National Institute for Health and Care Excellence

Draft for consultation

Maternal and child nutrition

[A] Evidence review for high-dose folic acid supplementation before and during the first 12 weeks of pregnancy

NICE guideline number tbc

Evidence review underpinning recommendations 1.1.4 to 1.1.6 and research recommendation on high-dose folic acid supplementation in the NICE guideline

June 2024

Draft for consultation

This evidence review was developed by NICE



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1 High-dose folic acid supplementation

before and during the first 12 weeks of

3 pregnancy

4 Review question

- 5 Which groups of women should be advised to take high-dose folic acid supplements before
- 6 and during the first 12 weeks of pregnancy?

7 Introduction

- 8 There is uncertainty as to which high risk groups may benefit from a higher dose (5 mg) of
- 9 folic acid to help reduce the risk of neural tube defects (NTDs) and this uncertainty has led to
- variations in international guidance. The WHO and NICE guidance both recommend a higher
- 11 dose folic acid supplement and increased food intake of folate for those with previous
- 12 pregnancy affected by NTDs, who have diabetes, or taking anticonvulsant treatment. The
- 13 aim of this review is to determine which groups should be advised to take high-dose folic acid
- supplements before and during the first 12 weeks of pregnancy.

15 Summary of the protocol

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

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

omen in the preconception stage trying to become pregnant or women uring the first 12 weeks of a single or multiple pregnancy gh-dose folic acid supplementation (≥5 mg daily) lone or in combination with other vitamins and minerals) Low-dose folic acid supplementation (<1 mg daily) Medium-dose folic acid supplementation (≥1 to <5 mg daily) Placebo No intervention lone or in combination with other vitamins and minerals)
lone or in combination with other vitamins and minerals) Low-dose folic acid supplementation (<1 mg daily) Medium-dose folic acid supplementation (≥1 to <5 mg daily) Placebo No intervention
Medium-dose folic acid supplementation (≥1 to <5 mg daily) Placebo No intervention
,
ritical: neural tube defects (NTDs) in the baby birthweight hypertensive disorders of pregnancy (pre-eclampsia and gestational hypertension) nportant: RBC (or serum if RBC is not reported) folate concentrations in the mother neurodevelopmental outcomes (dichotomous outcome, not continuous outcomes such as mean change in score): o severe (score of >2SD below normal on validated assessment scales, or Bayley assessment scale of MDI or PDI) o moderate (score of 1-2 SD below normal on validated assessment scales, or Bayley assessment scale MDI or PDI 70-84). congenital heart defects midline facial defects such as cleft lip or cleft palate
rrkhrc

- 1 MDI: mental development index; mg: milligrams; PDI: psychomotor developmental index; RBC: red blood cell;
- 1 MDI: mental developme 2 SD: standard deviation
- 3 For further details see the review protocol in appendix A.

4 Methods and process

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

10 Effectiveness evidence

11 Included studies

- 12 No evidence was found on high dose folic acid supplementation (≥5 mg/day), the dose
- 13 currently recommended in the UK for pregnant women at high risk of NTDs. The committee
- agreed to make a deviation to the protocol and relax the inclusion criteria to include folic acid
- 15 supplementation of 4 mg/day, in the absence of evidence on dose of ≥5 mg/day, but
- downgraded the evidence for indirectness.
- 17 Three randomised controlled trials were included in this review (Laurence 1981, MRC 1991,
- 18 Wen 2018).
- 19 The included studies are summarised in Table 2.
- Two studies reported on women with a history of births with NTDs (Laurence 1981, MRC
- 21 1991), and 1 study reported on women at high risk of pre-eclampsia (Wen 2018).
- Two studies compared 4 mg folic acid per day to placebo (Laurence 1981, Wen 2018). One
- 23 study compared 4 mg folic acid per day to placebo and also had a third arm comparing 4 mg
- folic acid with multivitamins per day to placebo (MRC 1991). In Wen 2018, participants in
- both arms included women taking daily prenatal vitamins or low dose folic acid supplements
- 26 (up to 1.1 mg of folic acid) throughout the study period.
- 27 One study was conducted in the United Kingdom (Laurence 1981), 1 study was conducted in
- Australia, Canada, France, Hungary, Israel, Russia and UK (MRC 1991) and 1 study was
- 29 conducted in Argentina, Australia, Canada, Jamaica and UK (Wen 2018).
- There was no evidence available for the following outcomes: birthweight,
- 31 neurodevelopmental outcomes, congenital heart defects and midline facial defects such as
- 32 cleft lip or cleft palate outcomes.
- 33 As per protocol the evidence was stratified according to women with a previous history of a
- pregnancy affected by NTDs. Pre-eclampsia was not a stratum in the protocol, however as
- 35 the risk of neural tube defects in this population is different to those with previous history of
- NTDs, the committee agreed that women with high risk of pre-eclampsia should be analysed
- 37 separately. There was not sufficient evidence to stratify evidence based on parity, women
- taking antiseizure medications, women taking antifolate medications, women with diabetes,
- and women who started folic acid pre-conception.
- 40 As per protocol, subgroup analyses for the following groups was agreed to be conducted if
- 41 there was heterogeneity: deprived socioeconomic group, age <40 versus ≥40, women and
- 42 parents with disabilities, including learning disabilities and other physical and mental health
- 43 conditions, women going through assisted conception, religion, cultural considerations and

- 1 ethnicity, and women who smoke versus not. There was no heterogeneity identified in the
- 2 evidence, hence subgroup analysis was not conducted.
- 3 See the literature search strategy in appendix B and study selection flow chart in appendix C.

4 Excluded studies

- 5 Studies not included in this review are listed, and reasons for their exclusion are provided in
- 6 appendix J.

7 Summary of included studies

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

9 Table 2: Summary of included studies

Study	Population	Intervention	Comparison	Outcomes	Comments
Laurence 1981 RCT UK	N=111 women with a history of births with NTDs Participant characteristics : NR	4mg folic acid Taken as a 2mg tablet, twice a day from the time contraception stopped until birth	Placebo	 Neural tube defects Red blood cell folate concentration 	Follow-up: at birth Strata: history of births with NTDs
MRC 1991 RCT Australia, Canada, France, Hungary, Israel, Russia, UK	N=893 women with a history of births with NTDs Maternal age, mean (SD: NR): Intervention Group A: 27 Group B: 27.4 Control: 26.8 Maternal BMI: NR Livebirths in previous pregnancies, mean: Group A (4mg folic acid): 0.88 Group B (4mg folic acid + multivitamin): 0.94 Group C (placebo): 0.91	Group A: 4mg folic acid Group B: 4mg folic acid + multivitamin Taken as 1 tablet, daily, from randomisatio n until 12 weeks gestational age	Group C: Placebo Contains dried ferrous sulphate (120mg) and di-calcium phosphate (240mg)	Neural tube defects	Multivitamin composition: vitamin A 4000 IU, D 400 IU, BI 1-5mg, B2 1.5 mg, B6 10mg, C 40mg, and nicotinamide 15mg Follow-up: at 12 weeks of pregnancy Strata: history of births with NTDs

