professionals should use their clinical judgement when

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- 1 implementing the recommendations, taking into account each person's circumstances, needs
- and preferences, and ensuring all people are treated with dignity and respect throughout
- 3 their care.

4 Recommendations supported by this evidence review

- 5 This evidence review supports the research recommendation on optimum vitamin D dose
- 6 during pregnancy for people with a BMI medically classified as overweight or obese.

7 References – included studies

- 8 **Alhomaid 2021**
- 9 Alhomaid, Raghad M, Mulhern, Maria S, Strain, Jj et al. (2021) Maternal obesity and baseline vitamin
- D insufficiency alter the response to vitamin D supplementation: a double-blind, randomized trial in
- 11 pregnant women. The American journal of clinical nutrition 114(3): 1208-1218
- 12 Corcoy 2020
- 13 Corcoy, Rosa, Mendoza, Lilian C, Simmons, David et al. (2020) The DALI vitamin D randomized
- controlled trial for gestational diabetes mellitus prevention: No major benefit shown besides vitamin D
- sufficiency. Clinical nutrition (Edinburgh, Scotland) 39(3): 976-984

Appendices

2 Appendix A Review protocols

- Review protocol for review question: What dose of vitamin D is appropriate during pregnancy for women with a BMI
- 4 medically classified as overweight or obese?

5 Table 3: Review protocol

Field	Content
PROSPERO registration number	CRD42022346781
Review title	Optimum vitamin D dose during pregnancy for women medically classified as overweight or obese
Review question	What dose of vitamin D is appropriate during pregnancy for women with a BMI medically classified as overweight or obese?
Objective	To determine the optimal vitamin D dose during pregnancy for women medically classified as overweight or obese.
Searches	The following databases will be searched:
	Cochrane Central Register of Controlled Trials (CENTRAL)
	Cochrane Database of Systematic Reviews (CDSR)
	Embase
	MEDLINE
	Epistemonikos
	• CINAHL
	International HTA Database
	Health Technology Assessment (HTA) database (last updated October 2016)
	Searches will be restricted by:
	No date restriction
	English language only

Field	Content
	Human studies only
	Other searches:
	Inclusion lists of systematic reviews
	The full search strategies for MEDLINE database will be published in the final review. For each search, the principal database search strategy is quality assured by a second information scientist using an adaptation of the PRESS 2015 Guideline Evidence-Based Checklist.
Condition or domain being studied	Vitamin D supplementation
Population	 Women during a single or multiple pregnancy, who have a BMI medically classified as overweight or obese (in line with <u>NICE guideline on Obesity: identification and classification of overweight and obesity</u> [update]) at booking or first scan
	Note: other anthropometric measures, such as weight to height ratio, that have been reported in the <u>NICE</u> <u>guidance on Obesity: identification and classification of overweight and obesity (update)</u> will be considered
	Note: if BMI is measured after the 1st trimester, the study will be included if eligible, but downgraded for indirectness.
Intervention	 Low dose vitamin D supplementation (<10μg/ 400IU daily)
	 Medium dose vitamin D supplementation (≥10μg/ 400IU to <20μg/800IU daily)
	 High dose vitamin D supplementation (≥20μg/800IU daily)
	In combination or not with other vitamins and minerals
Comparator	A different vitamin D supplementation dose daily (same dose category)
	A different vitamin D supplementation dose daily (different dose category)
	Standard care (as defined by the study)
	Placebo
	No intervention

Field	Content
Types of study to be included	Include published full-text papers: Systematic reviews of RCTs Parallel RCTs If insufficient parallel RCTs*: Quasi-randomised controlled trials Non-randomised controlled trials/prospective cohort studies Retrospective cohort studies Historically controlled studies *Non-randomised studies will be considered for inclusion if insufficient RCT evidence is available for guideline decision making. Sufficiency will be judged taking into account factors including number/quality/sample size of RCTs, outcomes reported and availability of data from subgroups of interest. Non-randomised studies will only be included if they adjust for confounding factors in the analysis. Conference abstracts will not be included because these do not typically have sufficient information to allow full critical appraisal.
Other exclusion criteria	Setting: • Countries other than high income countries (as defined by the OECD) If any study or systematic review includes <1/3 of women who received care in the above setting, it will be considered for inclusion but, if included, the evidence will be downgraded for indirectness.
Context	The population of this guideline may overlap with the population of women included in other NICE guidelines (such as postnatal care, antenatal care, intrapartum care, pregnancy and complex social factors or obesity prevention).
Primary outcomes (critical outcomes)	 Maternal serum 25(OH)D concentration Preterm birth (before 37 weeks gestational age) Low birth weight (<2500 g)

Field	Content
Secondary outcomes (important outcomes)	 Pre-eclampsia Gestational diabetes Bone mass in infant Childhood respiratory function
Data extraction (selection and coding)	All references identified by the searches and from other sources will be uploaded into EPPI and deduplicated. Titles and abstracts of the retrieved citations will be screened to identify studies that potentially meet the inclusion criteria outlined in the review protocol. Duplicate screening will not be undertaken for this question. Full versions of the selected studies will be obtained for assessment. Studies that fail to meet the inclusion criteria once the full version has been checked will be excluded at this stage. Each study excluded after checking the full version will be listed, along with the reason for its exclusion. A standardised form will be used to extract data from studies. The following data will be extracted: study details (reference, country where study was carried out, type and dates), participant characteristics, inclusion and exclusion criteria, details of the interventions if relevant, setting and follow-up, relevant outcome data and source of funding. One reviewer will extract relevant data into a standardised form, and this will be quality assessed by a senior reviewer.
Risk of bias (quality) assessment	Quality assessment of individual studies will be performed using the following checklists: ROBIS tool for systematic reviews Cochrane RoB tool v.2 for RCTs and quasi-RCTs Cochrane ROBINS-I tool for non-randomised (clinical) controlled trials The quality assessment will be performed by one reviewer and this will be quality assessed by a senior reviewer.
Strategy for data synthesis	Quantitative findings will be formally summarised in the review. Where multiple studies report on the same outcome for the same comparison, meta-analyses will be conducted using Cochrane Review Manager software. A fixed effect meta-analysis will be conducted and data will be presented as risk ratios if possible or odds ratios when required (for example, if only available in this form in included studies) for dichotomous

Field	Content
	outcomes, and mean differences or standardised mean differences for continuous outcomes. Heterogeneity in the effect estimates of the individual studies will be assessed using the I2 statistic. Alongside visual inspection of the point estimates and confidence intervals, I2 values of greater than 50% and 80% will be considered as significant and very significant heterogeneity, respectively. Heterogeneity will be explored as appropriate using sensitivity analyses and pre-specified subgroup analyses. If heterogeneity cannot be explained through subgroup analysis then a random effects model will be used for meta-analysis, or the data will not be pooled.
	The confidence in the findings across all available evidence will be evaluated for each outcome using an adaptation of the 'Grading of Recommendations Assessment, Development and Evaluation (GRADE) toolbox' developed by the international GRADE working group: http://www.gradeworkinggroup.org/ Minimally important differences:
	 Preterm birth (before 37 weeks gestational age), low birth weight (<2500g), pre-eclampsia, gestational diabetes, childhood respiratory function: statistical significance
	Validated scales/continuous outcomes: published MIDs where available
	 All other outcomes & where published MIDs are not available: 0.8 and 1.25 for all relative dichotomous outcomes; +/- 0.5x control group SD for continuous outcomes
Analysis of subgroups	Evidence will be stratified by:
	BMI thresholds on booking:
	o Overweight range: 25 to 29.99 kg/m²
	o Obesity range 1: 30 to 34.99 kg/m²
	o Obesity range 2: 35 to 39.99 kg/m ²
	o Obesity range 3: >40 kg/m²
	 Follow the <u>NICE guidance on Obesity: identification and classification of overweight and obesity</u> (<u>update</u>) for people with a South Asian, Chinese, other Asian, Middle Eastern, Black 24 African or African-Caribbean family background
	Single versus multiple pregnancy
	Comorbidities (yes versus no)
	Season (summer versus winter)
	• Ethnicity
	○ White/White British

Field	Content
	∘ Asian/Asian British
	○ Black/African/Caribbean/Black British
	 Mixed/Multiple ethnic groups
	o Other ethnic group
	Evidence will be sub-grouped by the following only in the event that there is significant heterogeneity in outcomes:
	Deprived socioeconomic group
	• Age
	○ Under 40 years of age
	○ Over 40 years of age
	 Women and parents with disabilities, including learning disabilities and other physical and mental health conditions
	Women going through assisted conception
	LGBTQ+ women and parents
	Children with developmental problems
	• Geographical variation e.g. places without adequate provision of primary care (outside cities).
	Religion and cultural considerations
	Where evidence is stratified or sub-grouped the committee will consider on a case by case basis if separate recommendations should be made for distinct groups. Separate recommendations may be made where there is evidence of a differential effect of interventions in distinct groups. If there is a lack of evidence in one group, the committee will consider, based on their experience, whether it is reasonable to extrapolate and assume the interventions will have similar effects in that group compared with others.
Type and method of review	
	□ Diagnostic
	□ Prognostic
	☐ Qualitative
	□ Epidemiologic
	- Epideimologio

Field	Content		
	□ Service	Delivery	
	☐ Other (p	lease specif	y)
Language	English		
Country	England		
Anticipated or actual start date	14/07/2022		
Anticipated completion date	22/11/2023		
Stage of review at time of this	Review stage	Started	Completed
submission	Preliminary searches		
	Piloting of the study selection process		
	Formal screening of search results against eligibility criteria		
	Data extraction		\boxtimes
	Risk of bias (quality) assessment		
	Data analysis		\boxtimes
Named contact	5a. Named contact National Institute 5b. Named contact mandcoutrition@r	for Health ar ct e-mail <u>nice.org.uk</u>	

Field	Content
	National Institute for Health and Care Excellence (NICE)
Review team members	From the National Guideline Alliance: • Senior Systematic Reviewer • Systematic Reviewer
Funding sources/sponsor	This systematic review is being completed by the National Institute for Health and Care Excellence
Conflicts of interest	All guideline committee members and anyone who has direct input into NICE guidelines (including the evidence review team and expert witnesses) must declare any potential conflicts of interest in line with NICE's code of practice for declaring and dealing with conflicts of interest. Any relevant interests, or changes to interests, will also be declared publicly at the start of each guideline committee meeting. Before each meeting, any potential conflicts of interest will be considered by the guideline committee Chair and a senior member of the development team. Any decisions to exclude a person from all or part of a meeting will be documented. Any changes to a member's declaration of interests will be recorded in the minutes of the meeting. Declarations of interests will be published with the final guideline.
Collaborators	Development of this systematic review will be overseen by an advisory committee who will use the review to inform the development of evidence-based recommendations in line with section 3 of Developing NICE guidelines: the manual . Members of the guideline committee are available on the NICE website: https://www.nice.org.uk/guidance/indevelopment/gid-ng10191
Other registration details	None
URL for published protocol	https://www.crd.york.ac.uk/prospero/display_record.php?RecordID=346781
Dissemination plans	NICE may use a range of different methods to raise awareness of the guideline. These include standard approaches such as: notifying registered stakeholders of publication publicising the guideline through NICE's newsletter and alerts issuing a press release or briefing as appropriate, posting news articles on the NICE website, using social media channels, and publicising the guideline within NICE.
Keywords	Vitamin D, pregnancy, overweight, obesity
Details of existing review of same topic by same authors	Not applicable

Field	Content
Current review status	
	□ Completed but not published
	□ Completed and published
	☐ Completed, published and being updated
	□ Discontinued
Additional information	None
Details of final publication	www.nice.org.uk

BMI: body mass index; CDSR: Cochrane Database of Systematic Reviews; CENTRAL: Cochrane Central Register of Controlled Trials; EPPI: Evidence for policy and practice information; GRADE: Grading of Recommendations Assessment, Development and Evaluation; HTA: Health Technology Assessment; MID: minimally important difference; NGA: National Guideline Alliance; NICE: National Institute for Health and Care Excellence; OECD: organisation for economic cooperation and development; (OH)D: hydroxyvitamin D; RCT: randomised controlled trial; RoB: risk of bias; ROBINS-I: risk of bias in non-randomised studies; ROBIS: risk of bias in systematic reviews; SD: standard deviation

Appendix B Literature search strategies

Literature search strategies for review question: What dose of vitamin D is appropriate during pregnancy for women with a BMI medically classified as overweight or obese?

Effectiveness Searches

Databases: MEDLINE

maternai/ or obesity, metabolically benign/ or obesity, morbid/ or waist circumference/ or waist-hip ratio/ or Waist-Heigit Ratio/ or Weight Gain/ or Skinfold Thickness/ or Body Fat Distribution/ exp Adipose Tissue/ (obes* or overweight or over weight or corpulen* or heavy or heavier or fat or adipos* or (weight adj2 (manag* or gain* or increas* or excess or chang*))).ti,ab,kf. (body mass index or BMI or quetelet index).ti,ab,kf. (waist circumference* or waist hip ratio* or waist height ratio* or weight height ratio*),ti,ab,kf. (skin fold* or skinfold* or body composition or (body fat adj3 percent*)).ti,ab,kf. ((arm or midarm or upperarm or brachial) adj4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)).ti,ab,kf. 10 or/4-10 12 3 and 11 13 exp Vitamin D/ (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calciftriol* or 24,25-dihydroxyvitamin D* or ergocalciferol* or ergosterol* or viosterol or vitamin d* or vitamind* or 25 hydroxy* or 25-?OH*).ti,ab,kf. 11 or/13-14 12 and 15 12 letter/ editorial/ news/ exp historical article/ 23 case report/ (letter or comment*).ti. or/17-24 randomized controlled trial/ or random*.ti,ab. 25 not 26 animals/ not humans/ exp Animals Laboratory/ exp Animals Laboratory/ exp Animals Laboratory/ exp Models, Animal/ exp Rodentia/	Date	of last search: 05/12/2023
2 (antenatal* or ante-natal* or gestational or maternal* or pregnan* or prenatal* or pre-natal*), ti, ab, kf. 1 or 2 4 adiposity/ or body mass index/ or body size/ or body weight/ or overweight/ or obesity, abdominal/ or obesity maternal/ or obesity, metabolically benign/ or obesity, morbid/ or waist circumference/ or waist-hip ratio/ or Waist-Heigh Ratio/ or Weight Gain/ or Skinfold Thickness/ or Body Fat Distribution/ exp Adipose Tissue/ (obes* or overweight or over weight or corpulen* or heavy or heavier or fat or adipos* or (weight adj2 (manag* or gain* or increas* or excess or chang*)), ti, ab, kf. (waist circumference* or waist hip ratio* or waist height ratio* or weight height ratio*), ti, ab, kf. (skin fold* or skinfold* or body composition or (body fat adj3 percent*)), ti, ab, kf. ((arm or midarm or upperarm or brachial) adj4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)), ti, ab, kf. (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24, 25-dihydroxyvitamin D* or ergocalciferol* or ergosterol* or viosterol or vitamin d* or vitamind* or 25 hydroxy* or 25-70H*), ti, ab, kf. 12 and 15 letter/ editorial/ news/ exp historical article/ Anecdotes as Topic/ comment/ 23 case report/ 4 (letter or comment*), ti. 6 or/17-24 6 randomized controlled trial/ or random*. ti, ab. 7 25 not 26 8 animals/ not humans/ exp Animals, Laboratory/ exp Animals, Laboratory/ exp Animals, Laboratory/ exp Rodentia/ exp Rodentia/	#	Searches
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28 animals/ not humans/ 29 exp Animals, Laboratory/ 30 exp Animal Experimentation/ 31 exp Models, Animal/ 32 exp Rodentia/	26	randomized controlled trial/ or random*.ti,ab.
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32 exp Rodentia/	30	exp Animal Experimentation/
	31	exp Models, Animal/
	32	exp Rodentia/
33 (rat or rats or mouse or mice).ti.		(rat or rats or mouse or mice).ti.
34 or/27-33	34	or/27-33