Mean number of previous NTD pregnancies: Group A (4mg folic acid): 1.03 Group B (4mg folic acid + multivitamin): 1.04 Group C (placebo): 1.01 Wen 2018 N=2301 women at high risk of pre-eclampsia Argentina, Australia, Canada, Jamaica, UK Jamaica, UK Maternal age, mean(SD): Intervention: 31(5.4) Placebo: 31(5.4) Placebo: 31(5.4) Pregnancy BMI, mean (SD) Intervention: 34 (8.6) Control: 34 (13) Parity, n (%): Intervention: 0 - 413 (33.7) 1 - 498 (40.6) ≥2 - 316 (25.7) Control: 0 - 420 (34.0) 1 - 499 (40.4) ≥2 - 317 (25.6) History of pre-eclampsia:	04	D	Inter-		0.1	0
of previous NTD pregnancies: Group A (4mg folic acid): 1.03 Group B (4mg folic acid): 1.04 Group C (placebo): 1.01 Wen 2018 N=2301 Argentina, Australia, Canada, Jamaica, UK Placebo 31(5.4) Pregnancy BMI, mean (SD) Intervention: 34 (8.6) Control: 34 (13) Parity, n (%): Intervention: 0 - 413 (33.7) 1 - 498 (40.6) ≥2 - 316 (25.7) Control: 0 - 420 (34.0) 1 - 499 (40.4) ≥2 - 317 (25.6) History of preeclampsia:	Study	Population	Intervention	Comparison	Outcomes	Comments
RCT high risk of pre-eclampsia Argentina, Australia, Canada, Jamaica, UK Pregnancy BMI, mean (SD) Intervention: 34 (8.6) Control: 34 (13) Parity, n (%): Intervention: 0 - 413 (33.7) 1 - 498 (40.6) ≥2 - 316 (25.7) Control: 0 - 420 (34.0) 1 - 499 (40.4) ≥2 - 317 (25.6) History of pre-eclampsia: Taken as a 1 mg tablet, 4 times a day from 8 to 16 weeks gestational age until birth Taken as a 1 mg tablet, 4 times a day from 8 to 16 weeks gestational age until birth Follow-up: at 42 days postpartur Strata: high risk of pre-eclampsia: both arms were allowed to take daily prenatal vitamins or low dose folic acid. Taken as a 1 mg tablet, 4 times a day from 8 to 16 weeks gestational age until birth Follow-up: at 42 days postpartur Strata: high risk of pre-eclampsia: both arms were allowed to take daily prenatal vitamins or low dose folic acid. Strata: high risk of pre-eclampsia: both arms were allowed to take daily prenatal vitamins or low dose folic acid. Strata: high risk of pre-eclampsia: both arms were allowed to take daily prenatal vitamins or low dose folic acid. Strata: high risk of pre-eclampsia: both arms were allowed to take daily prenatal vitamins or low dose folic acid. Strata: high risk of pre-eclampsia: both arms were allowed to take daily prenatal vitamins or low dose folic acid. Strata: high risk of pre-eclampsia: both arms were allowed to take daily prenatal vitamins or low dose folic acid. Strata: high risk of pre-eclampsia: both arms were allowed to take daily prenatal vitamins or low dose folic acid. Strata: high risk of pre-eclampsia: both arms were allowed to take daily prenatal vitamins or low dose folic acid.		of previous NTD pregnancies: Group A (4mg folic acid): 1.03 Group B (4mg folic acid + multivitamin): 1.04 Group C (placebo):				
Intervention: 308 (25.1) Control: 303 (24.5)	RCT Argentina, Australia, Canada,	N=2301 women at high risk of pre-eclampsia Maternal age, mean(SD): Intervention: 31(5.4) Placebo: 31(5.4) Pregnancy BMI, mean (SD) Intervention: 34 (8.6) Control: 34 (13) Parity, n (%): Intervention: 0 - 413 (33.7) 1 - 498 (40.6) ≥2 - 316 (25.7) Control: 0 - 420 (34.0) 1 - 499 (40.4) ≥2 - 317 (25.6) History of pre-eclampsia: Intervention: 308 (25.1) Control: 303	Taken as a 1mg tablet, 4 times a day from 8 to 16 weeks gestational age until	Placebo		both arms were allowed to take daily prenatal vitamins or low dose folic acid supplements (up to 1.1 mg of folic

BMI: body mass index; IU: international units; mg: milligrams; μg: micrograms; N/A: not applicable; NR: not reported; NTDs: neural tube defects; RCT: randomised controlled trial; SD: standard deviation

1

1 See the full evidence tables in appendix D and the forest plots in appendix E.

2 Summary of the evidence

- 3 Folic acid supplementation 4 mg/day versus placebo in women with a history of births
- 4 with neural tube defects (NTDs)
- 5 Two studies were included under this comparison.
- 6 Evidence for 4 mg folic acid per day versus placebo for women with a history of births with
- 7 NTDs showed that there was an important benefit for 4 mg folic acid per day compared to the
- 8 placebo for the outcomes of NTDs (at 12 weeks of pregnancy and birth follow-up) and red
- 9 blood cell folate concentration (at birth follow-up) favouring folic acid supplementation over
- 10 placebo.
- 11 The evidence was of low quality.
- 12 Folic acid supplementation 4 mg/day + multivitamin versus placebo for women with a
- 13 history of births with neural tube defects (NTDs)
- 14 One study was included under this comparison.
- 15 Evidence for 4 mg folic acid per day plus multivitamins versus placebo for women with a
- history of births with NTDs showed that there was an important benefit for daily 4 mg folic
- 17 acid plus multivitamins intake compared to the placebo for the outcome of NTDs at 12 weeks
- 18 of pregnancy follow-up.
- 19 The evidence was of very low quality.
- 20 Folic acid supplementation 4 mg/day versus placebo in women at high risk of
- 21 preeclampsia
- 22 One study was included under this comparison.
- 23 Evidence for 4 mg folic acid per day versus placebo for women at high risk of preeclampsia
- 24 showed that there was no important difference between daily 4 mg folic acid intake
- compared to placebo for the outcome of severe pre-eclampsia at 42 days post-partum follow-
- 26 up.
- The evidence was very low quality.
- 28 See appendix F for full GRADE tables.

29 Economic evidence

30 Included studies

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

34 Excluded studies

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

36 Economic model

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

1 Unit costs

2

Table 3: Unit costs of folic acid

Resource	Unit costs	Source
Folic acid 5 mg tablets	28 for £0.68	NHS Business Services Authority, NHS Prescription Services. NHS England and Wales. Electronic Drug Tariff. Issue: May 2024. Compiled on the behalf of the Department of Health and Social Care. Available from: https://www.nhsbsa.nhs.uk/pharmacies-gp- practices-and-appliance-contractors/drug-tariff
Folic acid 400 microgram tablets	90 for £3.27	NHS Business Services Authority, NHS Prescription Services. NHS England and Wales. Electronic Drug Tariff. Issue: May 2024. Compiled on the behalf of the Department of Health and Social Care. Available from: https://www.nhsbsa.nhs.uk/pharmacies-gp- practices-and-appliance-contractors/drug-tariff

3 The committee's discussion and interpretation of the evidence

4 The outcomes that matter most

- 5 Neural tube defects in the baby, birthweight, and hypertensive disorders of pregnancy were
- 6 prioritised as critical outcomes by the committee. These outcomes were identified as critical
- 7 because high dose maternal folic acid supplements can reduce the risk of NTDs in the baby
- 8 and observational studies have suggested a relationship between low maternal folic acid
- status in the mother with low birthweight and maternal hypertensive disease in pregnancy.
- 10 The committee agreed that red blood cell folate concentrations in the mother should be an
- important outcome as it is a critical biomarker for folate status. Congenital heart defects, and
- 12 midline facial defects were also deemed important outcomes because folic acid protects
- babies against some birth defects other than NTDs including heart and midline facial defects.
- 14 Relationships have also been shown between low maternal folate status and
- 15 neurodevelopmental disorders other the NTDs.
- No evidence was identified for relationships between high dose folic acid supplementation
- and congenital heart defects with midline facial defects, birthweight, neurodevelopmental,
- 18 outcomes other than NTDs.

19 The quality of the evidence

- The quality of the evidence was assessed using GRADE methodology. The quality of the
- 21 evidence ranged from very low to low quality. The main reasons for downgrading were risk of
- bias (for example, there were some concerns with bias in selection of the reported result,
- bias due to missing outcome data, and bias in the measurement of the outcome), imprecision
- due to small number of events rate and indirectness due to no intervention available of high
- 25 dose folic acid supplements (≥5 mg) and due to the population including people with body
- 26 mass index ≥35kg/m². Risk of bias was most commonly due to deviations from the intended
- 27 interventions and missing outcome data.
- 28 Studies were assessed for quality using the Cochrane Risk of Bias tool version 2. One study
- 29 (Wen 2018) included participants and settings that were not directly relevant to the study
- protocol (that is, women with a BMI ≥35kg/m² and trial centres in low and middle income
- 31 countries (LMICs)). Overall, the population from LMICs was very small and not considered to
- 32 affect the quality of the evidence, however the evidence was downgraded for indirectness as

- 1 >33% of the population had a BMI ≥35kg/m². All 3 included studies compared 4 mg/day folic
- 2 acid supplement with placebo. These studies were downgraded for indirectness as the dose
- 3 of folic acid did not meet the intervention high dose folic acid supplement threshold in the
- 4 protocol.

5

7

Benefits and harms

- 6 Overall, the committee agreed that the evidence base is sparse in this topic area, with no
 - studies identified on high dose folic acid (≥5 mg per day). The committee discussed that 4
- 8 mg of folic acid was close to 5 mg and clinically a high dose compared to the standard 400
- 9 micrograms.
- 10 The committee discussed that current recommendation advising those at higher risk of NTDs
- 11 to take 5 mg folic acid per day are based on clinical consensus rather than clinical evidence.
- 12 The committee were aware of information about how and when to take folic acid on the NHS
- website, which recommends 5 mg folic acid per day for people with folate deficiency
- anaemia, and discussed that this dose had probably been extended to include pregnant
- people with a higher risk of NTDs given the availability of this dose in pharmacies. However,
- 16 this is not only the case in the UK as 5mg dose is commonly recommended in other
- 17 countries as well to pregnant populations at higher risk of NTDs.
- 18 Low quality evidence showed that 4 mg folic acid per day in women with a history of births
- 19 with NTDs had an important benefit on reducing neural tube defects in the baby and
- 20 increasing red blood cell folate concentration in the pregnant woman. The committee agreed
- 21 that ideally they would recommend 4 mg of folic acid for those at higher risk of NTDs
- because that is based on the available evidence. However, in the UK, the only folic acid
- 23 formulations available are 400 micrograms and 5 mg and it would be impractical to suggest
- that people should take 10 of the 400-microgram tablets daily. Folic acid is not a narrow
- 25 therapeutic index drug, meaning that there is likely no big difference between the 4 mg and 5
- 26 mg dose. Furthermore, folic acid is generally well tolerated even in high doses. While
- evidence on folic acid toxicity among different populations was not reviewed by the
- committee, it is known that there is no evidence to suggest harm from 5 mg folic acid per day
- in different populations, although the evidence base is limited. Concerns have been raised
- 30 about the potential exacerbation of neuropathy in individuals with vitamin B12 deficiency
- when treated with high dose folic acid, instead of vitamin B12. However, this analysis
- 32 appears to have been incorrect (Wald 2018). Regardless, this has little relevance for our
- 33 population of interest. Because of no known harm of high dose folic acid, the
- recommendation continues as 5 mg of folic acid for those at higher risk of NTDs.
- 35 The evidence was only available for those with history of NTDs and those at risk of
- 36 preeclampsia so the committee agreed based on their expertise, the other groups who
- 37 should be offered high dose of folic acid. No evidence was identified investigating high dose
- 38 folic acid in pregnant women with pre-gestational type 1 or type 2 diabetes who are
- 39 anticipating pregnancy or in the first 12 weeks of pregnancy, however, the committee were
- 40 aware that pre-gestational diabetes is associated with increased risk of babies being born
- 41 with NTDs or other congenital malformations. The committee agreed that the mechanism
- 42 involves uncontrolled blood glucose which can lead to changes in methylation status in key
- 43 genes including raised oxidative stress, mitochondria dysfunction and rates of apoptosis and
- cell death in neurodevelopment pathways. Because of the higher risk of NTDs in diabetic
- 45 population, the committee agreed that these women should be advised to take high dose
- 46 folic acid.
- 47 No evidence was identified investigating high dose folic acid in pregnant women with sickle
- 48 cell anaemia and thalassaemia, in pregnant women with epilepsy taking anti-epileptic
- 49 medication or in women taking medication that affects the absorption of folic acid. Therefore,
- the committee made their recommendations based on their knowledge and experience. The
- 51 committee were aware of groups of people that require a higher dose of folic acid. For

- 1 example, those with sickle cell anaemia are at an increased risk of folate deficiency due to 2 haemolytic anaemia and therefore require a high dose to compensate for the increased 3 demand for folate during pregnancy. The committee noted that those with thalassaemia may 4 similarly have a greater risk of their baby developing neural tube defects. The committee also 5 discussed those with epilepsy taking anti-epileptic medications and was aware that these 6 medications interact with folate metabolism resulting in low levels of serum and red blood cell 7 folate concentrations. The committee discussed other groups of people taking medication 8 that interferes with folic acid absorption, such as those taking HIV medication. The committee 9 agreed that a high dose should be given to these groups of people to reduce the risk of births 10 with neural tube defects.
- 11 The committee agreed that since there is no evidence for 5 mg folic acid for women at 12 greater risk of delivering a baby with a NTD, it is important to determine whether this dose can result in over-supplementation. Especially given the UK government decision from 2021 13 of mandatory fortification of white flour with folic acid. Therefore, the committee agreed to 14 15 make a research recommendation on 5 mg folic acid per day in pregnant people with a high risk of NTDs. The committee also included pregnant people with a BMI of ≥25 kg/m² in the 16 17 research recommendation. Pregnant people with a BMI of ≥25 kg/m² were excluded from this review, however appropriate folic acid dose for this group was included in evidence review B. 18 but there was limited evidence. See Appendix K for more details. 19
- There was some very low-quality evidence identified from 1 RCT that reported 4 mg of folic acid per day beyond the first trimester showed no important difference for pre-eclampsia in women at high risk of developing this condition compared to placebo. The committee were aware that this group are not currently prescribed high dose folic acid in clinical practice. The committee agreed with the evidence that high dose folic acid did not reduce the risk of pre-eclampsia and therefore made a recommendation in line with the evidence.

Cost effectiveness and resource use

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27 No economic evidence was identified for this review question. Overall, the recommendations 28 reflect current practice, so no resource implications are expected. The daily cost of the high 29 dose 5 mg tablets (28 for £0.68) is slightly lower than the daily cost of the standard dose 400 microgram tablets (90 for £3.27). It is noted, however, that the 400 microgram tablets are 30 31 available over the counter (and are usually bought over the counter), whereas the 5 mg 32 tablets are available by prescription only (so, a person requiring the high dose tablets 33 normally needs to pay the prescription charge of currently £9.90 pre-conception unless they 34 are exempt, but that the prescription is free during pregnancy). Offering the higher dose 35 tablet to people who would benefit from it is expected to lead to important benefits and cost-36 savings resulting from prevention of babies born with neural tube defects.

Other factors the committee took into account

38 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 39 40 searches were not designed to specifically look for evidence on trans men or non-binary people but they were also not excluded. However, there is a small chance evidence on them 41 42 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, 43 44 and used gender inclusive language to promote equity, respect and effective communication with everyone. Healthcare professionals should use their clinical judgement when 45 46 implementing the recommendations, taking into account each person's circumstances, needs 47 and preferences, and ensuring all people are treated with dignity and respect throughout 48 their care.

1 Recommendations supported by this evidence review

- 2 This evidence review supports recommendations 1.1.4 and 1.1.6 and the research
- 3 recommendation on high-dose folic acid supplementation.

References – included studies 4

_	Fffectiveness	
7		

6 7

> 8 9

Laurence 1981

Laurence, K M, James, N, Miller, M H et al. (1981) Double-blind randomised controlled trial of folate treatment before conception to prevent recurrence of neural-tube defects. British medical journal (Clinical research ed.) 282(6275): 1509-11

10 11 12

MRC 1991

MRC (1991) Prevention of neural tube defects: results of the Medical Research Council 13 Vitamin Study. MRC Vitamin Study Research Group. Lancet (London, England) 338(8760): 14

15 131-7

16 17

Wen 2018

18 Wen, Shi Wu, White, Ruth Rennicks, Rybak, Natalie et al. (2018) Effect of high dose folic acid supplementation in pregnancy on pre-eclampsia (FACT): double blind, phase III, 20 randomised controlled, international, multicentre trial. BMJ (Clinical research ed.) 362: k3478

21

19

22 Other

23

24 Wald, N.J., Morris, J.K. & Blakemore, C. (2018) Public health failure in the prevention of 25 neural tube defects: time to abandon the tolerable upper intake level of folate. Public Health Rev 39(2) 26

27

Appendices

2 Appendix A Review protocols

- 3 Review protocol for review question: Which groups of women should be advised to take high-dose folic acid supplements
- 4 before and during the first 12 weeks of pregnancy?

5 Table 4: Review protocol

Field	Content
PROSPERO registration number	CRD42022346779
Review title	High-dose folic acid supplementation before and during the first 12 weeks of pregnancy
Review question	Which groups of women should be advised to take high-dose folic acid supplements before and during the first 12 weeks of pregnancy?
Objective	To determine which groups of women should be advised to take high-dose folic acid supplements before and during the first 12 weeks of pregnancy.
Searches	The following databases will be searched:
	Cochrane Central Register of Controlled Trials (CENTRAL)
	Cochrane Database of Systematic Reviews (CDSR)
	• Embase
	MEDLINE
	Epistemonikos
	CINAHLInternational HTA Database
	Health Technology Assessment (HTA) database (last updated October 2016).
	Searches will be restricted by:
	English language only
	human studies only.