#	Searches
35	16 not 34
36	limit 35 to English language
37	randomized controlled trial.pt.
38	controlled clinical trial.pt.
39	pragmatic clinical trial.pt.
40	randomi#ed.ab.
41	placebo.ab.
42	drug therapy.fs.
43	randomly.ab.
44	trial.ab.
45	groups.ab.
46	or/37-45
47	Meta-Analysis/
48	Meta-Analysis as Topic/
49	(meta analy* or metanaly* or metaanaly*).ti,ab.
50	((systematic* or evidence*) adj2 (review* or overview*)).ti,ab.
51	(reference list* or bibliograph* or hand search* or manual search* or relevant journals).ab.
52	(search strategy or search criteria or systematic search or study selection or data extraction).ab.
53	(search* adj4 literature).ab.
54	(medline or pubmed or cochrane or embase or psychlit or psychinfo or psycinfo or cinahl or science citation index or bids or cancerlit).ab.
55	cochrane.jw.
56	or/47-55
57	36 and (46 or 56)
58	Observational Studies as Topic/
59	Observational Study/
60	Epidemiologic Studies/
61	exp Case-Control Studies/
62	exp Cohort Studies/
63	Cross-Sectional Studies/
64	Controlled Before-After Studies/
65	Historically Controlled Study/
66	Interrupted Time Series Analysis/
67	Comparative Study.pt.
68	case control\$.tw.
69	case series.tw.
70	(cohort adj (study or studies)).tw.
71	cohort analy\$.tw.
72	(follow up adj (study or studies)).tw.
73	(observational adj (study or studies)).tw.
74	longitudinal.tw.
75	prospective.tw.
76	retrospective.tw.
77	cross sectional.tw.
78	or/58-77
79	36 and 78
80	afghanistan/ or africa/ or africa, northern/ or africa, central/ or africa, eastern/ or "africa south of the sahara"/ or africa, southern/ or africa, western/ or albania/ or algeria/ or andorra/ or angola/ or "antigua and barbuda"/ or argentina/ or armenia/ or azerbaijan/ or bahamas/ or bahrain/ or bangladesh/ or barbados/ or belize/ or benin/ or bhutan/ or bolivia/ or borneo/ or "bosnia and herzegovina"/ or botswana/ or brazil/ or brunei/ or bulgaria/ or burkina faso/ or burundi/ or cabo verde/ or cambodia/ or cameroon/ or central african republic/ or chad/ or exp china/ or comoros/ or congo/ or cote d'ivoire/ or croatia/ or cuba/ or "democratic republic of the congo"/ or cyprus/ or dibouti/ or dominica/ or dominical

d'ivoire/ or croatia/ or cuba/ or "democratic republic of the congo"/ or cyprus/ or djibouti/ or dominica/ or dominica/

Searches

republic/ or ecuador/ or egypt/ or el salvador/ or equatorial guinea/ or eritrea/ or eswatini/ or ethiopia/ or fiji/ or gabon/ or gambia/ or "georgia (republic)"/ or ghana/ or grenada/ or guatemala/ or guinea/ or guinea-bissau/ or guyana/ or haiti/ or honduras/ or independent state of samoa/ or exp india/ or indian ocean islands/ or indochina/ or indonesia/ or iran/ or iraq/ or jamaica/ or jordan/ or kazakhstan/ or kenya/ or kosovo/ or kuwait/ or kyrgyzstan/ or laos/ or lebanon/ or liechtenstein/ or lesotho/ or liberia/ or libya/ or madagascar/ or malaysia/ or malawi/ or mali/ or malar/ or mauritania/ or mauritius/ or mekong valley/ or melanesia/ or micronesia/ or monaco/ or mongolia/ or montenegro/ or morocco/ or mozambique/ or myanmar/ or namibia/ or nepal/ or nicaragua/ or niger/ or nigeria/ or oman/ or pakistan/ or palau/ or exp panama/ or papua new guinea/ or paraguay/ or peru/ or philippines/ or qatar/ or "republic of belarus"/ or "republic of north macedonia"/ or romania/ or exp russia/ or rwanda/ or "saint kitts and nevis"/ or saint lucia/ or "saint vincent and the grenadines"/ or "sao tome and principe"/ or saudi arabia/ or serbia/ or sierra leone/ or senegal/ or seychelles/ or singapore/ or somalia/ or south africa/ or south sudan/ or sri lanka/ or sudan/ or suriname/ or syria/ or taiwan/ or tajikistan/ or tanzania/ or thailand/ or timor-leste/ or togo/ or tonga/ or "trinidad and tobago"/ or tunisia/ or turkmenistan/ or uganda/ or ukraine/ or united arab emirates/ or uruguay/ or uzbekistan/ or vanuatu/ or venezuela/ or vietnam/ or west indies/ or yemen/ or zambia/ or zimbabwe/

- 81 "organisation for economic co-operation and development"/
- australasia/ or exp australia/ or austria/ or baltic states/ or belgium/ or exp canada/ or chile/ or colombia/ or costa rica/ or czech republic/ or exp denmark/ or estonia/ or europe/ or finland/ or exp france/ or exp germany/ or greece/ or hungary/ or iceland/ or ireland/ or israel/ or exp italy/ or exp japan/ or korea/ or latvia/ or lithuania/ or luxembourg/ or mexico/ or netherlands/ or new zealand/ or north america/ or exp norway/ or poland/ or portugal/ or exp "republic of korea"/ or "scandinavian and nordic countries"/ or slovakia/ or slovenia/ or spain/ or sweden/ or switzerland/ or turkey/ or exp united kingdom/ or exp united states/
- 83 european union/
- 84 developed countries/
- 85 or/81-84
- 86 80 not 85
- 87 57 not 86
- 88 79 not 86

Databases: Embase

#	Searches
1	exp pregnancy/ or pregnant woman/ or prenatal care/ or prenatal period/
2	(antenatal* or ante natal* or gestation* or maternal* or mother* or pregnan* or prenatal* or pre natal*).ti,ab,kf.
3	1 or 2
4	body mass/ or body size/ or body weight/ or body weight gain/ or obesity/ or abdominal obesity/ or fat mass/ or maternal obesity/ or metabolically benign obesity/ or morbid obesity/ or normal weight obesity/ or waist circumference/ or waist hip ratio/ or waist to height ratio/ or weight height ratio/ or skinfold thickness/ or body fat distribution/
5	exp adipose tissue/ or exp obese patient/
6	(obes* or overweight or over weight or corpulen* or heavy or heavier or fat or adipos* or (weight adj2 (manag* or gain* or increas* or excess or chang*))).ti,ab,kf.
7	(body mass index or BMI or quetelet index).ti,ab,kf.
8	(waist circumference* or waist hip ratio* or waist height ratio* or weight height ratio*).ti,ab,kf.
9	(skin fold* or skinfold* or body composition or (body fat adj3 percent*)).ti,ab,kf.
10	((arm or midarm or upperarm or brachial) adj4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)).ti,ab,kf.
11	or/4-10
12	3 and 11
13	exp vitamin D/
14	(calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24,25-dihydroxyvitamin D* or ergocalciferol* or ergosterol* or viosterol or vitamin d* or vitamind* or 25 hydroxy* or 25-?OH*).ti,ab,kf.
15	or/13-14
16	12 and 15
17	letter.pt. or letter/
18	note.pt.
19	editorial.pt.

#	Searches
20	case report/ or case study/
21	(letter or comment*).ti.
22	or/17-21
23	randomized controlled trial/ or random*.ti,ab.
24	22 not 23
25	animal/ not human/
26	nonhuman/
27	exp Animal Experiment/
28	exp Experimental Animal/
29	animal model/
30	exp Rodent/
31	(rat or rats or mouse or mice or rodent*).ti.
32	or/24-31
33	16 not 32
34	(conference abstract* or conference review or conference paper or conference proceeding).db,pt,su.
35	33 not 34
36	limit 35 to English language
37	random*.ti,ab.
38	factorial*.ti,ab.
39	(crossover* or cross over*).ti,ab.
40	((doubl* or singl*) adj blind*).ti,ab.
41	(assign* or allocat* or volunteer* or placebo*).ti,ab.
42	crossover procedure/
43	single blind procedure/
44	randomized controlled trial/
45	double blind procedure/
46	or/37-45
47	systematic review/
48	meta-analysis/
49	(meta analy* or metanaly* or metaanaly*).ti,ab.
50	((systematic or evidence) adj2 (review* or overview*)).ti,ab.
51	(reference list* or bibliograph* or hand search* or manual search* or relevant journals).ab.
52	(search strategy or search criteria or systematic search or study selection or data extraction).ab.
53	(search* adj4 literature).ab.
54	(medline or pubmed or cochrane or embase or psychlit or psyclit or psychinfo or psycinfo or cinahl or science citation index or bids or cancerlit).ab.
55	((pool* or combined) adj2 (data or trials or studies or results)).ab.
56	cochrane.jw.
57	or/47-56
58	36 and (46 or 57)
59	Clinical study/
60	Case control study/
61	Family study/
62	Longitudinal study/
63	Retrospective study/
64	comparative study/
65	Prospective study/
66	Randomized controlled trials/
67	65 not 66
68	Cohort analysis/
69	cohort analy\$.tw.

#	Searches
70	(Cohort adj (study or studies)).tw.
71	(Case control\$ adj (study or studies)).tw.
72	(follow up adj (study or studies)).tw.
73	(observational adj (study or studies)).tw.
74	(epidemiologic\$ adj (study or studies)).tw.
75	(cross sectional adj (study or studies)).tw.
76	case series.tw.
77	prospective.tw.
78	retrospective.tw.
79	or/59-64,67-78
80	36 and 79
81	afghanistan/ or africa/ or "africa south of the sahara"/ or albania/ or algeria/ or andorra/ or angola/ or argentina/ or "antigua and barbuda"/ or armenia/ or exp azerbaijan/ or bahamas/ or bahrain/ or bangladesh/ or barbados/ or belarus/ or belize/ or benin/ or bhutan/ or bolivia/ or borneo/ or exp "bosnia and herzegovina"/ or botswana/ or exp brazil/ or brunei darussalam/ or blugaria/ or burkina faso/ or burundi/ or cambodia/ or cameroon/ or cape verde/ or central africa/ or central african republic/ or chad/ or exp china/ or comoros/ or congo/ or cook islands/ or cote d'ivoire/ or croatia/ or cuba/ or cyprus/ or democratic republic congo/ or djibouti/ or dominica/ or dominican republic/ or ecuador/ or el salvador/ or egypt/ or equatorial guinea/ or eritrea/ or eswatini/ or ethiopia/ or exp "federated states of micronesia"/ or fiji/ or gabon/ or gambia/ or exp "georgia (republic)"/ or ghana/ or grenada/ or guatemala/ or guinea/ or guinea-bissau/ or guyana/ or haiti/ or honduras/ or exp india/ or exp indonesia/ or iran/ or exp iraq/ or jamaica/ or jordan/ or kazakhstan/ or kenya/ or kiribati/ or kosovo/ or kuwait/ or kyrgyzstan/ or laos/ or lebanon/ or liechtenstein/ or lesotho/ or liberia/ or libyan arab jamahiriya/ or madagascar/ or malawi/ or exp malaysia/ or maldives/ or mali/ or malta/ or mauritania/ or mauritius/ or melanesia/ or moldova/ or monaco/ or mongolia/ or "montenegro (republic)"/ or morocco/ or mozambique/ or myanmar/ or namibia/ or nauru/ or nepal/ or nicaragua/ or niger/ or nigeria/ or niue/ or north africa/ or oman/ or exp pakistan/ or palau/ or palestine/ or panama/ or papua new guinea/ or paraguay/ or peru/ or philippines/ or polynesia/ or qatar/ or "republic of north macedonia"/ or romania/ or exp russian federation/ or rwanda/ or sahel/ or "saint kitts and nevis"/ or "saint lucia"/ or "saint vincent and the grenadines"/ or saudi arabia/ or sexp somalia/ or south africa/ or south asia/ or south sudan/ or exp southeast asia/ or sri lanka/ or sudan/ or suriname/ or syrian arab republic/ or
82	exp "organisation for economic co-operation and development"/
83	exp australia/ or "australia and new zealand"/ or austria/ or baltic states/ or exp belgium/ or exp canada/ or chile/ or colombia/ or costa rica/ or czech republic/ or denmark/ or estonia/ or europe/ or exp finland/ or exp france/ or exp germany/ or greece/ or hungary/ or iceland/ or ireland/ or israel/ or exp italy/ or japan/ or korea/ or latvia/ or lithuania/ or luxembourg/ or exp mexico/ or netherlands/ or new zealand/ or north america/ or exp norway/ or poland/ or exp portugal/ or scandinavia/ or sweden/ or slovakia/ or slovenia/ or south korea/ or exp spain/ or switzerland/ or "Turkey (republic)"/ or exp united kingdom/ or exp united states/ or western europe/
84	european union/
85	developed country/
86	or/82-85
87	81 not 86
88	58 not 87
89	81 not 87

Database: The Cochrane Library: Cochrane Database of Systematic Reviews, Issue 12 of 12, December 2023 and Cochrane Central Register of Controlled Trials, Issue 12 of 12, December 2023

#	Searches
#1	MeSH descriptor: [Pregnancy] explode all trees
#2	MeSH descriptor: [Pregnant Women] this term only
#3	MeSH descriptor: [Prenatal Care] this term only
#4	(antenatal* or ante NEXT natal* or gestation* or maternal* or mother* or pregnan* or prenatal* or pre NEXT natal*):ti,ab,kw
#5	{OR #1-#4}