Field	Content
	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	Folic acid supplementation
Population	Women in the preconception stage trying to become pregnant or women during the first 12 weeks of a single or multiple pregnancy
Intervention	 High-dose folic acid supplementation (≥5 mg daily) In combination or not with other vitamins and minerals
Comparator	 Low-dose folic acid supplementation (<1 mg daily) Medium-dose folic acid supplementation (≥1 to <5 mg daily) Placebo No intervention In combination or not with other vitamins and minerals
Types of study to be included	Include published full-text papers: • systematic reviews of RCTs • parallel RCTs • if insufficient parallel RCTs*: • observational studies: - systematic reviews of observational studies - prospective cohort studies - retrospective cohort studies

Field	Content
	 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 cofounding 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	Population: • women with a BMI ≥25kg/m². Setting: • countries other than high income countries (as defined by the OECD). If any study or systematic review includes <1/3 of women with the above characteristics 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, diabetes in pregnancy, antenatal care, intrapartum care, pregnancy and complex social factors or obesity prevention).
Primary outcomes (critical outcomes)	 Neural tube defects (NTDs) in the baby Birthweight Hypertensive disorders of pregnancy (pre-eclampsia and gestational hypertension).
Secondary outcomes (important outcomes)	 Red blood cell (RBC) (or serum if RBC is not reported) folate concentrations in the mother Neurodevelopmental outcomes (dichotomous outcome, not continuous outcomes such as mean change in score):

Field	Content
	 severe (score of >2SD below normal on validated assessment scales, or Bayley assessment scale of mental development index [MDI] or psychomotor developmental index [PDI]) moderate (score of 1-2 SD below normal on validated assessment scales, or Bayley assessment scale MDI or PDI 70-84). congenital heart defects midline facial defects such as cleft lip or cleft palate. If there is a composite outcome that includes any of the outcomes above, the composite outcome will be reported and given priority over individually reported outcomes
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.

Field	Content
	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 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 prespecified 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:
	 NTDs, hypertensive disorders of pregnancy, neurodevelopmental outcomes, congenital heart defects, cleft palate: 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	The following factors will be of interest in any meta-regression or subgroup analyses:
	single versus multi-fetal pregnancies
	women with a previous history of a pregnancy affected by a neural tube defect versus not
	women taking antiseizure medications versus not
	women taking antifolate medications versus not

Field	Content
	women with diabetes (type 1 and type 2) versus not
	women who started folic acid pre-conception versus not (specify timing if reported).
	If the evidence identifies additional groups which have not been listed here, these groups will be reported in the evidence.
	Evidence will be sub-grouped by the following only in the event that there is significant heterogeneity in outcomes:
	deprived socioeconomic group
	• age <40 versus ≥40
	 women and parents with disabilities, including learning disabilities and other physical and mental health conditions
	women going through assisted conception
	religion and cultural considerations
	• ethnicity:
	○ White/White British
	∘ Asian/Asian British
	o Black/African/Caribbean/Black British
	Mixed/Multiple ethnic groups
	o other ethnic group.
	women who smoke versus not.
	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.

Field	Content		
Type and method of review		tic tic ve	·y)
Language	English		
Country	England		
Anticipated or actual start date	14/07/2023		
Anticipated completion date	22/11/2023	_	
age of review at time of this submission	Review stage	Started	Completed
	Preliminary searches		\boxtimes
	Piloting of the study selection process		
	Formal screening of search results against eligibility criteria		
	Data extraction		\boxtimes
	Risk of bias (quality) assessment		

Field	Content				
	Data analysis		\boxtimes		
Named contact	5a. Named contact				
	National Institute for Health and Care Excellence (NICE)				
	5b. Named contact e-mail				
	mandcnutrition@r				
	5c. Organisational affiliation of the review				
	National Institute t				
Review team members				Excellence (NICE):	
	senior Systema	atic Reviewe	r		
	systematic Rev	viewer.			
Funding sources/sponsor	This systematic re (NICE).	eview is being	g completed by th	e 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=346779				
Dissemination plans	NICE may use a r standard approac		rent methods to ra	aise awareness of the guideline. These include	

Field	Content		
	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	Folic acid, pregnancy, overweight, obesity		
Details of existing review of same topic by same authors	Not applicable		
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			

CDSR: Cochrane Database of Systematic Reviews; CENTRAL: Cochrane Central Register of Controlled Trials; DARE: Database of Abstracts of Reviews of Effects; EPPI: Evidence for Policy & Practice Information; GRADE: Grading of Recommendations Assessment, Development and Evaluation; HTA: Health Technology Assessment; m: metre; MID: minimally important difference; mg: milligrams; NGA: National Guideline Alliance; NHS: National health service; NICE: National Institute for Health and Care Excellence; OECD: Organization for Economic Cooperation and Development; RCT: randomised controlled trial; RoB: risk of bias; SD: standard deviation

Appendix B Literature search strategies

Literature search strategies for review question: Which groups of women should be advised to take high-dose folic acid supplements before and during the first 12 weeks of pregnancy?

Effectiveness searches

Database: Medline

Date	of last search: 04/12/2023
#	Searches
1	exp Pregnancy/ or Pregnant Women/ or Prenatal Care/
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	Preconception Care/
5	(periconcept* or peri concept* or preconcept* or pre concept* or prepregnan* or pre pregnan*).ti,ab,kf.
6	((before or plan* or intend* or intention* or wish* or desir* or want* or prior or prepar* or try* or becom* or get* or start*) adj3 (baby or babies or conceiving or pregnan* or conception* or conceive*)).ti,ab,kf.
7	(start* adj2 family).ti,ab.
8	or/4-7
9	3 or 8
10	Folic Acid/ or Folic Acid Deficiency/
11	(folic acid* or folate* or folacin or vitamin b9 or vitamin b 9 or vitamin m or pteroylglutamic acid* or folvite).ti,ab,kf.
12	or/10-11
13	9 and 12
14	letter/
15	editorial/
16	news/
17	exp historical article/
18	Anecdotes as Topic/
19	comment/
20	case report/
21	(letter or comment*).ti.
22	or/14-21
23	randomized controlled trial/ or random*.ti,ab.
24	22 not 23
25	animals/ not humans/
26	exp Animals, Laboratory/
27	exp Animal Experimentation/
28	exp Models, Animal/
29	exp Rodentia/
30	(rat or rats or mouse or mice or rodent*).ti.
31	or/24-30
32	13 not 31
33	limit 32 to English language
34	randomized controlled trial.pt.
35	controlled clinical trial.pt.
36	pragmatic clinical trial.pt.
37	randomi#ed.ab.
38	placebo.ab.
39	drug therapy.fs.
40	randomly.ab.
41	trial.ab.
42	groups.ab.

Searches

43	or/34-42
44	Meta-Analysis/
45	Meta-Analysis as Topic/
46	(meta analy* or metanaly* or metaanaly*).ti,ab.
47	((systematic* or evidence*) adj2 (review* or overview*)).ti,ab.
48	(reference list* or bibliograph* or hand search* or manual search* or relevant journals).ab.
49	(search strategy or search criteria or systematic search or study selection or data extraction).ab.
50	(search* adj4 literature).ab.
51	(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.
52	cochrane.jw.
53	or/44-52
54	33 and (43 or 53)
55	Observational Studies as Topic/
56	Observational Study/
57	Epidemiologic Studies/
58	exp Case-Control Studies/
59	exp Cohort Studies/
60	Cross-Sectional Studies/
61	Controlled Before-After Studies/
62	Historically Controlled Study/
63	Interrupted Time Series Analysis/
64	Comparative Study.pt.
65	case control\$.tw.
66	case series.tw.
67	(cohort adj (study or studies)).tw.
68	cohort analy\$.tw.
69	(follow up adj (study or studies)).tw.
70	(observational adj (study or studies)).tw.
71	longitudinal.tw.
72	prospective.tw.
73	retrospective.tw.
74	cross sectional.tw.
75	or/55-74
76	33 and 75
77	76 not 54
78	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 djibouti/ or dominica/ or dominican 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-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 malta/ 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 serbia/ or serral leone/ or sergal/ or seychelles/ or singapore/ or somalia/ or south africa/ or south sudan/ or su tonga/ or tonga/ or surianame/ or syria/ or turkme

79 "organisation for economic co-operation and development"/

indies/ or yemen/ or zambia/ or zimbabwe/

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

tajikistan/ or tanzania/ or thailand/ or timor-leste/ or togo/ or tonga/ or "trinidad and tobago"/ or turkmenistan/ or uganda/ or ukraine/ or united arab emirates/ or uruguay/ or uzbekistan/ or vanuatu/ or venezuela/ or vietnam/ or west