## Saarches MeSH descriptor: [Adiposity] this term only MeSH descriptor: [Body Size] this term only MeSH descriptor: [Body Size] this term only MeSH descriptor: [Overweight] this term only MeSH descriptor: [Overweight] this term only MeSH descriptor: [Obesity, Abdominal] this term only MeSH descriptor: [Obesity, Maternal] this term only MeSH descriptor: [Obesity, Metabolically Benign] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Waist-Height Ratio] MeSH descriptor: [Waist-Heigh		
MeSH descriptor: [Body Mass Index] this term only MeSH descriptor: [Body Size] this term only MeSH descriptor: [Overweight] this term only MeSH descriptor: [Overweight] this term only MeSH descriptor: [Obesity, Abdominal] this term only MeSH descriptor: [Obesity, Abdominal] this term only MeSH descriptor: [Obesity, Maternal] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Waist-Hip Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Maist-Height Ratio] this term only MeSH descriptor: [Maist-Height Ratio] this term only MeSH descriptor: [Adipose Tissue] explode all trees M	#	Searches
MeSH descriptor: [Body Weight] this term only MeSH descriptor: [Overweight] this term only MeSH descriptor: [Overweight] this term only MeSH descriptor: [Obesity, Abdominal] this term only MeSH descriptor: [Obesity, Maternal] this term only MeSH descriptor: [Obesity, Maternal] this term only MeSH descriptor: [Obesity, Maternal] this term only MeSH descriptor: [Obesity, Metabolically Benign] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Waist-Hip Ratio] this term only MeSH descriptor: [Waist-Hip Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Sody Fat Distribution] this term only MeSH descriptor: [Sody Fat Distribution] this term only MeSH descriptor: [Adipose Tissue] explode all trees (obes' or overweight or "over weight" or corpulent or heavy or heavier or fat or adipos" or (weight NEAR/2 (manag" or gain 'or increas' or excess or chang")).il.ab.kw Mest (waist NEXT ratio" triably was index" or BMI or "quetelet index").tl.ab.kw (okin NEXT fold" or skinfold* or "body composition" or ("body fat" NEAR/3 percent")):tl.ab.kw (okin NEXT fold" or skinfold* or "body composition" or ("body fat" NEAR/3 percent"):tl.ab.kw (okin NEXT fold" or skinfold* or "body composition" or ("body fat" NEAR/3 percent"):tl.ab.kw (okin NEXT fold" or skinfold* or "body composition" or ("body fat" NEAR/3 percent"):tl.ab.kw (okin NEXT fold" or skinfold* or "body composition" or ("body fat" NEAR/3 percent"):tl.ab.kw (okin NEXT fold" or skinfold* or "body composition" or "body fat" NEAR/3 percent"):tl.ab.kw (okin NEXT fold" or skinfold* or "body composition" or "body fat" NEAR/3 percent"):tl.ab.kw (okin NEXT fold" or skinfold* or "body composition" or "body fat" NEAR/3 percent"):tl.ab.kw (okin NEXT fold" or skinfold* or "body composition" or "body fat" NEAR/3 percent"):tl.ab.kw (okin NEXT fold" or skinfold* or calciol" or cholecalcifer	#6	MeSH descriptor: [Adiposity] this term only
#9 MeSH descriptor: [Doesity] this term only #1 MeSH descriptor: [Overweight] this term only #1 MeSH descriptor: [Overweight] this term only #1 MeSH descriptor: [Obesity, Abdominal] this term only #1 MeSH descriptor: [Obesity, Maternal] this term only #1 MeSH descriptor: [Obesity, Metabolically Benign] this term only #1 MeSH descriptor: [Obesity, Metabolically Benign] this term only #2 MeSH descriptor: [Obesity, Morbid] this term only #3 MeSH descriptor: [Waist-Hip Ratio] this term only #4 MeSH descriptor: [Waist-Hip Ratio] this term only #5 MeSH descriptor: [Waist-Hight Ratio] this term only #6 MeSH descriptor: [Waist-Hight Ratio] this term only #6 MeSH descriptor: [Waist-Hight Ratio] this term only #7 MeSH descriptor: [Waist-Hight Ratio] this term only #8 MeSH descriptor: [Waist-Hight Ra	#7	MeSH descriptor: [Body Mass Index] this term only
#1 MeSH descriptor: [Obesity, Abdominal] this term only MeSH descriptor: [Obesity, Abdominal] this term only MeSH descriptor: [Obesity, Maternal] this term only MeSH descriptor: [Obesity, Maternal] this term only MeSH descriptor: [Obesity, Metabolically Benign] this term only MeSH descriptor: [Obesity, Metabolically Benign] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Waist-Hip Ratio] this term only MeSH descriptor: [Waist-Hip Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Adjose Tissue] explode all trees MeSH descriptor: [Adjose Tissu	#8	MeSH descriptor: [Body Size] this term only
MeSH descriptor: [Obesity] this term only MeSH descriptor: [Obesity, Matemal] this term only MeSH descriptor: [Obesity, Matemal] this term only MeSH descriptor: [Obesity, Matemal] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Vaist Circumference] this term only MeSH descriptor: [Waist-Hip Ratio] this term only MeSH descriptor: [Waist-Hip Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Skinfold Thickness] this term only MeSH descriptor: [Body Fat Distribution] this term only MeSH descriptor: [Gody Fat Distribution] this term only MeSH descriptor: [Adipose Tissue] explode all trees (obes* or overweight or *over weight* or corpulen* or heavy or heavier or fat or adipos* or (weight NEAR/2 (manag* or gain* or increas* or excess or chang*)));ti,ab,kw (waist NEXT circumference* or waist NEXT hip NEXT ratio* or waist NEXT height NEXT ratio* or weight NEXT height NEXT ratio*);ti,ab,kw (amor midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*);ti,ab,kw (amor midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*);ti,ab,kw (amor midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*);ti,ab,kw (amor or midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*);ti,ab,kw (acicliferol* or calclifediol* or calclof* or cholecalcliferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or vitamin of or Xiamin of 25 NEXT hydroxy* or 25OH*);ti,ab,kw 3 #30 OR #31 #3 #29 AND #32 4 conference or or (clinicaltrials or trialsearch);so	#9	MeSH descriptor: [Body Weight] this term only
MeSH descriptor: [Obesity, Matemal] this term only MeSH descriptor: [Obesity, Matemal] this term only MeSH descriptor: [Obesity, Matemal] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Waist-Hip Ratio] this term only MeSH descriptor: [Waist-Hip Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Skinfold Thickness] this term only MeSH descriptor: [Skinfold Thickness] this term only MeSH descriptor: [Rody Fat Distribution] this term only MeSH descriptor: [Adipose Tissue] explode all trees MeSH descriptor: [Vitamin D] explode all trees		MeSH descriptor: [Overweight] this term only
MeSH descriptor: [Obesity, Maternal] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Obesity, Morbid] this term only MeSH descriptor: [Waist Circumference] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Weight Gain] this term only MeSH descriptor: [Skinfold Thickness] this term only MeSH descriptor: [Body Fat Distribution] this term only MeSH descriptor: [Adipose Tissue] explode all trees MeSH descriptor: [Adipose Tissue]		MeSH descriptor: [Obesity] this term only
## MeSH descriptor: [Obesity, Metabolically Benign] this term only ## MeSH descriptor: [Obesity, Morbid] this term only ## MeSH descriptor: [Waist Circumference] this term only ## MeSH descriptor: [Waist-Height Ratio] this term only ## MeSH descriptor: [Waist-Height Ratio] this term only ## MeSH descriptor: [Waist-Height Ratio] this term only ## MeSH descriptor: [Weight Gain] this term only ## MeSH descriptor: [Skinfold Thickness] this term only ## MeSH descriptor: [Skinfold Thickness] this term only ## MeSH descriptor: [Body Fat Distribution] this term only ## MeSH descriptor: [Adipose Tissue] explode all trees ## MeSH descriptor: [Vitamin D] explode all trees ## AND ## MeSH descriptor: [Vitamin D] explode all trees ## AND ## MeSH descriptor: [Vitamin D] explode all trees ## AND ## MeSH descriptor: [Vitamin D] explode all trees ## AND ## MeSH descriptor: [Vitamin D] explode all trees ## AND ## MeSH descriptor: [Vitamin D] explode all trees ## AND ## MeSH descriptor: [Vitamin D] explode all trees ## AND ##		MeSH descriptor: [Obesity, Abdominal] this term only
#1 MeSH descriptor: [Obesity, Morbid] this term only #2 MeSH descriptor: [Waist-Hip Ratio] this term only #3 MeSH descriptor: [Waist-Height Ratio] this term only #4 MeSH descriptor: [Weight Gain] this term only #4 MeSH descriptor: [Weight Gain] this term only #5 MeSH descriptor: [Skinfold Thickness] this term only #6 MeSH descriptor: [Skinfold Thickness] this term only #7 MeSH descriptor: [Skinfold Thickness] this term only #8 MeSH descriptor: [Adipose Tissue] explode all trees #8 MeSH descriptor: [Adipose Tissue] explode all trees #8 (obes* or overweight or "over weight" or corpulen* or heavy or heavier or fat or adipos* or (weight NEAR/2 (manag* or gain* or increas* or excess or chang*)):iti,ab,kw #8 (waist NEXT circumference* or waist NEXT hip NEXT ratio* or waist NEXT height NEXT ratio* or weight NEXT height NEXT ratio*):ti,ab,kw #8 (skin NEXT fold* or skinfold* or "body composition" or ("body fat" NEAR/3 percent")):ti,ab,kw #8 (QR #6-#27) #9 MeSH descriptor: [Vitamin D] explode all trees #9 MeSH descriptor: [Vitamin D] explode all trees #9 (acliderof* or calcifediol* or calciol* or cholecalciferof* or hydroxycholecalciferof* or dihydroxycholecalciferof* or calciferof* or calcifediol* or calciol* or cholecalciferof* or hydroxycholecalciferof* or dihydroxycholecalciferof* or dilation* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferof* or dihydroxycholecalciferof* or visamind or 25 NEXT hydroxy* or 25OH*):ti,ab,kw #9 #9 AND #32 #9 #9 AND #33 #9 Onference:pt or (clinicaltrials or trialsearch):so		MeSH descriptor: [Obesity, Maternal] this term only
MeSH descriptor: [Waist-Hip Ratio] this term only MeSH descriptor: [Waist-Hip Ratio] this term only MeSH descriptor: [Waist-Height Ratio] this term only MeSH descriptor: [Weight Gain] this term only MeSH descriptor: [Skinfold Thickness] this term only MeSH descriptor: [Skinfold Thickness] this term only MeSH descriptor: [Body Fat Distribution] this term only MeSH descriptor: [Adipose Tissue] explode all trees dobes* or overweight or "over weight" or corpulen* or heavy or heavier or fat or adipos* or (weight NEAR/2 (manag* or gain* or increas* or excess or chang*))):ti, ab, kw ("body mass index" or BMI or "quetelet index"):ti, ab, kw (waist NEXT circumference* or waist NEXT hip NEXT ratio* or waist NEXT height NEXT ratio* or weight NEXT height NEXT ratio*):ti, ab, kw (waist NEXT fold* or skinfold* or "body composition" or ("body fat" NEAR/3 percent")):ti, ab, kw ((arm or midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)):ti, ab, kw (OR #6-#27) #2 #3 MeSH descriptor: [Vitamin D] explode all trees (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferol* or dihydroxycholecalciferol* or vitamind or 25 NEXT dihydroxyv* or 25OH*):ti, ab, kw #3 #30 OR #31 33 NOT #34		MeSH descriptor: [Obesity, Metabolically Benign] this term only
## MeSH descriptor: [Waist-Hip Ratio] this term only ## MeSH descriptor: [Weight Ratio] this term only ## MeSH descriptor: [Weight Gain] this term only ## MeSH descriptor: [Skinfold Thickness] this term only ## MeSH descriptor: [Skinfold Thickness] this term only ## MeSH descriptor: [Body Fat Distribution] this term only ## MeSH descriptor: [Adipose Tissue] explode all trees ## (obes* or overweight or "over weight" or corpulen* or heavy or heavier or fat or adipos* or (weight NEAR/2 (manag* or gain* or increas* or excess or chang*)):ti,ab,kw ## ("body mass index" or BMI or "quetelet index"):ti,ab,kw ## (waist NEXT circumference* or waist NEXT hip NEXT ratio* or waist NEXT height NEXT ratio* or weight NEXT height NEXT ratio*) thi,ab,kw ## (skin NEXT fold* or skinfold* or "body composition" or ("body fat" NEAR/3 percent*)):ti,ab,kw ## ((arm or midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)):ti,ab,kw ## (OR #6-#27) ## 5 AND #28 ## 3 MeSH descriptor: [Vitamin D] explode all trees ## 3 MeSH descriptor: [Vitamin D] explode all trees ## 3 Moor ## 31 ## 3 #3 OOR ## 31 ## 3 #3 NOT #34		MeSH descriptor: [Obesity, Morbid] this term only
## MeSH descriptor: [Waist-Height Ratio] this term only ## MeSH descriptor: [Weight Gain] this term only ## MeSH descriptor: [Skinfold Thickness] this term only ## MeSH descriptor: [Skinfold Thickness] this term only ## MeSH descriptor: [Body Fat Distribution] this term only ## MeSH descriptor: [Adipose Tissue] explode all trees ## Clobes or overweight or "over weight" or corpulent or heavy or heavier or fat or adipos" or (weight NEAR/2 (manag" or gain" or increas" or excess or chang"))):ti,ab,kw ## Clobes or overweight or "over weight" or corpulent or heavy or heavier or fat or adipos" or (weight NEAR/2 (manag" or gain" or increas" or excess or chang"))):ti,ab,kw ## Clobes or overweight or "over weight" or corpulent or waist NEXT height NEXT ratio" or weight NEXT ratio" and the second or weight NEXT height NEXT ratio" or weight NEXT height NEXT ratio" or weight NEXT height NEXT ratio"):ti,ab,kw ## Clobes or overweight or "over weight" or corpulent or waist NEXT height NEXT ratio" or weight NEXT height NEXT ratio" or weight NEXT height NEXT ratio"):ti,ab,kw ## Clobes or overweight or "overweight" or overweight or weight NEXT height NEXT ratio" or weight NEXT height NEXT ratio" or weight NEXT height NEXT ratio" or weight NEXT height NEXT height NEXT ratio" or weight NEXT height NEXT height NEXT or weight NEXT height NEXT atio", bi,ab,kw ## Clobes or overweight or "overweight" or overweight or weight NEXT or weight NEXT or weight NEXT or weight NEXT atio" or area" or fat" or diameter" or anthropometr");ti,ab,kw ## Clobes or overweight or overweight or "overweight" or weight NEXT or diameter" or anthropometr");ti,ab,kw ## Clobes or overweight or excess or chang"));ti,ab,kw ## Clobes or overweight or excess or chang");ti,ab,kw ## Clobes or overweight or beavy or heavier or heavy or heavier or fat or adipos" or (eliminater or anthropometr");ti,ab,kw ## Clobes or overweight or		MeSH descriptor: [Waist Circumference] this term only
### MeSH descriptor: [Weight Gain] this term only ### MeSH descriptor: [Skinfold Thickness] this term only ### MeSH descriptor: [Body Fat Distribution] this term only ### MeSH descriptor: [Body Fat Distribution] this term only ### MeSH descriptor: [Adipose Tissue] explode all trees ### (obes* or overweight or "over weight" or corpulen* or heavy or heavier or fat or adipos* or (weight NEAR/2 (manag* or gain* or increas* or excess or chang*)));ti,ab,kw ### (waist NEXT circumference* or waist NEXT hip NEXT ratio* or waist NEXT height NEXT ratio* or weight NEXT height NEXT ratio*;ti,ab,kw ### (skin NEXT fold* or skinfold* or "body composition" or ("body fat" NEAR/3 percent*));ti,ab,kw ### (arm or midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*));ti,ab,kw ### (OR #6-#27) ### ### MeSH descriptor: [Vitamin D] explode all trees ### (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or vitamin dr or vitamin dr 25 NEXT dihydroxy* or 250H*);ti,ab,kw ### ### ### ### ### ### ### ### ### #		MeSH descriptor: [Waist-Hip Ratio] this term only
MeSH descriptor: [Skinfold Thickness] this term only MeSH descriptor: [Body Fat Distribution] this term only MeSH descriptor: [Adipose Tissue] explode all trees dobes or overweight or "over weight" or corpulent or heavy or heavier or fat or adipos or (weight NEAR/2 (manag* or gain* or increas* or excess or chang*))):ti,ab,kw ("body mass index" or BMI or "quetelet index"):ti,ab,kw (waist NEXT circumference* or waist NEXT hip NEXT ratio* or waist NEXT height NEXT ratio* or weight NEXT height NEXT ratio*);ti,ab,kw (skin NEXT fold* or skinfold* or "body composition" or ("body fat" NEAR/3 percent*)):ti,ab,kw ((arm or midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*));ti,ab,kw (OR #6-#27) MeSH descriptor: [Vitamin D] explode all trees (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or or calcifrol* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferol* or ergosterol* or viosterol or "vitamin d" or vitamind or 25 NEXT hydroxy* or 25OH*);ti,ab,kw 33 #30 OR #31 #33 NOT #34		MeSH descriptor: [Waist-Height Ratio] this term only
MeSH descriptor: [Body Fat Distribution] this term only #2 MeSH descriptor: [Adipose Tissue] explode all trees #3 (obes* or overweight or "over weight" or corpulen* or heavy or heavier or fat or adipos* or (weight NEAR/2 (manag* or again* or increas* or excess or chang*))):ti,ab,kw #4 ("body mass index" or BMI or "quetelet index"):ti,ab,kw #5 (waist NEXT circumference* or waist NEXT hip NEXT ratio* or waist NEXT height NEXT ratio* or weight NEXT height NEXT ratio* or weight NEXT height NEXT height NEXT fold* or skinfold* or "body composition" or ("body fat" NEAR/3 percent*)):ti,ab,kw #6 ((arm or midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)):ti,ab,kw #7 (OR #6-#27) #8 MeSH descriptor: [Vitamin D] explode all trees #8 MeSH descriptor: [Vitamin D] explode all trees #8 (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or vitamin or 25 NEXT hydroxy* or 25OH*):ti,ab,kw #8 #3 0R #31 #9 ON #31 #9 ON #33 #9 Conference:pt or (clinicaltrials or trialsearch):so		MeSH descriptor: [Weight Gain] this term only
MeSH descriptor: [Adipose Tissue] explode all trees (obes* or overweight or "over weight" or corpulen* or heavy or heavier or fat or adipos* or (weight NEAR/2 (manag* or gain* or increas* or excess or chang*))):ti,ab,kw ("body mass index" or BMI or "quetelet index"):ti,ab,kw (waist NEXT circumference* or waist NEXT hip NEXT ratio* or waist NEXT height NEXT ratio* or weight NEXT height NEXT ratio*):ti,ab,kw (skin NEXT fold* or skinfold* or "body composition" or ("body fat" NEAR/3 percent*)):ti,ab,kw ((arm or midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)):ti,ab,kw (OR #6-#27) #5 AND #28 #3 MeSH descriptor: [Vitamin D] explode all trees (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or vitamind or 25 NEXT hydroxy* or 25OH*):ti,ab,kw #3 #30 OR #31 #3 #29 AND #32 #3 conference:pt or (clinicaltrials or trialsearch):so #3 #3 NOT #34		MeSH descriptor: [Skinfold Thickness] this term only
2 (obes* or overweight or "over weight" or corpulen* or heavy or heavier or fat or adipos* or (weight NEAR/2 (manag* or 3 gain* or increas* or excess or chang*))):ti,ab,kw #2 ("body mass index" or BMI or "quetelet index"):ti,ab,kw #2 (waist NEXT circumference* or waist NEXT hip NEXT ratio* or waist NEXT height NEXT ratio* or weight NEXT height NEXT ratio*):ti,ab,kw #2 (skin NEXT fold* or skinfold* or "body composition" or ("body fat" NEAR/3 percent*)):ti,ab,kw #2 ((arm or midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)):ti,ab,kw #2 (PR #6-#27) #3 MeSH descriptor: [Vitamin D] explode all trees (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferol* or ergosterol* or viosterol or "vitamin d" or vitamind or 25 NEXT hydroxy* or 25OH*):ti,ab,kw #3 #3 0 OR #31 #3 #29 AND #32 *3 conference:pt or (clinicaltrials or trialsearch):so #3 #3 #33 NOT #34		MeSH descriptor: [Body Fat Distribution] this term only
gain* or increas* or excess or chang*))):ti,ab,kw ("body mass index" or BMI or "quetelet index"):ti,ab,kw (waist NEXT circumference* or waist NEXT hip NEXT ratio* or waist NEXT height NEXT ratio* or weight NEXT height NEXT height NEXT ratio* or weight NEXT height NEXT height NEXT part or weight NEXT height NEXT height NEXT or area* or fat* or diameter* or anthropometr*)):ti,ab,kw ((arm or midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)):ti,ab,kw (OR #6-#27) #5 AND #28 #6 (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferol* or ergosterol* or viosterol or "vitamin d" or vitamind or 25 NEXT hydroxy* or 25OH*):ti,ab,kw #3 #30 OR #31 #3 #30 OR #31 #3 conference:pt or (clinicaltrials or trialsearch):so #3 #3 #33 NOT #34		MeSH descriptor: [Adipose Tissue] explode all trees
(waist NEXT circumference* or waist NEXT hip NEXT ratio* or waist NEXT height NEXT ratio*):ti,ab,kw (skin NEXT fold* or skinfold* or "body composition" or ("body fat" NEAR/3 percent*)):ti,ab,kw ((arm or midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)):ti,ab,kw (OR #6-#27) #5 AND #28 (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calciol* or vitamin D or vitamin D or vitamin D or 25 NEXT hydroxy* or 25OH*):ti,ab,kw #3 #30 OR #31 #3 #29 AND #32 #3 conference:pt or (clinicaltrials or trialsearch):so #3 #33 NOT #34		
NEXT ratio*):ti,ab,kw (skin NEXT fold* or skinfold* or "body composition" or ("body fat" NEAR/3 percent*)):ti,ab,kw ((arm or midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)):ti,ab,kw (OR #6-#27) #5 AND #28 MeSH descriptor: [Vitamin D] explode all trees (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferol* or viosterol or "vitamin d" or vitamind or 25 NEXT hydroxy* or 250H*):ti,ab,kw #3 #30 OR #31 #3 #29 AND #32 #3 #33 NOT #34		
6 #2 ((arm or midarm or upperarm or brachial) NEAR/4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)):ti,ab,kw #2 {OR #6-#27} #3 #5 AND #28 MeSH descriptor: [Vitamin D] explode all trees (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferol* or ergosterol* or viosterol or "vitamind or 25 NEXT hydroxy* or 25OH*):ti,ab,kw #3 #30 OR #31 #3 #29 AND #32 #3 conference:pt or (clinicaltrials or trialsearch):so #3 #33 NOT #34		
anthropometr*)):ti,ab,kw #2 {OR #6-#27} #3 #5 AND #28 #4 MeSH descriptor: [Vitamin D] explode all trees (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferol* or vitamind or 25 NEXT hydroxy* or 25OH*):ti,ab,kw #3 #30 OR #31 #3 #29 AND #32 conference:pt or (clinicaltrials or trialsearch):so #3 #33 NOT #34		(skin NEXT fold* or skinfold* or "body composition" or ("body fat" NEAR/3 percent*)):ti,ab,kw
#2 #5 AND #28 #3 MeSH descriptor: [Vitamin D] explode all trees (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferol* or ergosterol* or viosterol or "vitamind or 25 NEXT hydroxy* or 25OH*):ti,ab,kw #3 #30 OR #31 #3 #29 AND #32 #3 conference:pt or (clinicaltrials or trialsearch):so #3 #33 NOT #34		
 MeSH descriptor: [Vitamin D] explode all trees (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferol* or ergosterol* or viosterol or "vitamin d" or vitamind or 25 NEXT hydroxy* or 25OH*):ti,ab,kw #3 #30 OR #31 #3 #29 AND #32 conference:pt or (clinicaltrials or trialsearch):so #3 #33 NOT #34 		{OR #6-#27}
#3 (calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferol* or ergosterol* or viosterol or "vitamin d" or vitamind or 25 NEXT hydroxy* or 25OH*):ti,ab,kw #3 #30 OR #31 #3 #29 AND #32 #3 conference:pt or (clinicaltrials or trialsearch):so #3 #33 NOT #34		#5 AND #28
1 calcitriol* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferol* or ergosterol* or viosterol or "vitamin d" or vitamind or 25 NEXT hydroxy* or 25OH*):ti,ab,kw #3 #30 OR #31 #3 #29 AND #32 #3 conference:pt or (clinicaltrials or trialsearch):so #3 #33 NOT #34		MeSH descriptor: [Vitamin D] explode all trees
2 #3 #29 AND #32 3 #3 conference:pt or (clinicaltrials or trialsearch):so #3 #33 NOT #34		calcitriol* or 24 NEXT 25 NEXT dihydroxyvitamin NEXT D or ergocalciferol* or ergosterol* or viosterol or "vitamin d" or
3 #3 conference:pt or (clinicaltrials or trialsearch):so #3 #33 NOT #34		#30 OR #31
4 #3 #33 NOT #34		#29 AND #32
		conference:pt or (clinicaltrials or trialsearch):so
		#33 NOT #34

Database: CINAHL

Date of last search: 05/12/2023

	of fast search: 05/12/2025
#	Searches
1	(MH "Pregnancy+")
2	(MH "Expectant Mothers")
3	(MH "Prenatal Care")
4	TI ((antenatal* or ante-natal* or gestational or maternal* or pregnan* or prenatal* or pre-natal*)) OR AB ((antenatal* or ante-natal* or gestational or maternal* or pregnan* or prenatal* or pre-natal*))
5	S1 OR S2 OR S3 OR S4
6	(MH "Body Mass Index")
7	(MH "Body Size")
8	(MH "Body Weight")
9	(MH "Obesity")
10	(MH "Obesity, Maternal") OR (MH "Obesity, Morbid")
11	(MH "Waist Circumference")
12	(MH "Waist-Hip Ratio")
13	(MH "Body Height")
14	(MH "Weight Gain")
15	(MH "Skinfold Thickness")
16	(MH "Adipose Tissue Distribution")
17	(MH "Adipose Tissue+")
18	TI ((obes* or overweight or "over weight" or corpulen* or heavy or heavier or fat or adipos*)) OR AB ((obes* or overweight or "over weight" or corpulen* or heavy or heavier or fat or adipos*))
19	TI ((weight N2 (manag* or gain* or increas* or excess or chang*))) OR AB ((weight N2 (manag* or gain* or increas* or excess or chang*)))
20	TI (("body mass index" or BMI or quetelet index)) OR AB (("body mass index" or BMI or quetelet index))
21	TI (("waist circumference*" or "waist hip ratio*" or "waist height ratio*" or "weight height ratio*")) OR AB (("waist circumference*" or "waist hip ratio*" or "waist height ratio*" or "weight height ratio*"))
22	TI (("skin fold*" or skinfold* or "body composition")) OR AB (("skin fold*" or skinfold* or "body composition"))
23	TI (((arm or midarm or upperarm or brachial) N4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*))) OR AB (((arm or midarm or upperarm or brachial) N4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)))
24	S6 OR S7 OR S8 OR S9 OR S10 OR S11 OR S12 OR S13 OR S14 OR S15 OR S16 OR S17 OR S18 OR S19 OR S20 OR S21 OR S22 OR S23
25	S5 AND S24
26	(MH "Vitamin D+")
27	TI ((calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or "24,25-dihydroxyvitamin D*" or ergocalciferol* or ergosterol* or viosterol or "vitamin d*" or vitamind* or 25 hydroxy* or 25-?OH*)) OR AB ((calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or "24,25-dihydroxyvitamin D*" or ergocalciferol* or ergosterol* or viosterol or "vitamin d*" or vitamind* or 25 hydroxy* or 25-?OH*))
28	S26 OR S27
29	S25 AND S28

Database: Epistemonikos

#	Searches
1	(advanced_title_en:((antenatal* OR ante-natal* OR gestational OR maternal* OR pregnan* OR prenatal* OR prenatal*)) OR advanced_abstract_en:((antenatal* OR ante-natal* OR gestational OR maternal* OR pregnan* OR prenatal* OR pre-natal*)))
2	((obese OR overweight OR adipos* OR BMI OR skinfold* OR "arm circumference")) OR advanced_abstract_en:((obese OR overweight OR adipos* OR BMI OR skinfold* OR "arm circumference")))
3	(advanced_title_en:("vitamin d") OR advanced_abstract_en:("vitamin d"))

DRAFT FOR CONSULTATION

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

Searches

4 1 AND 2 AND 3 [Filters: classification=systematic-review, cochrane=missing, protocol=no]

Economic searches

Databases: MEDLINE

#	Searches	
1	exp Pregnancy/ or Pregnant Women/ or Prenatal Care/	
2	(antenatal* or ante-natal* or gestational or maternal* or pregnan* or prenatal* or pre-natal*).ti,ab,kf.	
3	1 or 2	
1	adiposity/ or body mass index/ or body size/ or body weight/ or overweight/ or obesity/ or obesity, abdominal/ or obesity, maternal/ or obesity, metabolically benign/ or obesity, morbid/ or waist circumference/ or waist-hip ratio/ or Waist-He Ratio/ or Weight Gain/ or Skinfold Thickness/ or Body Fat Distribution/	
5	exp Adipose Tissue/	
3	(obes* or overweight or over weight or corpulen* or heavy or heavier or fat or adipos* or (weight adj2 (manag* or gain* or increas* or excess or chang*))).ti,ab,kf.	
7	(body mass index or BMI or quetelet index).ti,ab,kf.	
3	(waist circumference* or waist hip ratio* or waist height ratio* or weight height ratio*).ti,ab,kf.	
9	(skin fold* or skinfold* or body composition or (body fat adj3 percent*)).ti,ab,kf.	
10	((arm or midarm or upperarm or brachial) adj4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)).ti,ab,kf.	
11	or/4-10	
12	3 and 11	
13	exp Vitamin D/	
14	(calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24,25-dihydroxyvitamin D* or ergocalciferol* or ergosterol* or viosterol or vitamin d* or vitamind* or 25 hydroxy* or 25-?OH*).ti,ab,kf.	
15	or/13-14	
16	12 and 15	
17	letter/	
18	editorial/	
19	news/	
20	exp historical article/	
21	Anecdotes as Topic/	
22	comment/	
23	case reports/	
24	(letter or comment*).ti.	
25	or/17-24	
26	randomized controlled trial/ or random*.ti,ab.	
27	25 not 26	
28	animals/ not humans/	
29	exp Animals, Laboratory/	
30	exp Animal Experimentation/	
31	exp Models, Animal/	
32	exp Rodentia/	
33	(rat or rats or mouse or mice).ti.	
34	or/27-33	
35	16 not 34	
36	limit 35 to English language	
37	Economics/	
38	Value of life/	