#	Searches
	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/
81	european union/
82	developed countries/
83	or/79-82
84	78 not 83
85	54 not 84
86	77 not 84

Database: 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	prepregnancy care/
5	(periconcept* or peri concept* or preconcept* or pre concept* or prepregnan* or pre pregnan*).ti,ab,kf.
6	((before or plan* or intend* or intention* or wish* or desir* or want* or prior or prepar* or try* or becom* or get* or start*) adj3 (baby or babies or conceiving or pregnan* or conception* or conceive*)).ti,ab,kf.
7	(start* adj2 family).ti,ab.
8	or/4-7
9	3 or 8
10	folic acid/ or folic acid deficiency/
11	(folic acid* or folate* or folacin or vitamin b9 or vitamin b 9 or vitamin m or pteroylglutamic acid* or folvite).ti,ab,kf.
12	or/10-11
13	9 and 12
14	letter.pt. or letter/
15	note.pt.
16	editorial.pt.
17	case report/ or case study/
18	(letter or comment*).ti.
19	or/14-18
20	randomized controlled trial/ or random*.ti,ab.
21	19 not 20
22	animal/ not human/
23	nonhuman/
24	exp Animal Experiment/
25	exp Experimental Animal/
26	animal model/
27	exp Rodent/
28	(rat or rats or mouse or mice or rodent*).ti.
29	or/21-28
30	13 not 29
31	(conference abstract* or conference review or conference paper or conference proceeding).db,pt,su.
32	30 not 31
33	limit 32 to English language
34	random*.ti,ab.
35	factorial*.ti,ab.
36	(crossover* or cross over*).ti,ab.
37	((doubl* or singl*) adj blind*).ti,ab.
38	(assign* or allocat* or volunteer* or placebo*).ti,ab.
39	crossover procedure/

#	Searches
40	single blind procedure/
41	randomized controlled trial/
42	double blind procedure/
43	or/34-42
44	systematic review/
45	meta-analysis/
46	(meta analy* or metanaly* or metaanaly*).ti,ab.
47	((systematic or evidence) adj2 (review* or overview*)).ti,ab.
48	(reference list* or bibliograph* or hand search* or manual search* or relevant journals).ab.
49	(search strategy or search criteria or systematic search or study selection or data extraction).ab.
50	(search* adj4 literature).ab.
51	(medline or pubmed or cochrane or embase or psychlit or psychinfo or psychinfo or cinahl or science citation index or bids or cancerlit).ab.
52	((pool* or combined) adj2 (data or trials or studies or results)).ab.
53	cochrane.jw.
54	or/44-53
55	33 and (43 or 54)
56	Clinical study/
57	Case control study/
58	Family study/
59	Longitudinal study/
60	Retrospective study/
61	comparative study/
62	Prospective study/
63	Randomized controlled trials/
64	62 not 63
65	Cohort analysis/
66	cohort analy\$.tw.
67	(Cohort adj (study or studies)).tw.
68	(Case control\$ adj (study or studies)).tw.
69	(follow up adj (study or studies)).tw.
70	(observational adj (study or studies)).tw.
71	(epidemiologic\$ adj (study or studies)).tw.
72	(cross sectional adj (study or studies)).tw.
73	case series.tw.
74	prospective.tw.
75	retrospective.tw.
76	or/56-61,64-75
77	33 and 76
78	77 not 55
79	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 bahrain/ 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 bulgaria/ 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 senegal/ or exp serbia/ or seychelles/ or sierra leone/ or singapore/ or "sao tome and principe"/ or solomon islands/ or exp 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 taiwan/ or tajikistan/ or tanzania/ or thailand/ or timor-leste/ or togo/ or tonga/ or "trinidad and tobago"/ or tunisia/ or

#	Searches
	turkmenistan/ or tuvalu/ or uganda/ or exp ukraine/ or exp united arab emirates/ or uruguay/ or exp uzbekistan/ or vanuatu/ or venezuela/ or viet nam/ or western sahara/ or yemen/ or zambia/ or zimbabwe/
80	exp "organisation for economic co-operation and development"/
81	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/
82	european union/
83	developed country/
84	or/80-83
85	79 not 84
86	55 not 85
87	78 not 85

Database: Cochrane Database of Systematic Reviews Issue 12 of 12, December 2023 & Cochrane Central Register of Controlled Trials Issue 12 of 12, December 2023

Date of last search: 04/12/2023

#	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}
#6	MeSH descriptor: [Preconception Care] this term only
#7	(periconcept* or peri NEXT concept* or preconcept* or pre NEXT concept* or prepregnan* or pre NEXT pregnan*):ti,ab,kw
#8	((before or plan* or intend* or intention* or wish* or desir* or want* or prior or prepar* or try* or becom* or get* or start*) near/3 (baby or babies or conceiving or pregnan* or conception* or conceive*)):ti,ab,kw
#9	(start* NEAR/2 family):ti,ab
#10	{OR #6-#9}
#11	#5 OR #10
#12	MeSH descriptor: [Folic Acid] this term only
#13	MeSH descriptor: [Folic Acid Deficiency] this term only
#14	(folic acid* or folate* or folacin or vitamin b9 or "vitamin b 9" or "vitamin m" or pteroylglutamic NEXT acid* or folvite):ti,ab,kw
#15	{OR #12-#14}
#16	#11 AND #15
#17	conference:pt or (clinicaltrials or trialsearch):so
#18	#16 NOT #17

Database: CINAHL (Cumulative Index to Nursing and Allied Health Literature)

#	Searches	
S1	(MH "Pregnancy+")	
S2	62 (MH "Expectant Mothers")	
S3	(MH "Prenatal Care")	
S4	TI ((antenatal* or ante natal* or gestation* or maternal* or mother* or pregnan* or prenatal* or pre natal*)) OR AB ((antenatal* or ante natal* or gestation* or maternal* or mother* or pregnan* or prenatal* or pre natal*))	
S5	5 S1 OR S2 OR S3 OR S4	
S6	(MH "Prepregnancy Care")	
S7	TI ((periconcept* or peri concept* or preconcept* or pre concept* or prepregnan* or pre pregnan*)) OR AB ((periconcept* or peri concept* or preconcept* or pre concept* or prepregnan* or pre pregnan*))	

#	Searches	
S8	TI (((before or plan* or intend* or intention* or wish* or desir* or want* or prior or prepar* or try* or becom* or get* or start*) N3 (baby or babies or conceiving or pregnan* or conception* or conceive*))) OR AB (((before or plan* or intend* or intention* or wish* or desir* or want* or prior or prepar* or try* or becom* or get* or start*) N3 (baby or babies or conceiving or pregnan* or conception* or conceive*)))	
S9	TI (start* N2 family) OR AB (start* N2 family)	
S10	S10 S6 OR S7 OR S8 OR S9	
S11	S5 OR S10	
S12	(MH "Folic Acid") OR (MH "Folic Acid Deficiency")	
S13 TI ((folic acid* or folate* or folacin or vitamin b9 or vitamin b 9 or vitamin m or pteroylglutamic acid* or AB ((folic acid* or folate* or folacin or vitamin b9 or vitamin b 9 or vitamin m or pteroylglutamic acid*		
S14	S12 OR S13	
S15	S11 AND S14	
S16	S11 AND S14	
S17	7 S11 AND S14	

Database: Epistemonikos

Date of last search: 04/12/2023

#	Searches
1	title:((pregnan* OR antenatal OR prenatal OR periconcept* OR preconcept* OR prepregnan*)) or abstract:((pregnan* OR antenatal OR prenatal OR periconcept* OR preconcept* OR prepregnan*))
2	title:((folic acid* OR folate* OR folacin OR vitamin b9 OR pteroylglutamic acid* OR folvite)) OR abstract:((folic acid* OR folacin OR vitamin b9 OR pteroylglutamic acid* OR folvite)
3	#1 and #2 [Filters: classification=systematic-review, cochrane=missing, protocol=no]

Economic searches

Database: Medline

#	Searches	
1	exp Pregnancy/ or Pregnant Women/ or Prenatal Care/	
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	Preconception Care/	
5	(periconcept* or peri concept* or preconcept* or pre concept* or prepregnan* or pre pregnan*).ti,ab,kf.	
6	((before or plan* or intend* or intention* or wish* or desir* or want* or prior or prepar* or try* or becom* or get* or start*) adj3 (baby or babies or conceiving or pregnan* or conception* or conceive*)).ti,ab,kf.	
7	(start* adj2 family).ti,ab.	
8	or/4-7	
9	3 or 8	
10	Folic Acid/ or Folic Acid Deficiency/	
11	(folic acid* or folate* or folacin or vitamin b9 or vitamin b 9 or vitamin m or pteroylglutamic acid* or folvite).ti,ab,kf.	
12	or/10-11	
13	9 and 12	
14	letter/	
15	editorial/	
16	news/	
17	exp historical article/	
18	Anecdotes as Topic/	
19	comment/	
20	case reports/	
21	(letter or comment*).ti.	
22	or/14-21	
23	randomized controlled trial/ or random*.ti,ab.	