#	Searches	
39	exp "Costs and Cost Analysis"/	
40	exp Economics, Hospital/	
41	· ·	
42	2 exp Resource Allocation/	
43	Economics, Nursing/	
44	Economics, Pharmaceutical/	
45	exp "Fees and Charges"/	
46	exp Budgets/	
47	budget*.ti,ab.	
48	cost*.ti,ab.	
49	(economic* or pharmaco?economic*).ti,ab.	
50	(price* or pricing*).ti,ab.	
51	(financ* or fee or fees or expenditure* or saving*).ti,ab.	
52	(value adj2 (money or monetary)).ti,ab.	
53	resourc* allocat*.ti,ab.	
54	(fund or funds or funding* or funded).ti,ab.	
55	(ration or rations or rationing* or rationed).ti,ab.	
56	ec.fs.	
57	or/37-56	
58	exp models, economic/	
59	*Models, Theoretical/	
60	*Models, Organizational/	
61	markov chains/	
62	monte carlo method/	
63	exp Decision Theory/	
64	(markov* or monte carlo).ti,ab.	
65	econom* model*.ti,ab.	
66	(decision* adj2 (tree* or analy* or model*)).ti,ab.	
67	or/58-66	
68	quality-adjusted life years/	
69	sickness impact profile/	
70	(quality adj2 (wellbeing or well being)).ti,ab.	
71	sickness impact profile.ti,ab.	
72	disability adjusted life.ti,ab.	
73	(qal* or qtime* or qwb* or daly*).ti,ab.	
74	(euroqol* or eq5d* or eq 5*).ti,ab.	
75	(qol* or hql* or hqol* or h qol* or hrqol* or hr qol*).ti,ab.	
76	(health utility* or utility score* or disutilit* or utility value*).ti,ab.	
77	(hui or hui1 or hui2 or hui3).ti,ab.	
78	(health* year* equivalent* or hye or hyes).ti,ab.	
79	discrete choice*.ti,ab.	
80	rosser.ti,ab.	
81	(willingness to pay or time tradeoff or time trade off or tto or standard gamble*).ti,ab.	
82	(sf36* or sf 36* or short form 36* or shortform 36* or shortform36*).ti,ab.	
83	(sf20 or sf 20 or short form 20 or shortform 20 or shortform20).ti,ab.	
84	(sf12* or sf 12* or short form 12* or shortform 12* or shortform12*).ti,ab.	
85	(sf8* or sf 8* or short form 8* or shortform 8* or shortform8*).ti,ab.	
86	(sf6* or sf 6* or short form 6* or shortform 6* or shortform6*).ti,ab.	
87	or/68-86	
88	36 and (57 or 67 or 87)	

Databases: Embase

1 2	Searches exp pregnancy/ or pregnant woman/ or prenatal care/ or prenatal period/	
	exp pregnancy/ or pregnant woman/ or prenatal care/ or prenatal period/	
2		
2 (antenatal* or ante natal* or gestation* or maternal* or mother* or pregnan* or prenatal* or pre natal*).ti,al		
3	1 or 2	
4	body mass/ or body size/ or body weight/ or body weight gain/ or obesity/ or abdominal obesity/ or fat mass/ or maternal obesity/ or metabolically benign obesity/ or morbid obesity/ or normal weight obesity/ or waist circumference/ or waist hip ratio/ or waist to height ratio/ or weight height ratio/ or skinfold thickness/ or body fat distribution/	
5	exp adipose tissue/ or exp obese patient/	
6	(obes* or overweight or over weight or corpulen* or heavy or heavier or fat or adipos* or (weight adj2 (manag* or gain* or increas* or excess or chang*))).ti,ab,kf.	
7	(body mass index or BMI or quetelet index).ti,ab,kf.	
8	(waist circumference* or waist hip ratio* or waist height ratio* or weight height ratio*).ti,ab,kf.	
9	(skin fold* or skinfold* or body composition or (body fat adj3 percent*)).ti,ab,kf.	
10	((arm or midarm or upperarm or brachial) adj4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)).ti,ab,kf.	
11	or/4-10	
12	3 and 11	
13	exp vitamin D/	
14	(calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24,25-dihydroxyvitamin D* or ergocalciferol* or ergosterol* or viosterol or vitamin d* or vitamind* or 25 hydroxy* or 25-?OH*).ti,ab,kf.	
15	or/13-14	
16	12 and 15	
17	letter.pt. or letter/	
18	note.pt.	
19	editorial.pt.	
20	case report/ or case study/	
21	(letter or comment*).ti.	
22	or/17-21	
23	randomized controlled trial/ or random*.ti,ab.	
24	22 not 23	
25	animal/ not human/	
26	nonhuman/	
27	exp Animal Experiment/	
28	exp Experimental Animal/	
29	animal model/	
30	exp Rodent/	
31	(rat or rats or mouse or mice or rodent*).ti.	
32	or/24-31	
33	16 not 32	
34	(conference abstract* or conference review or conference paper or conference proceeding).db,pt,su.	
35	33 not 34	
36	limit 35 to English language	
37	health economics/	
38	exp economic evaluation/	
39	exp health care cost/	
40	exp fee/	
41	budget/	
42	funding/	
43	resource allocation/	

#	Searches	
#		
	budget*.ti,ab.	
45 46	cost*.ti,ab. (economic* or pharmaco?economic*).ti,ab.	
47	(price* or pricing*).ti,ab.	
48		
	(financ* or fee or fees or expenditure* or saving*).ti,ab.	
49 50	(value adj2 (money or monetary)).ti,ab.	
51	resourc* allocat*.ti,ab.	
52	(fund or funds or funding* or funded).ti,ab. (ration or rations or rationing* or rationed).ti,ab.	
53	or/37-52	
54	statistical model/	
55	exp economic aspect/	
56	54 and 55	
57	*theoretical model/	
58	*nonbiological model/	
59	stochastic model/	
60	decision theory/	
61	decision tree/	
62	monte carlo method/	
63	(markov* or monte carlo).ti,ab.	
64	econom* model*.ti,ab.	
65	(decision* adj2 (tree* or analy* or model*)).ti,ab.	
66	or/56-65	
67	quality adjusted life year/	
68	"quality of life index"/	
69	short form 12/ or short form 20/ or short form 36/ or short form 8/	
70	sickness impact profile/	
71	(quality adj2 (wellbeing or well being)).ti,ab.	
72	sickness impact profile.ti,ab.	
73	disability adjusted life.ti,ab.	
74	(qal* or qtime* or qwb* or daly*).ti,ab.	
75	(qal* or qtime* or qwb* or daly*).ti,ab.	
76	(qol* or hql* or hqol* or h qol* or hrqol* or hr qol*).ti,ab.	
77	(health utility* or utility score* or disutilit* or utility value*).ti,ab.	
78	(hui or hui1 or hui2) or hui3).ti,ab.	
79	(health* year* equivalent* or hye or hyes).ti,ab.	
80	discrete choice*.ti,ab.	
81	rosser.ti,ab.	
82	(willingness to pay or time tradeoff or time trade off or tto or standard gamble*).ti,ab.	
83	(sf36* or sf 36* or short form 36* or shortform 36* or shortform36*).ti,ab.	
84	(sf20 or sf 20 or short form 20 or shortform 20 or shortform20).ti,ab.	
85 86	(sf12* or sf 12* or short form 12* or shortform 12* or shortform12*).ti,ab.	
87	(sf8* or sf 8* or short form 8* or shortform 8* or shortform8*).ti,ab. (sf6* or sf 6* or short form 6* or shortform 6* or shortform6*).ti,ab.	
88	or/67-87	
89	36 and (53 or 66 or 88)	
09	55 dire (55 5) 50 00 00 00)	

Database: INAHTA HTA

	of last search: U5/12/2023	
#	Searches	
1	"Pregnancy"[mhe]	
	2 "Pregnant Women"[mh]	
3	"Prenatal Care"[mh]	
4	((antenatal* or ante-natal* or gestational or maternal* or pregnan* or prenatal* or pre-natal*))[Title] OR ((antenatal* or ante-natal* or gestational or maternal* or pregnan* or prenatal* or pre-natal*))[abs]	
5	#1 OR #2 OR #3 OR #4	
6	"Adiposity"[mh]	
7	"Body Mass Index"[mh]	
8	"Body Size"[mh]	
9	"Body Weight"[mh]	
10	"Overweight"[mh]	
11	"Obesity"[mh]	
12	"Obesity, Abdominal"[mh]	
13	"Obesity, Maternal"[mh]	
14	"Obesity, Metabolically Benign"[mh]	
15	"Obesity, Morbid"[mh]	
16	"Waist Circumference"[mh]	
17	"Waist-Hip Ratio"[mh]	
18	"Waist-Height Ratio"[mh]	
19	"Weight Gain"[mh]	
20 "Skinfold Thickness"[mh]		
21	"Body Fat Distribution"[mh]	
22	"Adipose Tissue"[mhe]	
23	((obes* or overweight or "over weight" or corpulen* or heavy or heavier or fat or adipos*))[Title] OR ((obes* or overweight or "over weight" or corpulen* or heavy or heavier or fat or adipos*))[abs]	
24	((weight AND (manag* or gain* or increas* or excess or chang*)))[Title] OR ((weight AND (manag* or gain* or increas* or excess or chang*)))[abs]	
25	(("body mass index" or BMI or "quetelet index"))[Title] OR (("body mass index" or BMI or "quetelet index"))[abs]	
26	(("waist circumference*" or "waist hip ratio*" or "waist height ratio*" or "weight height ratio*"))[Title] OR (("waist circumference*" or "waist hip ratio*" or "waist height ratio*" or "weight height ratio*"))[abs]	
27	(("skin fold*" or skinfold* or "body composition"))[Title] OR (("skin fold*" or skinfold* or "body composition"))[abs]	
28	(("body fat" AND percent*))[Title] OR (("body fat" AND percent*))[abs]	
29	(((arm or midarm or upperarm or brachial) AND (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)))[Title] OR (((arm or midarm or upperarm or brachial) AND (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)))[abs]	
30	#6 OR #7 OR #8 OR #9 OR #10 OR #11 OR #12 OR #13 OR #14 OR #15 OR #16 OR #17 OR #18 OR #19 OR #20 OR #21 OR #22 OR #23 OR #24 OR #25 OR #26 OR #27 OR #28 OR #29	
31	"Vitamin D"[mhe]	
32	((calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24,25-dihydroxyvitamin D* or ergocalciferol* or ergosterol* or viosterol or vitamin d* or vitamind* or 25 hydroxy*))[Title] OR ((calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24,25-dihydroxyvitamin D* or ergocalciferol* or ergosterol* or viosterol or vitamin d* or vitamind* or 25 hydroxy*))[abs]	
33	#31 OR #32	
34	#5 AND #30	
35	#33 AND #34	

Database: CRD HTA (last updated October 2016)

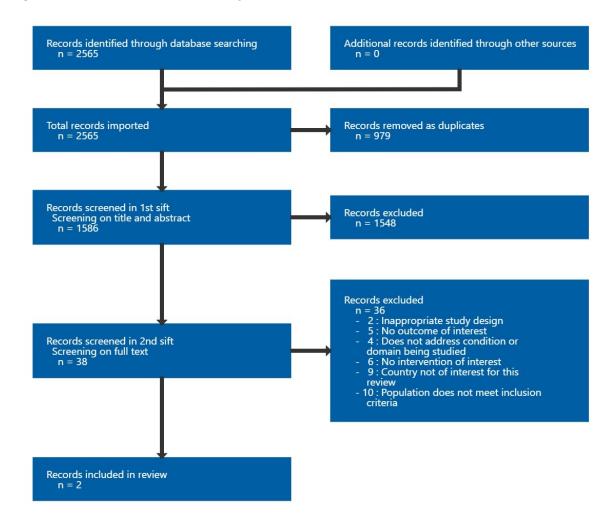
Date of last search: 14/07/2022

	of last search: 14/07/2022	
#	Searches	
1	MeSH DESCRIPTOR pregnancy EXPLODE ALL TREES IN HTA	
2	MeSH DESCRIPTOR pregnant women IN HTA	
3	MeSH DESCRIPTOR prenatal care IN HTA	
4	(((antenatal* or ante-natal* or gestational or maternal* or pregnan* or prenatal* or pre-natal*))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
5	#1 OR #2 OR #3 OR #4	
6	MeSH DESCRIPTOR adiposity IN HTA	
7	MeSH DESCRIPTOR body mass index IN HTA	
8	MeSH DESCRIPTOR body size IN HTA	
9	MeSH DESCRIPTOR body weight IN HTA	
10	MeSH DESCRIPTOR overweight IN HTA	
11	MeSH DESCRIPTOR obesity IN HTA	
12	MeSH DESCRIPTOR obesity, abdominal IN HTA	
13	MeSH DESCRIPTOR obesity, maternal IN HTA	
14	MeSH DESCRIPTOR obesity, metabolically benign IN HTA	
15	MeSH DESCRIPTOR obesity, morbid IN HTA	
16	MeSH DESCRIPTOR waist circumference IN HTA	
17	MeSH DESCRIPTOR waist-hip ratio IN HTA	
18	MeSH DESCRIPTOR waist-height ratio IN HTA	
19	MeSH DESCRIPTOR weight gain IN HTA	
20	MeSH DESCRIPTOR skinfold thickness IN HTA	
21	MeSH DESCRIPTOR body fat distribution IN HTA	
22	MeSH DESCRIPTOR adipose tissue EXPLODE ALL TREES IN HTA	
23	(((obes* or overweight or over weight or corpulen* or heavy or heavier or fat or adipos*))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
24	(((weight adj2 (manag* or gain* or increas* or excess or chang*)))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
25	(((body mass index or BMI or quetelet index))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
26	(((waist circumference* or waist hip ratio* or waist height ratio* or weight height ratio*))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
27	(((skin fold* or skinfold* or body composition))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
28	(((body fat adj3 percent*))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
29	((((arm or midarm or upperarm or brachial) adj4 (circumferen* or measure* or area* or fat* or diameter* or anthropometr*)))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
30	#6 OR #7 OR #8 OR #9 OR #10 OR #11 OR #12 OR #13 OR #14 OR #15 OR #16 OR #17 OR #18 OR #19 OR #20 OR #21 OR #22 OR #23 OR #24 OR #25 OR #26 OR #27 OR #28 OR #29	
31	#5 AND #30	
32	MeSH DESCRIPTOR VITAMIN D EXPLODE ALL TREES IN HTA	
33	(((calciferol* or calcifediol* or calciol* or cholecalciferol* or hydroxycholecalciferol* or dihydroxycholecalciferol* or calcitriol* or 24,25-dihydroxyvitamin D* or ergocalciferol* or ergosterol* or viosterol or vitamin d* or vitamind*))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
34	#32 OR #33	
35	#31 AND #34	

Appendix C Effectiveness evidence study selection

Study selection for: What dose of vitamin D is appropriate during pregnancy for women with a BMI medically classified as overweight or obese?

Figure 1: Effective evidence study selection flow chart



Appendix D Evidence tables

Evidence tables for review question: What dose of vitamin D is appropriate during pregnancy for women with a BMI medically classified as overweight or obese?

Table 4: Evidence tables

Alhomaid, 2021

Bibliographic Reference

Alhomaid, Raghad M; Mulhern, Maria S; Strain, Jj; Laird, Eamon; Healy, Martin; Parker, Michael J; McCann, Mary T; Maternal obesity and baseline vitamin D insufficiency alter the response to vitamin D supplementation: a double-blind, randomized trial in pregnant women.; The American journal of clinical nutrition; 2021; vol. 114 (no. 3); 1208-1218

Study details

Country/ies where study was carried out	Northern Ireland
Study type	Randomised controlled trial (RCT)
Study dates	Not reported
Inclusion criteria	 pregnant women ≥12 weeks gestational age aged ≥18 years BMI ≥18.5kg/m² without current pregnancy-related complications having a singleton pregnancy
Exclusion criteria	 multiple pregnancy involvement in another research study

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

	 history of gastrointestinal, hepatic, renal, vascular, or hematological disorders in vitro fertilization treatment a history of neural tube defect pregnancies active thyroid disease
Patient characteristics	Age, (mean \pm SD), years Intervention = 29.5 \pm 5.5 Control = 29.7 \pm 5.1
	BMI, (mean ±SD), kg/m ²
	Intervention = 27.8 ±5.4
	Control = 28.1 ±5.7
	BMI categories
	Normal weight, %
	Intervention = 33.9
	Control = 33.1
	Overweight, %
	Intervention = 33.1
	Control = 33.1
	Obese, %
	Intervention = 33.1

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

	Control = 33.9
	Gestation week (mean ±SD) Intervention = 12.8 ±1.4 Control = 13.0 ±1.4
Intervention(s)/control	Intervention: 1 multivitamin tablet containing 10-μg vitamin D and a 10-μg vitamin D tablet Control: 1 multivitamin tablet containing 10-μg vitamin D and a 0-μg vitamin D (placebo) tablet
Duration of follow-up	No follow-up (immediately post-intervention)
Sources of funding	Not industry funded
Sample size	N = 240 Intervention group, n =122 (n=1 participant withdrew from 10mcg group after 12-week baseline) Control group, n =118 Note:
Other information	Recruitment took place at the first antenatal visit. Women were enrolled in the study after 12 weeks gestational age, and anthropometric measurements (height and weight) were taken at this point. Women taking any vitamin D containing supplement before 12 weeks gestational age were asked to only take the study supplements after enrolment.

BMI: Body mass index; RCT: randomised controlled trial; SD: standard deviation

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

Study arms

Intervention arm: $20\mu g/d$ Vitamin D (N = 121) Comparison arm: $10\mu g/d$ Vitamin D (N = 181)

Outcomes

Study timepoints

• 28 weeks (gestational age)

• 36 weeks (gestational age)

Maternal serum 25(OH)D, nmol/L concentration at various timepoints

Outcome	28 week, Intervention arm: 20ug/d Vitamin D, N = 121	28 week, Comparison arm: 10ug/d Vitamin D, N = 118	36 week, Intervention arm: 20ug/d Vitamin D, N = 121	36 week, Comparison arm: 10ug/d Vitamin D, N = 118
Serum 25(OH)D nmol/L in overweight (25.0 to 29.9 kg/m²) women Mean (SD)	81.6 (29.6)	72.9 (25.8)	93.9 (31.3)	81.9 (24.9)
Serum 25(OH)D nmol/L in obese (BMI ≥ 30 kg/m²) women Mean (SD)	74.6 (32.2)	64.3 (27)	82.1 (33.4)	74.8 (30.6)

25 (OH)D: 25 hydroxyvitamin D; BMI: Body mass index; nmol/L: nanomoles per litre; SD: standard deviation

Critical appraisal

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

Section	Question	Answer
Domain 1: Bias arising from the randomisation process	Risk of bias judgement for the randomisation process	Low (No serious concerns about the randomisation process)
Domain 2a: Risk of bias due to deviations from the intended interventions (effect of assignment to intervention)	Risk of bias for deviations from the intended interventions (effect of assignment to intervention)	Low (Intention-to-treat analysis was used to account for missing data)
Domain 2b: Risk of bias due to deviations from the intended interventions (effect of adhering to intervention)	Risk of bias judgement for deviations from the intended interventions (effect of adhering to intervention)	Low (No serious concerns on the effect of adhering to interventions)
Domain 3. Bias due to missing outcome data	Risk-of-bias judgement for missing outcome data	Low (No serious concerns about missing outcome data, Intention-to-treat analysis was carried out)
Domain 4. Bias in measurement of the outcome	Risk-of-bias judgement for measurement of the outcome	Low (No serious concerns about outcome measurement)
Domain 5. Bias in selection of the reported result	Risk-of-bias judgement for selection of the reported result	Low (No serious concerns on the reported results)
Overall bias and Directness	Risk of bias judgement	Low (No serious concerns in any domain)
Overall bias and Directness	Overall Directness	Directly applicable
Overall bias and Directness	Risk of bias variation across outcomes	N/A

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

Corcoy, 2020

Bibliographic Reference

Corcoy, Rosa; Mendoza, Lilian C; Simmons, David; Desoye, Gernot; Adelantado, J M; Chico, Ana; Devlieger, Roland; van Assche, Andre; Galjaard, Sander; Timmerman, Dirk; Lapolla, Annunziata; Dalfra, Maria G; Bertolotto, Alessandra; Harreiter, Jurgen; Wender-Ozegowska, Ewa; Zawiejska, Agnieszka; Kautzky-Willer, Alexandra; Dunne, Fidelma P; Damm, Peter; Mathiesen, Elisabeth R; Jensen, Dorte M; Andersen, Lise Lotte T; Tanvig, Mette; Hill, David J; Jelsma, Judith G; Snoek, Frank J; Kofeler, Harald; Trotzmuller, Martin; Lips, Paul; van Poppel, Mireille N M; The DALI vitamin D randomized controlled trial for gestational diabetes mellitus prevention: No major benefit shown besides vitamin D sufficiency.; Clinical nutrition (Edinburgh, Scotland); 2020; vol. 39 (no. 3); 976-984

Study details

Country/ies where	7 European countries:		
study was carried out	United Kingdom		
	Ireland		
	Austria		
	• Poland		
	Italy (Padua, Pisa)		
	Spain		
	Belgium		
Study type	Cluster randomised controlled trial		
Study dates	2012 to 2015		
Inclusion criteria	 pregnant women with a pre-pregnancy BMI ≥29 kg/m² 		
	≤19 weeks + 6 days of gestation		

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

	singleton pregnancy
	aged ≥18 years
Exclusion criteria	GDM diagnosed by oral glucose tolerance test (OGTT) using IADPSG/WHO 2013 criteria
	pre-existing diabetes, chronic medical conditions or a psychiatric disorder
	being unable to walk ≥ 100 m safely
	requiring complex diets
	not fluent in the major language of the country
	 having current or past abnormal calcium metabolism (hypo/hyperparathyroidism, nephrolithiasis, hypercalciuria), or having hypercalciuria (>0.6 mmol/mmol creatinine in morning spot samples)
	hypercalcemia detected at baseline measurement according to trimester cut-offs
Patient	Age (mean±SD), years
characteristics	Intervention = 32.2±5.2
	Control = 32.8 ± 5.4
	Parity, %
	Multiparous
	Intervention = 53
	Control = 57
	History of GDM, %
	Intervention = 15

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

Control = 15

Gestational age at entry (mean±SD), weeks

Intervention = 15.0 ± 2.9

Control = 15.4 ± 2.5

Pre-pregnancy BMI, (mean±SD), kg/m²

Intervention = 33.7 ± 4.3

Control = 33.3 ± 4.3

BMI at entry, (mean±SD), kg/m²

Intervention = 34.2 ± 4.3

Control = 34.2 ± 4.2

Taking (multi)vitamins at entry, %

Intervention = 86

Control = 84

Intervention(s)/control Intervention: Vitamin D3 supplementation (1600 IU/day) (with and without lifestyle counselling)

Control: placebo (with and without lifestyle counselling)

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

Sample size $N = 154$ Intervention group, $n = 79$ Placebo group, $n=75$ Design effect = $1+ (7-1) \times 0.231 = 2.386$ Adjusted sample size = $154/2.386 = 64.5 \sim 65$	-	
Sample size N = 154 Intervention group, n = 79 Placebo group, n=75 Design effect = 1+ (7-1) x 0.231 = 2.386 Adjusted sample size = 154/2.386 = 64.5 ~ 65 Other information Study randomised participants into 4 groups: Lifestyle counselling (healthy eating and physical activity) and placebo, Lifestyle counselling and vitamin D supplementation, vitamin D supplementation alone and placebo alone The study reported that there was no interaction between the lifestyle intervention and the vitamin D supplementation and compared them with the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment		the chosen intervention dose was 1600 IU/day. The intervention was delivered in a tablet form. Each tablet contained 400 IU of vitamin D3. Placebo tablets, identical to the intervention tablets in appearance, were produced especially for
N = 154 Intervention group, n = 79 Placebo group, n=75 Design effect = 1+ (7-1) x 0.231 = 2.386 Adjusted sample size = 154/2.386 = 64.5 ~ 65 Other information Study randomised participants into 4 groups: Lifestyle counselling (healthy eating and physical activity) and placebo, Lifestyle counselling and vitamin D supplementation, vitamin D supplementation alone and placebo alone The study reported that there was no interaction between the lifestyle intervention and the vitamin D intervention for any of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment	Duration of follow-up	Immediately post-intervention
Intervention group, n = 79 Placebo group, n=75 Design effect = 1+ (7-1) x 0.231 = 2.386 Adjusted sample size = 154/2.386 = 64.5 ~ 65 Other information Study randomised participants into 4 groups: Lifestyle counselling (healthy eating and physical activity) and placebo, Lifestyle counselling and vitamin D supplementation, vitamin D supplementation alone and placebo alone The study reported that there was no interaction between the lifestyle intervention and the vitamin D intervention for any of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment	Sources of funding	Not industry funded
Placebo group, n=75 Design effect = 1+ (7-1) x 0.231 = 2.386 Adjusted sample size = 154/2.386 = 64.5 ~ 65 Other information Study randomised participants into 4 groups: Lifestyle counselling (healthy eating and physical activity) and placebo, Lifestyle counselling and vitamin D supplementation, vitamin D supplementation alone and placebo alone The study reported that there was no interaction between the lifestyle intervention and the vitamin D intervention for any of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment	Sample size	N = 154
Design effect = 1+ (7-1) x 0.231 = 2.386 Adjusted sample size = 154/2.386 = 64.5 ~ 65 Other information Study randomised participants into 4 groups: Lifestyle counselling (healthy eating and physical activity) and placebo, Lifestyle counselling and vitamin D supplementation, vitamin D supplementation alone and placebo alone The study reported that there was no interaction between the lifestyle intervention and the vitamin D intervention for any of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment		Intervention group, n = 79
Adjusted sample size = 154/2.386 = 64.5 ~ 65 Other information Study randomised participants into 4 groups: Lifestyle counselling (healthy eating and physical activity) and placebo, Lifestyle counselling and vitamin D supplementation, vitamin D supplementation alone and placebo alone The study reported that there was no interaction between the lifestyle intervention and the vitamin D intervention for any of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment		Placebo group, n=75
Adjusted sample size = 154/2.386 = 64.5 ~ 65 Other information Study randomised participants into 4 groups: Lifestyle counselling (healthy eating and physical activity) and placebo, Lifestyle counselling and vitamin D supplementation, vitamin D supplementation alone and placebo alone The study reported that there was no interaction between the lifestyle intervention and the vitamin D intervention for any of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment		
Study randomised participants into 4 groups: • Lifestyle counselling (healthy eating and physical activity) and placebo, • Lifestyle counselling and vitamin D supplementation, • vitamin D supplementation alone and • placebo alone The study reported that there was no interaction between the lifestyle intervention and the vitamin D intervention for any of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment		Design effect = 1+ (7-1) x 0.231 = 2.386
 Lifestyle counselling (healthy eating and physical activity) and placebo, Lifestyle counselling and vitamin D supplementation, vitamin D supplementation alone and placebo alone The study reported that there was no interaction between the lifestyle intervention and the vitamin D intervention for any of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment 		Adjusted sample size = 154/2.386 = 64.5 ~ 65
 Lifestyle counselling and vitamin D supplementation, vitamin D supplementation alone and placebo alone The study reported that there was no interaction between the lifestyle intervention and the vitamin D intervention for any of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment 	Other information	Study randomised participants into 4 groups:
 vitamin D supplementation alone and placebo alone The study reported that there was no interaction between the lifestyle intervention and the vitamin D intervention for any of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment 		Lifestyle counselling (healthy eating and physical activity) and placebo,
 placebo alone The study reported that there was no interaction between the lifestyle intervention and the vitamin D intervention for any of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment 		
of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and compared them with the 2 groups randomised to have placebo. Pre-pregnancy BMI (used as an inclusion criterion) was self-reported at recruitment		• •
		of the outcomes, and therefore the authors combined the 2 groups randomised to vitamin D supplementation and

BMI: Body mass index; IADPSG: the international association of the diabetes and pregnancy study groups; IU: international units; OGTT: oral glucose tolerance test; (OH)D: hydroxyvitamin D: SD: standard deviation; WHO: world health organisation

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

Study arms

Intervention arm: 1600IU/day (N = 79)

Control arm: Placebo (N = 75)

Outcomes

Study timepoints

- 24 week gestational age (Study reports 24-28weeks)
- 35 week gestational age (Study reports 35-37 weeks)
- 38 week gestational age (Study reports at delivery)