ш	Consider	
#	Searches	
24	22 not 23	
25	animals/ not humans/	
26	exp Animals, Laboratory/	
27	exp Animal Experimentation/	
28	exp Models, Animal/	
29	exp Rodentia/	
30	(rat or rats or mouse or mice or rodent*).ti.	
31	or/24-30	
32	13 not 31	
33	limit 32 to English language	
34	Economics/	
35	Value of life/	
36	exp "Costs and Cost Analysis"/	
37	exp Economics, Hospital/	
38	exp Economics, Medical/	
39	exp Resource Allocation/	
40	Economics, Nursing/	
41	Economics, Pharmaceutical/	
42	exp "Fees and Charges"/	
43	exp Budgets/	
44	budget*.ti,ab.	
45	cost*.ti,ab.	
46	(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	(value adj2 (money or monetary)).ti,ab.	
50	resourc* allocat*.ti,ab.	
51	(fund or funds or funding* or funded).ti,ab.	
52	(ration or rations or rationing* or rationed).ti,ab.	
53	ec.fs.	
54	or/34-53	
55	exp models, economic/	
56	*Models, Theoretical/	
57	*Models, Organizational/	
58	markov chains/	
59	monte carlo method/	
60	exp Decision Theory/	
61	(markov* or monte carlo).ti,ab.	
62	econom* model*.ti,ab.	
63	(decision* adj2 (tree* or analy* or model*)).ti,ab.	
64	or/55-63	
65	quality-adjusted life years/	
66	sickness impact profile/	
67	(quality adj2 (wellbeing or well being)).ti,ab.	
68	sickness impact profile.ti,ab.	
69	disability adjusted life.ti,ab.	
70	(qal* or qtime* or qwb* or daly*).ti,ab.	
71	(euroqol* or eq5d* or eq 5*).ti,ab.	
72	(qol* or hql* or hqol* or h qol* or hrqol* or hr qol*).ti,ab.	
73	(health utility* or utility score* or disutilit* or utility value*).ti,ab.	
74	(hui or hui1 or hui2 or hui3).ti,ab.	
75	(health* year* equivalent* or hye or hyes).ti,ab.	
76	discrete choice*.ti,ab.	

#	Searches	
77	rosser.ti,ab.	
78	(willingness to pay or time tradeoff or time trade off or tto or standard gamble*).ti,ab.	
79	(sf36* or sf 36* or short form 36* or shortform 36* or shortform36*).ti,ab.	
80	(sf20 or sf 20 or short form 20 or shortform 20 or shortform20).ti,ab.	
81	(sf12* or sf 12* or short form 12* or shortform 12* or shortform12*).ti,ab.	
82	(sf8* or sf 8* or short form 8* or shortform 8* or shortform8*).ti,ab.	
83	(sf6* or sf 6* or short form 6* or shortform 6* or shortform6*).ti,ab.	
84	or/65-83	
85	33 and (54 or 64 or 84)	

Database: Embase

	01 IdSt SedICI1. 04/12/2023		
#	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	prepregnancy care/		
5	(periconcept* or peri concept* or preconcept* or pre concept* or prepregnan* or pre pregnan*).ti,ab,kf.		
6	((before or plan* or intend* or intention* or wish* or desir* or want* or prior or prepar* or try* or becom* or get* or start*) adj3 (baby or babies or conceiving or pregnan* or conception* or conceive*)).ti,ab,kf.		
7	(start* adj2 family).ti,ab.		
8	or/4-7		
9	3 or 8		
10	folic acid/ or folic acid deficiency/		
11	(folic acid* or folate* or folacin or vitamin b9 or vitamin b 9 or vitamin m or pteroylglutamic acid* or folvite).ti,ab,kf.		
12	or/10-11		
13	9 and 12		
14	letter.pt. or letter/		
15	note.pt.		
16	editorial.pt.		
17	case report/ or case study/		
18	(letter or comment*).ti.		
19	or/14-18		
20	randomized controlled trial/ or random*.ti,ab.		
21	19 not 20		
22	animal/ not human/		
23	nonhuman/		
24	exp Animal Experiment/		
25	exp Experimental Animal/		
26	animal model/		
27	exp Rodent/		
28	(rat or rats or mouse or mice or rodent*).ti.		
29	or/21-28		
30	13 not 29		
31	(conference abstract* or conference review or conference paper or conference proceeding).db,pt,su.		
32	30 not 31		
33	limit 32 to English language		
34	health economics/		
35	exp economic evaluation/		
36	exp health care cost/		
37	exp fee/		
38	budget/		
39	funding/		
	-		

#	Searches		
40	resource allocation/		
41	budget*.ti,ab.		
42	cost*.ti,ab.		
43	(economic* or pharmaco?economic*).ti,ab.		
44			
45	(price* or pricing*).ti,ab. (financ* or fee or fees or expenditure* or saving*).ti,ab.		
46	(value adj2 (money or monetary)).ti,ab.		
47	resourc* allocat*.ti,ab.		
48	(fund or funds or funding* or funded).ti,ab.		
49	(ration or rations or rationing* or rationed).ti,ab.		
50	or/34-49		
51	statistical model/		
52			
	exp economic aspect/ 51 and 52		
53 54	*theoretical model/		
55	*nonbiological model/		
56	stochastic model/		
57	decision theory/		
58	decision tree/		
59	monte carlo method/		
60	(markov* or monte carlo).ti,ab.		
61	econom* model*.ti,ab.		
62	(decision* adj2 (tree* or analy* or model*)).ti,ab.		
63	or/53-62		
64	quality adjusted life year/		
65	"quality of life index"/		
66	short form 12/ or short form 20/ or short form 36/ or short form 8/		
67	sickness impact profile/		
68	(quality adj2 (wellbeing or well being)).ti,ab.		
69	sickness impact profile.ti,ab.		
70	disability adjusted life.ti,ab.		
71	(qal* or qtime* or qwb* or daly*).ti,ab.		
72	(qal* or qtime* or qwb* or daly*).ti,ab.		
73	(qol* or hql* or hqol* or h qol* or hrqol*).ti,ab.		
74	(health utility* or utility score* or disutilit* or utility value*).ti,ab.		
75	(hui or hui1 or hui2 or hui3).ti,ab.		
76	(health* year* equivalent* or hye or hyes).ti,ab.		
77	discrete choice*.ti,ab.		
78	rosser.ti,ab.		
79	(willingness to pay or time tradeoff or time trade off or tto or standard gamble*).ti,ab.		
80	(sf36* or sf 36* or short form 36* or shortform 36* or shortform36*).ti,ab.		
81	(sf20 or sf 20 or short form 20 or shortform 20 or shortform20).ti,ab.		
82	(sf12* or sf 12* or short form 12* or shortform 12* or shortform12*).ti,ab.		
83	(sf8* or sf 8* or short form 8* or shortform 8* or shortform8*).ti,ab.		
84	(sf6* or sf 6* or short form 6* or shortform 6* or shortform6*).ti,ab.		
85	or/64-84		
86	33 and (50 or 63 or 85)		

Database: INAHTA HTA

#	Searches
1	"Pregnancy"[mhe]

#	Searches	
2	"Pregnant Women"[mh]	
3 "Prenatal Care"[mh]		
4	((antenatal* or ante natal* or gestation* or maternal* or mother* or pregnan* or prenatal* or pre natal*))[Title] OR ((antenatal* or ante natal* or gestation* or maternal* or mother* or pregnan* or prenatal* or pre natal*))[abs]	
5	#4 OR #3 OR #2 OR #1	
6	"Preconception Care"[mh]	
7	((periconcept* or peri concept* or preconcept* or pre concept* or prepregnan* or pre pregnan*))[Title] OR ((periconcept* or peri concept* or preconcept* or preconcept* or prepregnan*))[abs]	
8	(((before or plan* or intend* or intention* or wish* or desir* or want* or prior or prepar* or try* or becom* or get* or start*) AND (baby or babies or conceiving or pregnan* or conception* or conceive*)))[Title] OR (((before or plan* or intend* or intention* or wish* or desir* or want* or prior or prepar* or try* or becom* or get* or start*) AND (baby or babies or conceiving or pregnan* or conception* or conceive*)))[abs]	
9	((start* AND family))[Title] OR ((start* AND family))[abs]	
10	#9 OR #8 OR #7 OR #6	
11	#10 OR #5	
12	"Folic Acid"[mh]	
13	"Folic Acid Deficiency"[mh]	
14	((folic acid* or folate* or folacin or "vitamin b9" or "vitamin b 9" or "vitamin m" or "pteroylglutamic acid*" or folvite))[Title] OR ((folic acid* or folate* or folacin or "vitamin b9" or "vitamin b 9" or "vitamin m" or "pteroylglutamic acid*" or folvite))[abs]	
15	#14 OR #13 OR #12	
16	#15 AND #11 Limit by English language	