Total serum 25(OH)D concentrations

Outcome	Intervention arm: 1600IU/day, 24 week, N = 32	Intervention arm: 1600IU/day, 35 week, N = 29	Intervention arm: 1600IU/day, 38 week, N = 31	Control arm: Placebo, 24 week, N = 28	Control arm: Placebo, 35 week, N = 25	Control arm: Placebo, 38 week, N = 27
25(OH)D nmol/L level in women with BMI ≥ 29kg/m ²	119.4 (35.5)	122.9 (38.8)	101.2 (32.2)	81.9 (39.4)	84.5 (39.8)	76.9 (34.8)
Mean (SD)						

BMI: body mass index; nmol/L: nanomoles per litre; (OH)D: hydroxyvitamin D; IU: international units Sample sizes and estimates are adjusted for clustering effect

Preterm birth

Outcome	Intervention arm: 1600IU/day, 38 week, N = 31	Control arm: Placebo, 38 week, N = 27
Preterm birth	n = 1; % = 3	n = 0; % = 2
No of events		

IU: international units

Sample sizes and estimates are adjusted for clustering effect

Birth weight <2.5kg

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

Outcome	Intervention arm: 1600IU/day, 38 week, N = 31	Control arm: Placebo, 38 week, N = 26
Birth weight <2.5kg	n = 0; % = 0	n = 0; % = 2
No of events		

IU: international units

Sample sizes and estimates are adjusted for clustering effect

Preeclampsia

Outcome	Intervention arm: 1600IU/day, 38 week, N = 28	Control arm: Placebo, 38 week, N = 24
Preeclampsia	n = 0; % = 2	n = 1; % = 5
No of events		

IU: international units

Sample sizes and estimates are adjusted for clustering effect

Gestational diabetes mellitus

Outcome	Intervention arm: 1600IU/day, 24 week, N = 31	Intervention arm: 1600IU/day, 35 week, N = 29	Control arm: Placebo, 24 week, N = 28	Control arm: Placebo, 35 week, N = 22
Gestational Diabetes mellitus	n = 6; % = 20	n = 11; % = 37	n = 7; % = 26	n = 10; % = 39
No of events				

IU: international units

Sample sizes and estimates are adjusted for clustering effect

Critical appraisal - Cochrane Risk of Bias tool (RoB 2.0) Cluster randomised trials NGA

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

Section	Question	Answer
1a. Bias arising from the randomisation process	Risk of bias judgement for the randomisation process	Some concerns (No information on the likelihood that allocation sequence was subverted)
1b. Bias arising from the timing of identification and recruitment of individual participants in relation to timing of randomisation	Risk of bias judgement for the timing of identification and recruitment of individual participants in relation to timing of randomisation	Low
2. Bias due to deviations from intended interventions (If your aim is to assess the effect of assignment to intervention, answer the following questions).	Risk of bias judgement for deviations from intended interventions	Low
3. Bias due to missing outcome data	Risk of bias judgement for missing outcome data	Some concerns (16% loss to follow up but ITT analysis conducted)
4. Bias in measurement of the outcome	Risk of bias judgement for measurement of the outcome	Low
5. Bias in selection of the reported result	Risk of bias for selection of the reported result	Low
Overall bias and Directness	Risk of bias judgement	Some concerns (Some concerns about the randomisation process and missing outcome data)
Overall bias and Directness	Overall Directness	Directly applicable

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

Appendix E Forest plots

Forest plots for review question: What dose of vitamin D is appropriate during pregnancy for women with a BMI medically classified as overweight or obese?

No meta-analysis was conducted for this review question and so there are no forest plots.

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

Appendix F GRADE tables

GRADE tables for review question: What dose of vitamin D is appropriate during pregnancy for women with a BMI medically classified as overweight or obese?

Table 5: Evidence profile for comparison between 20µg/day and 10µg/day vitamin D supplementation in pregnant women medically

classified as overweight (25.0 to 29.9 kg/m²) or obese (BMI ≥30.0 kg/m²)

	Ciassilled	as overv	vergiit (25.0 ti	0 29.9 kg/III) or ones	e (BMI 230.0 I	kg/III)		ı			
			Quality assess	ment			No of p	oatients	Effect		Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	20µg/day vitamin D	10µg/day vitamin D	Relative (95% CI)	Absolute		
			n (nmol/L) in preg ilable kit; Better ir			fied as overweigh	t (BMI 25.0 to	29.9k/m²) at 2	28 weeks	gestational age (mea	sured with: LC-ta	andem MS
1 (Alhomaid 2021)	randomised trials		no serious inconsistency	no serious indirectness	serious ¹	none	40	39	-	MD 8.7 higher (3.54 lower to 20.94 higher)	MODERATE	CRITICAL
											NO IMP. DIFF.	
	Maternal serum 25(OH)D concentration (nmol/L) in pregnant women medically classified as obese (BMI ≥30.0kg/m²) at 28 weeks gestational age (measured with: LC-tandem MS (LC-MS/MS) using a commercially available kit; Better indicated by higher values)											
1 (Alhomaid 2021)	randomised trials		no serious inconsistency	no serious indirectness	serious ¹	none	40	40	-	MD 10.3 higher (2.72 lower to 23.32 higher)	MODERATE	CRITICAL
											NO. EV. OF. IMP. DIFF.	
			n (nmol/L) in wom e kit; Better indica			erweight (BMI 25.0	0 to 29.9kg/m	²) at 36 weeks	gestatio	nal age (measured w	ith: LC-tandem N	IS (LC-
1 (Alhomaid 2021)	randomised trials		no serious inconsistency	no serious indirectness	serious ¹	none	40	39	-	MD 12 higher (0.46 lower to 24.46 higher)	MODERATE	CRITICAL
											POSS. IMP. BENEFIT.	
			n (nmol/L) in preg e kit; Better indica			fied as obese (BM	l≥30.0kg/m²) :	at 36 weeks g	estationa	al age (measured with	: LC-tandem MS	(LC-
1 (Alhomaid 2021)	randomised trials		no serious inconsistency	no serious indirectness	serious ¹	none	40	40	-	MD 7.3 higher (6.74 lower to 21.34 higher)	MODERATE	CRITICAL
			tenual: MD: mean di								NO. IMP. DIFF.	

BMI: Body mass index; CI: confidence interval; MD: mean difference; nmol/L: nanomoles per litre; 25(OH)D: 25 hydroxyvitamin D

¹ 95% CI crosses 1 MID (0.5 x control group SD for '25 (OH)D concentration in overweight pregnant women' = ±11.1); (0.5 x control group SD for '25 (OH)D concentration in obese pregnant women' = ±9.85)

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

Table 6: Evidence profile for comparison between 40μg /day vitamin D supplementation and no supplementation in pregnant women medically classified as overweight or obese (BMI > 29 0kg/m²)

	medica	lly clas	sified as ove	rweight or	obese (BMI	≥ 29.0kg/m²)) 		l			
			Quality as	sessment			No of patient				Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	1600IU/day vitamin D versus Placebo	Control	Relative (95% CI)	Absolute		
	erum 25(OH) indicated by			nol/L) at 24 to 28	8 weeks (measi	ured with: ClinMas	ss liquid chromatog	raphy-m	ass spectron	netry/mass spectrome	etry (LC-MS/MS) complete
	randomised trials		no serious inconsistency	no serious indirectness	no serious imprecision	none	32	28	-	MD 37.5 higher (18.41 to 56.59 higher)	MODERATE IMP. BENEFIT	CRITICAL
	erum 25(OH) indicated by			nol/L) at 35 to 3	7 weeks (measi	ured with: ClinMas	ss liquid chromatog	raphy-m	ass spectron	netry/mass spectrome	etry (LC-MS/MS) complete
1 (Corcoy		serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	29	25	-	MD 38.4 higher (17.36 to 59.44 higher)	MODERATE	CRITICAL
			concentration (nr			eeks not specified	d) (measured with: C	linMass	liquid chrom	atography-mass spec	ctrometry/mass	
1 (Corcoy		serious ¹	no serious	no serious indirectness	serious ²	none	31	27	-	MD 24.3 higher (6.96 to 41.64 higher)	LOW	CRITICAL
											IMP. BENEFIT	
Preterm bi	irth (not defir		dy) at birth									
	randomised trials		no serious inconsistency	no serious indirectness	very serious ³	none	1/31 (3.2%)	0/27 (0%)	POR 6.49 (0.13 to 330.38)	-	MODERATE NO EV. OF IMP. DIFF.	CRITICAL
ow birth	weight (<2.5k	(g)						,				
`	randomised trials		no serious inconsistency	no serious indirectness	very serious ⁴	none	0/31 (0%)	0/26 (0%)	RD 0.00 (- 0.07 to +0.07)	-	VERY LOW	CRITICAL
											NO IMP. DIFF.	
Preeclamp	sia at birth											
	randomised trials		no serious inconsistency	no serious indirectness	very serious ³	none	0/28 (0%)	1/24 (4.2%)	POR 0.11 (0 to 5.84)	37 fewer per 1000 (from 42 fewer to 161 more)	VERY LOW	IMPORTANT
										,	NO IMP. DIFF.	
Gestationa	al diabetes m	ellitus at	24 to 28 weeks									

Optimum vitamin D dose during pregnancy for those medically classified as overweight or obese

1 (Corcoy 2020)	randomised trials		no serious inconsistency	no serious indirectness	very serious ³	none	6/31 (19.4%)	7/28 (25%)	RR 0.77 (0.3 to 2.03)	58 fewer per 1000 (from 175 fewer to 257 more)	VERY LOW NO EV. OF IMP. DIFF.	IMPORTANT
Gestation	al diabetes m	nellitus at	35 to 37 weeks									
1 (Corcoy 2020)	randomised trials		no serious inconsistency	no serious indirectness	very serious ³	none	11/29 (37.9%)	10/22 (45.5%)	`	(from 259 fewer to 273 more)	VERY LOW NO IMP. DIFF.	IMPORTANT

BMI: Body mass index; CI: confidence interval; LC-MS/MS: liquid chromatography-mass spectrometry/mass spectrometry; MD: mean difference; nmol/L: nanomoles per litre; (OH)D: hydroxyvitamin D: POR: Peto odds ratio; RD: risk difference: RR: relative risk

¹ Serious risk of bias in the evidence contributing to the outcomes as per RoB 2 ² 95% CI crosses 1 MID (0.5xcontrol group SD at baseline for maternal serum 25(0H)D serum concentration at birth = ±13.4)

³ 95% CI crosses 2 MIDs (0.8 and 1.25) ⁴ Event rate <150

Appendix G Economic evidence study selection

Study selection for: What dose of vitamin D is appropriate during pregnancy for women with a BMI medically classified as overweight or obese?

Figure 2. Flow diagram of selection process for economic evaluations



Appendix H Economic evidence tables

Economic evidence tables for review question: What dose of vitamin D is appropriate during pregnancy for women with a BMI medically classified as overweight or obese?

No economic evidence was identified which was applicable to these review question.

Appendix I Economic model

Economic model for review question: What dose of vitamin D is appropriate during pregnancy for women with a BMI medically classified as overweight or obese?

No economic analysis was conducted for this review question.

Appendix J Excluded studies

Excluded studies for review question: What dose of vitamin D is appropriate during pregnancy for women with a BMI medically classified as overweight or obese?

Excluded effectiveness studies

Table 7: Excluded studies and reasons for their exclusion

Study	Code [Reason]
Amberntsson, Anna, Barebring, Linnea, Winkvist, Anna et al. (2023) Vitamin D intake and determinants of vitamin D status during pregnancy in The Norwegian Mother, Father and Child Cohort Study. Frontiers in nutrition 10: 1111004	- Inappropriate study design Not an intervention study. Study assessed the total vitamin D intake from food and supplements in pregnant women
Amberntsson, Anna, Papadopoulou, Eleni, Winkvist, Anna et al. (2021) Maternal vitamin D intake and BMI during pregnancy in relation to child's growth and weight status from birth to 8 years: a large national cohort study. BMJ open 11(10): e048980	- No appropraite intervention Study assessed total vitamin D intake from both diets and supplements. Vitamin D from supplements alone was not reported.
Anonymous. (2023) Erratum: Maternal vitamin D intake and BMI during pregnancy in relation to child's growth and weight status from birth to 8 years: a large national cohort study (BMJ Open (2021) 11 (e048980) DOI: 10.1136/bmjopen-2021-048980). BMJ Open 13(3): e048980corr1	- No outcome of interest Study assessed total vitamin D intake from both diets and supplements. Vitamin D from supplements alone was not reported.
Begum, R, Coutinho, M L, Dormandy, T L et al. (1968) Maternal malabsorption presenting as congenital rickets. Lancet (London, England) 1(7551): 1048-52	- Inappropriate study design Case report
Bhowmik, B., Siddiquee, T., Mdala, I. et al. (2021) Vitamin D3 and B12 supplementation in pregnancy. Diabetes Research and Clinical Practice 174: 108728	- Country not of interest for this review Study conducted in Bangladesh which is not a high income country as stated in the protocol
Brustad, Nicklas, Garland, Juri, Thorsen, Jonathan et al. (2020) Effect of High-Dose vs Standard-Dose Vitamin D Supplementation in Pregnancy on Bone Mineralization in Offspring Until Age 6 Years: A Prespecified Secondary Analysis of a Double-Blinded, Randomized Clinical Trial. JAMA pediatrics 174(5): 419-427	- Population does not meet inclusion criteria All pregnant women were included. Results were not stratified by weight status or BMI categories
Catov, J.M., Bodnar, L.M., Olsen, J. et al. (2011) Periconceptional multivitamin use and risk of	- No outcome of interest

Study	Code [Reason]
preterm or small-for-gestational-age Births in the Danish National birth cohort. American Journal of Clinical Nutrition 94(3): 906-912	Observational study which reports an outcome that is already reported in the included RCTs
Chatzakis, Christos, Goulis, Dimitrios G, Mareti, Evangelia et al. (2019) Prevention of gestational diabetes mellitus in overweight or obese pregnant women: A network meta-analysis. Diabetes research and clinical practice 158: 107924	- Does not address condition or domain being studied Studies included in this systematic review do not assess vitamin D supplementation except 1(Corcoy 2020), which has been included in this review
Chawes BL, Bønnelykke K, Stokholm J et al. (2016) Effect of Vitamin D3 Supplementation During Pregnancy on Risk of Persistent Wheeze in the Offspring: A Randomized Clinical Trial. JAMA 315(4): 353-361	- Population does not meet inclusion criteria All pregnant women were included. Results were not separated by weight status or BMI category.
Christesen, Henrik T, Falkenberg, Tine, Lamont, Ronald F et al. (2012) The impact of vitamin D on pregnancy: a systematic review. Acta obstetricia et gynecologica Scandinavica 91(12): 1357-67	- Population does not meet inclusion criteria Systematic review has a mixed study population including all pregnant women irrespective of weight category and outcome data is not stratified by BMI categories. There are no studies relevant for inclusion.
Dieberger, A.M., Obermayer-Pietsch, B., Harreiter, J. et al. (2023) Physical activity and sedentary time across pregnancy and associations with neonatal weight, adiposity and cord blood parameters: a secondary analysis of the DALI study. International Journal of Obesity 47(9): 873-881	- No outcome of interest Secondary analysis of DALI trial reported in Corcoy 2020 with no additional relevant outcomes as per protocol.
Egan, Aoife M, Vellinga, Akke, Harreiter, Jurgen et al. (2017) Epidemiology of gestational diabetes mellitus according to IADPSG/WHO 2013 criteria among obese pregnant women in Europe. Diabetologia 60(10): 1913-1921	- No intervention of interest Study assessed risk factors for gestational diabetes and there was no intervention.
Gerveieeha, Z., Siassi, F., Qorbani, M. et al. (2023) The effect of vitamin D supplementation on body composition in nursing mothers with overweight or obesity: a randomized double-blind placebo-controlled clinical trial. BMC Nutrition 9(1): 1	- Population does not meet inclusion criteria Study included postpartum nursing mothers who delivered at term, and was conducted in Iran which is not a high income country as stated in the protocol.
Gould, J.F., Gibson, R.A., Green, T.J. et al. (2022) A Systematic Review of Vitamin D during Pregnancy and Postnatally and Symptoms of Depression in the Antenatal and Postpartum Period from Randomized Controlled Trials and Observational Studies. Nutrients 14(11): 2300	- Country not of interest for this review Systematic review including 2 RCTs conducted in Iran which is not a high income country as stated in the protocol. Other included studies were case control and cohort studies.