Database: CRD HTA (last updated October 2016)

Date of last search: 20/07/2022

#	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 gestation* or maternal* or mother* 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 preconception care IN HTA	
7	(((periconcept* or peri concept* or preconcept* or pre concept* or prepregnan* or pre pregnan*))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
8	((((before or plan* or intend* or intention* or wish* or desir* or want* or prior or prepar* or try* or becom* or get* or start*) adj3 (baby or babies or conceiving or pregnan* or conception* or conceive*)))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
9	(((start* adj2 family))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
10	#6 OR #7 OR #8 OR #9	
11	#5 OR #10	
12	MeSH DESCRIPTOR folic acid IN HTA	
13	MeSH DESCRIPTOR folic acid deficiency IN HTA	
14	(((folic acid* or folate* or folacin or vitamin b9 or vitamin b 9 or vitamin m or pteroylglutamic acid* or folvite))) and (Project record:ZDT OR Full publication record:ZDT) IN HTA	
15	#12 OR #13 OR #14	
16	#11 AND #15	

Appendix C Effectiveness evidence study selection

Study selection for review question: Which groups of women should be advised to take high-dose folic acid supplements before and during the first 12 weeks of pregnancy?

Records identified through database searching Additional records identified through other sources = 12081 Records removed as duplicates n = 4920 Total records imported = 12081 Records screened in 1st sift Records excluded n = 6995 Screening on title and abstract n = 7161 Records excluded
n = 163
- 5: Conference abstract
- 2: Study population did not supplement diet with folic acid
- 17: Ineligible study outcomes
- 5: No information on folic acid dose available
- 1: Article unavailable
- 8: Ineligible population
- 1: Study has no comparator group
- 2: Trial protocol
- 1: Study data presented graphically 2: Trial protocol
1: Study data presented graphically and cannot be used in analysis
4: Not in English language
1: Unclear how many participants randomised to each arm
7: Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not compare different doses of folic acid.
1: Systematic review. Included Records screened in 2nd sift unrerent doses of folic acid.

1: Systematic review. Included
studies checked for eligibility. No
studies identified for inclusion
because none of the papers were from
high income countries (as defined by
the OECD). Screening on full text n = 166 the OECD).
22 : Ineligible study design
11 : Study conducted in general
population and dose already
established for this group established for this group

5: Duplicate

7: Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not have eligible outcomes. outcomes.

10 : Systematic review. Included studies checked for eligibility. No studies identified for inclusion or have already been included through main search. - 16 : Ineligible country - 37 : Ineligible intervention Records included in review

Figure 1: Effectiveness evidence study selection flow chart

Appendix D Evidence tables

Evidence tables for review question: Which groups of women should be advised to take high-dose folic acid supplements before and during the first 12 weeks of pregnancy?

Table 5: Evidence tables

Laurence, 1981

Bibliographic Reference

Laurence, K M; James, N; Miller, M H; Tennant, G B; Campbell, H; Double-blind randomised controlled trial of folate treatment before conception to prevent recurrence of neural-tube defects.; British medical journal (Clinical research ed.); 1981; vol. 282 (no. 6275); 1509-11

Study details

Country/ies where study was carried out	UK
Study type	Randomised controlled trial (RCT)
Study dates	Not reported
Inclusion criteria	 History of birth with NTDs Under 35 years
Exclusion criteria	Not reported
Patient characteristics	Not reported
Intervention(s)/control	Intervention: 4mg folic acid* Placebo *2x2mg tablets per day Taken from the time contraception stopped until birth
Duration of follow-up	At birth

Sources of funding	Not industry funded
Sample size	N=111
	Intervention: n=60 Placebo: n=51
Other information	Subgroup in analysis: risk of preeclampsia or history of births with NTDs

mg: milligrams; n: number of participants; NTDs: neural tube defects

Study arms

4mg folic acid/day (n = 60)

Placebo (n = 51)

Outcomes

Outcome	4mg folic acid/day, n = 60	Placebo, n = 51
Neural tube defects measured at birth Number for intervention is combined for compliers and non-compliers No of events	n = 2; % = 1.2	n = 4; % = 2.04
Red blood cell folate concentration at birth Mean for intervention arm is combined for compliers and non-compliers Mean (SD) (µg/l red blood cells)	609 (218)	278 (16)

mg: milligrams; n: number of participants; SD: standard deviation

Critical appraisal - Cochrane Risk of Bias tool v2.0 for RCTs

Section	Question	Answer
Domain 1: Bias arising from the randomisation process	Risk of bias judgement for the randomisation process	Low

Section	Question	Answer
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
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
Domain 3. Bias due to missing outcome data	Risk-of-bias judgement for missing outcome data	Low
Domain 4. Bias in measurement of the outcome	Risk-of-bias judgement for measurement of the outcome	Low
Domain 5. Bias in selection of the reported result	Risk-of-bias judgement for selection of the reported result	Some concerns (No study protocol reported)
Overall bias and Directness	Risk of bias judgement	Some concerns (The study is judged to have some concerns of bias in one domain)
Overall bias and Directness	Overall Directness	Directly applicable (Study selected pregnant women with history of births with NTDs)
Overall bias and Directness	Risk of bias variation across outcomes	Low risk of bias in variation across outcomes

NTDs: neural tube defects; RCT: randomised controlled trial.

MRC, 1991

Bibliographic Reference

MRC; Prevention of neural tube defects: results of the Medical Research Council Vitamin Study. MRC Vitamin Study Research Group.; Lancet (London, England); 1991; vol. 338 (no. 8760); 131-7

Study details

Country/ies where study was carried out	UK, Hungary, Australia, Israel, Canada, Russia, France						
Study type	Randomised controlled trial (RCT)						
Study dates	Not reported						
Inclusion criteria	 Those with a history of a pregnancy with NTDs Planning another pregnancy Not taking vitamin supplements 						
Exclusion criteria	Women with epilepsy						
Patient characteristics	Age in years, mean (SD: NR)						
	Group A (4mg folic acid): 27						
	Group B (4mg folic acid + multivitamin): 27.4						
	Group C (placebo): 26.8						
	Maternal BMI – NR						
	Livebirths in previous pregnancies, mean						
	Group A (4mg folic acid): 0.88						
	Group B (4mg folic acid + multivitamin): 0.94						
	Group C (placebo): 0.91						
	Mean number of previous NTD pregnancies						
	Group A (4mg folic acid): 1.03						

	Group B (4mg folic acid + multivitamin): 1.04
	Group C (placebo): 1.01
Intervention(s)/control	Group A: 4mg folic acid
	Group B: 4mg folic acid + multivitamin
	Group C: Placebo (contains dried ferrous sulphate (120mg) and di-calcium phosphate (240mg)
	*1 capsule daily from date of randomisation until 12 weeks gestational age
Duration of follow-up	12 weeks of pregnancy
Sources of funding	Not industry funded
Sample size	N=893
	Group A: n=298
	Group B: n=295
	Group C: n=300
Other information	Multivitamin composition: vitamin A 4000 IU, D 400 IU, BI 1-5 mg, B2 1.5 mg, B6 10 mg, C 40 mg, and nicotinamide 15 mg.
	Subgroup in analysis: risk of preeclampsia or history of births with NTDs

BMI: body mass index; IU: international units; mg: milligrams; NR: not reported; NTDs: neural tube defects; SD: standard deviation

Study arms

Folic acid 4mg/day (n = 298)

Folic acid 4mg + multivitamins/day (n = 295)

Placebo (n = 300)