Study	Code [Reason]
Griffith, Rebecca J, Alsweiler, Jane, Moore, Abigail E et al. (2020) Interventions to prevent women from developing gestational diabetes mellitus: an overview of Cochrane Reviews. The Cochrane database of systematic reviews 6: cd012394	- Does not address condition or domain being studied Included systematic reviews are not relevant to this review as they either assessed dietary or exercise interventions, metformin or pharmacological interventions, which did not include vitamin D. One systematic review had vitamin D as an intervention but individual studies included that stratified population by weight category were carried out in LMICs
Groth, Susan W, Stewart, Patricia A, Ossip, Deborah J et al. (2017) Micronutrient Intake Is Inadequate for a Sample of Pregnant African- American Women. Journal of the Academy of Nutrition and Dietetics 117(4): 589-598	- No intervention of interest Study examined dietary micronutrient intake in pregnancy.
Harreiter, Jurgen, Mendoza, Lilian C, Simmons, David et al. (2022) Vitamin D3 Supplementation in Overweight/Obese Pregnant Women: No Effects on the Maternal or Fetal Lipid Profile and Body Fat Distribution-A Secondary Analysis of the Multicentric, Randomized, Controlled Vitamin D and Lifestyle for Gestational Diabetes Prevention Trial (DALI). Nutrients 14(18)	- No outcome of interest Secondary analysis of DALI trial reported in Corcoy 2020 with no additional relevant outcomes as per protocol.
Harvey, Nicholas C, Holroyd, Christopher, Ntani, Georgia et al. (2014) Vitamin D supplementation in pregnancy: a systematic review. Health technology assessment (Winchester, England) 18(45): 1-190	- Does not address condition or domain being studied Systematic review has a mixed study population including all pregnant women irrespective of weight category and outcome data is not stratified by BMI categories. There are no studies relevant for inclusion.
Luo, T., Lin, Y., Lu, J. et al. (2022) Effects of vitamin D supplementation during pregnancy on bone health and offspring growth: A systematic review and meta-analysis of randomized controlled trials. PLoS ONE 17(10october): e0276016	- Population does not meet inclusion criteria Systematic review has a mixed study population including all pregnant women irrespective of weight category and outcome data is not stratified by BMI categories. Individual studies were conducted in HICs and LMICs. There are no studies relevant for inclusion.
Lutke-dorhoff, M, Schulz, J, Westendarp, H et al. (2022) Comparative Study of the Effects of Two Dietary Sources of Vitamin D on the Bone Metabolism, Welfare and Birth Progress of Sows Fed Protein- and Phosphorus-Reduced Diets. Animals 12(13)	- Does not address condition or domain being studied Non-human study. Study was on pigs.

Study	Code [Reason]
Ma, Shuangshuang, Yin, Wanjun, Wang, Peng et al. (2023) Effect of vitamin D supplementation on glucose control in mid-late gestation: A randomized controlled trial. Clinical nutrition (Edinburgh, Scotland) 42(6): 929-936	- Country not of interest for this review Study conducted in China, which is not a high income country as stated in the protocol
Mead, Molly J, McWhorter, Caroline A, Rodgers, Megan D et al. (2023) Does maternal vitamin D status influence placental weight or vascular and inflammatory pathology? Secondary analysis from the Kellogg Pregnancy Study. The Journal of steroid biochemistry and molecular biology 233: 106358	- Population does not meet inclusion criteria All pregnant women were included. Results were not stratified by weight status or BMI categories. Primary study of this secondary analysis equally assessed and was not eligible for inclusion
Moon, Rebecca J, Harvey, Nicholas C, Cooper, Cyrus et al. (2016) Determinants of the Maternal 25-Hydroxyvitamin D Response to Vitamin D Supplementation During Pregnancy. The Journal of clinical endocrinology and metabolism 101(12): 5012-5020	- Population does not meet inclusion criteria All pregnant women were included. Results were not stratified by weight status or BMI category
Morse, Nancy L (2012) Benefits of docosahexaenoic acid, folic acid, vitamin D and iodine on foetal and infant brain development and function following maternal supplementation during pregnancy and lactation. Nutrients 4(7): 799-840	- Inappropriate study design Not an intervention study. Study compared 25(OH)D concentrations in women who developed GDM to women who did not develop GDM
Nadeem, Amna, Saeed, Muniza, Sadiqa, Ayesha et al. (2023) The Effect of Vitamin D3 Intervention on the Association Among Vitamin D3, Adiponectin, and Body Mass Index in Pregnant Women With Gestational Diabetes. Cureus 15(8): e43506	- Country not of interest for this review Study conducted in Pakistan which is not a high income country as stated in the protocol
Pena, Homero Rabelo, de Lima, Marilia Carvalho, Brandt, Katia Galeao et al. (2015) Influence of preeclampsia and gestational obesity in maternal and newborn levels of vitamin D. BMC pregnancy and childbirth 15: 112	- Country not of interest for this review Study conducted in Brazil which is not a high income country as stated in the protocol
Rached, Veronica, Diogenes, Maria Eduarda Leao, Donangelo, Carmen Marino et al. (2023) Calcium plus vitamin D supplementation during pregnancy reduces postpartum fat mass in adolescents: A randomized trial. American journal of human biology: the official journal of the Human Biology Council 35(9): e23911	- Country not of interest for this review Study conducted in Brazil which is not a high income country as stated in the protocol
Redfern, Kathy M., Hollands, Heidi J., Welch, C. Ross et al. (2022) Dietary Intakes of Folate,	- No intervention of interest

Study	Code [Reason]
Vitamin D and Iodine during the First Trimester of Pregnancy and the Association between Supplement Use and Demographic Characteristics amongst White Caucasian Women Living with Obesity in the UK. Nutrients 14(23): 5135	Study examined dietary micronutrient intake in pregnancy
Samimi, M, Kashi, M, Foroozanfard, F et al. (2016) The effects of vitamin D plus calcium supplementation on metabolic profiles, biomarkers of inflammation, oxidative stress and pregnancy outcomes in pregnant women at risk for pre-eclampsia. Journal of human nutrition and dietetics 29(4): 505-515	- Country not of interest for this review Study conducted in Iran which is not a high income country as stated in the protocol
Sauder, K.A., Harte, R.N., Ringham, B.M. et al. (2021) Disparities in Risks of Inadequate and Excessive Intake of Micronutrients during Pregnancy. Journal of Nutrition 151(11): 3555-3569	- No outcome of interest Study outcome was the risk of inadequate and excessive nutrients from diet and supplements
Shahgheibi, S.; Farhadifar, F.; Pouya, B. (2016) The effect of vitamin D supplementation on gestational diabetes in high-risk women: Results from a randomized placebo-controlled trial. Journal of Research in Medical Sciences 21(1)	- Country not of interest for this review Study conducted in Iran which is not a high income country as stated in the protocol
Sunarno, Rita Dewi, Kartasurya, Martha Irene, Suwondo, Ari et al. (2023) Vitamin D Supplementation and Sun Exposure Maintain Blood Pressures of Pregnant Women and Increase Birth Weight in a Randomized Controlled Trial. Iranian journal of public health 52(10): 2148-2156	- Country not of interest for this review Study conducted in Indonesia which is not a high income country as stated in the protocol
Valkama, Anita J, Meinila, Jelena M, Koivusalo, Saila B et al. (2018) Body size modifies the relationship between maternal serum 25-hydroxyvitamin D concentrations and gestational diabetes in high-risk women. European journal of clinical nutrition 72(3): 460-463	- No intervention of interest Study assessed the change in 25(OH)D during pregnancy in relation to maternal BMI
Vanderlelie, Jessica, Scott, Rani, Shibl, Rania et al. (2016) First trimester multivitamin/mineral use is associated with reduced risk of preeclampsia among overweight and obese women. Maternal & Child Nutrition 12(2): 339-348	- No intervention of interest Vitamin D reported as part of multivitamin and the specific dose of vitamin D was not reported
Vestergaard, Anna Louise, Justesen, Signe, Volqvartz, Tabia et al. (2021) Vitamin D insufficiency among Danish pregnant women-	- No intervention of interest

Study	Code [Reason]
Prevalence and association with adverse obstetric outcomes and placental vitamin D metabolism. Acta obstetricia et gynecologica Scandinavica 100(3): 480-488	Study assessed the correlation between vitamin D levels in early pregnancy and pregnancy outcomes
Zerofsky, Melissa S, Jacoby, Bryon N, Pedersen, Theresa L et al. (2016) Daily Cholecalciferol Supplementation during Pregnancy Alters Markers of Regulatory Immunity, Inflammation, and Clinical Outcomes in a Randomized Controlled Trial. The Journal of nutrition 146(11): 2388-2397	- Population does not meet inclusion criteria All pregnant women were included. Results were not stratified by weight status or BMI category

Excluded economic studies

No economic study was reviewed at full text and excluded from this review.

Appendix K Research recommendations – full details

Research recommendations for review question: What dose of vitamin D is appropriate during pregnancy for people with a BMI medically classified as overweight or obese?

K.1.1 Research recommendation

What dose of vitamin D is appropriate during pregnancy for people with a BMI that is within the overweight or obese weight categories?

K.1.2 Why this is important

In the UK it is recommended that all adults take a daily 10µg supplement of Vitamin D. Optimum vitamin D status is associated with better bone health and may be linked to reduced risk of gestational diabetes. Those with higher adiposity may need additional vitamin D, especially during pregnancy, but the identified evidence was not clear.

K.1.3 Rationale for research recommendation

Table 8: Research recommendation rationale

Importance to 'patients' or the population	Those starting pregnancy with a BMI >25 kg/m² may be at a greater risk of some adverse pregnancy outcomes compared to those starting pregnancy with a healthy BMI. The risk of some of these outcomes (e.g. gestational diabetes) could potentially be reduced by having a better vitamin D status. Supplementing with vitamin D over and above what is recommended for the general UK population may be a simple preventable action that could improve some pregnancy outcomes for people with a BMI medically classified as overweight or obese
Relevance to NICE guidance	Vitamin supplementation and weight status during pregnancy are important aspects of these guidelines. Only two RCTs were identified for this research question which did not provide enough evidence on the optimum Vitamin D dose in this population group.
Relevance to the NHS	Optimal vitamin D status during pregnancy may be associated with reduced risk of gestational diabetes. In turn, gestational diabetes increases the risk of subsequent type 2 diabetes. Both of these aspects result in poorer health outcomes and greater use of NHS resources.
National priorities	High
Current evidence base	Two RCTs were included in the review. One study compared medium dose vitamin D supplementation of 10µg (400IU)/day with another medium dose vitamin D supplementation of 20µg (800IU)/day (Alhomaid 2021). However, no important difference was found in the same group of women at 28 weeks gestational age. No important difference or no evidence of an important difference was found in women medically classified as obese at the same time points. No other relevant outcomes were reported for this comparison group. Another study compared no vitamin D supplementation to a high dose vitamin D supplementation of 40µg (1600IU)/day (Corcoy 2020). There was important benefit for Vitamin D supplementation compared to placebo in terms of maternal serum 25(OH)D concentration at 24 to 28 and 35 to 37 weeks gestational age, and at delivery. No important differences were identified in this comparison for other outcomes identified, such as preterm birth, low birth weight (<2.5kg), preeclampsia or gestational diabetes. Given the lack of robust evidence, the committee did not make a recommendation and decided to make a research recommendation to help inform future guidelines.

Equality
considerations

Ethnicity and socio economic factors

K.1.4 Modified PICO table

Table 9: Research recommendation modified PICO table

Population	Inclusion:
	Women during a single or multiple pregnancy, for women with a BMI
	that is within the overweight or obese weight categories
Intervention	 Vitamin D supplementation (20μg/800IU daily)
	Vitamin D supplementation (40µg/1600IU daily)
	• Vitamin D supplementation (60-100µg/2400-4000IU daily)
	Exclude: vitamin D in combination with other vitamins and minerals
Comparator	comparisons
	 different doses in the intervention vs Vitamin D supplementation (10μg/ 400IU to 20μg/800IU daily)
	different doses in the intervention compared to each other
Outcome	Primary outcomes
	Maternal serum 25(OH)D concentration and neonatal (umbilical cord)
	25(OH) D concentration
	Gestational diabetes
	Low birth weight (<2500 g)
	Secondary outcomes
	Pre-eclampsia
	Bone mass and bone density in infant Childhand annuas and require to metion.
	Childhood eczema and respiratory function Infent hyposologomia
	Infant hypocalcaemia
	Cost-effectiveness (including resource use measurements and QALY
	estimations using a validated preference-based measure such as the EQ-5D or SF-6D).
Study design	• RCTs
Timeframe	For the secondary outcomes listed (bone mass, childhood eczema, infant hypocalcaemia), follow up would need to be until the offspring is age 2 years old at a minimum
Additional	Sub-group analysis:
information	1. BMI thresholds on booking:
	∘ Overweight range: 25 to 29.99 kg/m²
	o Obesity range 1: 30 to 34.99 kg/m²
	o Obesity range 2: 35 to 39.99 kg/m²
	o Obesity range 3: >40 kg/m²
	Single versus multiple pregnancy

- 3. Comorbidities (yes versus no)
- 4. Ethnicity
 - o White/White British
 - o Asian/Asian British
 - o Black/African/Caribbean/Black British
 - o Mixed/Multiple ethnic groups
 - o Other ethnic group
- 5. Socioeconomic groups (high vs low) (deprivation measured using IMD)
- 6. Sunlight exposure and use of SPF (sufficient vs not sufficient). To also consider effect of seasons on Vitamin D sufficiency.
- 7. Baseline vitamin D level (low vs high)

BMI: body mass index; (OH)D: hydroxyvitamin D; QALY: quality-adjusted life year; RCTs: randomised controlled trials; SPF: sun protection factor