Outcomes

Outcome	Folic acid 4mg/day, n = 298	Folic acid 4mg + multivitamins/day, n = 295	Placebo, n = 300
Neural tube defects measured at 12 weeks of pregnancy	n = 2; % = 0.67	n = 4; % = 1.4	n = 13; % = 4.3
No of events			

mg: milligrams; n: number of participants

Critical appraisal - Cochrane Risk of Bias tool v2.0 for RCTs

Section	Question	Answer
Domain 1: Bias arising from the randomisation process	Risk of bias judgement for the randomisation process	Low
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
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
Domain 3. Bias due to missing outcome data	Risk-of-bias judgement for missing outcome data	High (Outcome data were available for 64% of randomised participants. There is no evidence that the result was not biased by missing outcome data. Missingness in the outcome could depend on its true value, however this is not likely.)
Domain 4. Bias in measurement of the outcome	Risk-of-bias judgement for measurement of the outcome	Low
Domain 5. Bias in selection of the reported result	Risk-of-bias judgement for selection of the reported result	Some concerns (No study protocol reported)

Section	Question	Answer
Overall bias and Directness	Risk of bias judgement	High (The study is judged to raise some concerns in one domain and high risk of bias in one domain)
Overall bias and Directness	Overall Directness	Directly applicable (Study selected pregnant women with history of births with NTDs)
Overall bias and Directness	Risk of bias variation across outcomes	Low risk of bias in variation across outcomes

NTDs: neural tube defects; RCT: randomised controlled trial.

Wen, 2018

Bibliographic Reference

Wen, Shi Wu; White, Ruth Rennicks; Rybak, Natalie; Gaudet, Laura M; Robson, Stephen; Hague, William; Simms-Stewart, Donnette; Carroli, Guillermo; Smith, Graeme; Fraser, William D; Wells, George; Davidge, Sandra T; Kingdom, John; Coyle, Doug; Fergusson, Dean; Corsi, Daniel J; Champagne, Josee; Sabri, Elham; Ramsay, Tim; Mol, Ben Willem J; Oudijk, Martijn A; Walker, Mark C; FACT Collaborating, Group; Effect of high dose folic acid supplementation in pregnancy on pre-eclampsia (FACT): double blind, phase III, randomised controlled, international, multicentre trial.; BMJ (Clinical research ed.); 2018; vol. 362; k3478

Study details

Country/ies where study was carried out	Canada, Argentina, Australia, Jamaica, UK
Study type	Randomised controlled trial (RCT)
Study dates	April 2011 to November 2015
Inclusion criteria	 8-16 gestational weeks Confirmed viable fetus Risk factor for pre-eclampsia (pre-existing hypertension, prepregnancy diabetes (type 1 or 2), twin pregnancy, history of pre-eclampsia during pregnancy, BMI >=35 kg/m²)
Exclusion criteria	Known fetal anomaly or fetal deathHistory of maternal medical complications

Epilepsy Cancer Current use of folic acid antagonists Substance misuse Hypersensitivity to folic acid Multiple pregnancy • Previous participation in current trial History or presence of a disease or condition that would preclude use of high dose folic acid **Patient** Maternal age in years, n (%) characteristics Intervention: <20- 10 (0.8) 20-29-439 (35.8) 30-34-411 (33.5) ≥35- 367 (29.9) Age in years, mean (SD): 31 (5.4) Placebo: <20- 10 (0.8) 20-29-447 (36.2) 30-34-441 (35.7) ≥35- 338 (27.3)

```
Age in years, mean (SD): 31 (5.4)
Pregnancy BMI, mean (SD)
Intervention: 34 (8.6)
Control: 34 (13)
BMI ≥35, n (%)
Intervention: 606 (49.4)
Control: 656 (53.1)
Parity, n (%)
Intervention:
0 - 413 (33.7)
1 - 498 (40.6)
≥2 - 316 (25.7)
Control:
0 - 420 (34.0)
1 - 499 (40.4)
≥2 - 317 (25.6)
History of pre-eclampsia, n (%)
```

Intervention: 308 (25.1)

Control: 303 (24.5)

Chronic hypertension, n (%)

Intervention: 203 (16.5)

Control: 241 (19.5)

Type 1 diabetes, n (%)

Intervention: 84 (6.8)

Control: 72 (5.8)

Type 2 diabetes, n (%)

Intervention: 98 (8.0)

Control: 84 (6.8)

Twin pregnancy, n (%)

Intervention: 233 (19.0)

Control: 229 (18.5)

Maternal ethnicity, n (%)

Intervention:

Native/Aboriginal 35 (2.85)

White 970 (79.0) Black 93 (7.6) Asian 45 (3.7) Latino/Hispanic 31 (2.5) Indian/South Asian 45 (3.7) Declined to answer 8 (0.65) Control: Native/Aboriginal 24 (1.94) White 987 (79.8) Black 107 (8.7) Asian 50 (4.05) Latino/Hispanic 22 (1.8) Indian/South Asian 36 (2.9) Declined to answer 10 (0.8) Smoking during pregnancy, n (%) Intervention: Yes 98 (8.0)

	No 1046 (85.2)
	Quit during pregnancy 83 (6.8)
	Control:
	Yes 95 (7.7)
	No 1035 (83.7)
	Quit during pregnancy 106 (8.6)
Intervention(s)/control	Intervention: 4mg folic acid (4x1mg tablets daily)
	Control: Placebo
	Taken from 8 to 16 weeks gestational age until birth.
	*Participants continued taking vitamins or folic acid supplements (up to 1.1 mg of folic acid).
Duration of follow-up	42 days post-partum
Sources of funding	Not industry funded
Sample size	N=2301
	Intervention: n=1144
	Control: n=1157
Other information	Participants who took folic acid supplementation additional to intervention:
	Folic acid supplementation (includes multivitamin containing folic acid) at randomisation
	Intervention: 080 (80.6)
	Intervention: 989 (80.6)
	Control: 1016 (82.2)

Supplementation of high dose (≥4.0 mg/d folic acid) at randomisation

Intervention: 346 (28.2)

Control: 335 (27.1)

Subgroup in analysis: risk of preeclampsia or history of births with NTDs

BMI: body mass index; d: day; kg: kilogram; m: metre; mg: milligrams; n: number of participants; SD: standard deviation

Study arms

4mg folic acid (n = 1144)

Placebo (n = 1157)

Outcomes

Outcome	4mg folic acid, n = 1144	Placebo, n = 1157
Preeclampsia measured at 42 days post-partum	n = 24; % = 2.1	n = 16; % = 1.4
No of events		

n: number of participants; mg: milligrams

Critical appraisal - Cochrane Risk of Bias tool v2.0 for RCTs

Section	Question	Answer
Domain 1: Bias arising from the randomisation process	Risk of bias judgement for the randomisation process	Low
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
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

Section	Question	Answer
Domain 3. Bias due to missing outcome data	Risk-of-bias judgement for missing outcome data	Low
Domain 4. Bias in measurement of the outcome	Risk-of-bias judgement for measurement of the outcome	Some concerns (Participants could continue taking prenatal vitamins or low dose folic acid supplements (up to 1.1 mg of folic acid).)
Domain 5. Bias in selection of the reported result	Risk-of-bias judgement for selection of the reported result	Low
Overall bias and Directness	Risk of bias judgement	Some concerns (The study is judged to raise some concerns one domain)
Overall bias and Directness	Overall Directness	Directly applicable (Study selected pregnant women at high risk of developing pre-eclampsia)
Overall bias and Directness	Risk of bias variation across outcomes	Low risk of bias in variation across outcomes

mg: milligrams; RCT: randomised controlled trial.

Appendix E Forest plots

Forest plots for review question: Which groups of women should be advised to take high-dose folic acid supplements before and during the first 12 weeks of pregnancy?

This section includes forest plots only for outcomes that are meta-analysed. Outcomes from single studies are not presented here; the quality assessment for such outcomes is provided in the GRADE profiles in appendix F.

Comparison between folic acid supplementation 4mg/day and placebo in women with a history of births with neural tube defects (NTDs)

Figure 2: Neural tube defects (follow-up: at 12 weeks of pregnancy and birth)

	4mg folic aci	d/day	Place	bo		Risk Ratio	Risk Ratio	
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% CI	M-H, Fixed, 95% CI	
Laurence 1981	2	60	4	51	25.0%	0.42 [0.08, 2.23]		
MRC 1991	2	298	13	300	75.0%	0.15 [0.04, 0.68]		
Total (95% CI)		358		351	100.0%	0.22 [0.07, 0.66]	-	
Total events	4		17					
Heterogeneity: Chi² = 0.82, df = 1 (P = 0.37); l² = 0%				0.01 0.1 10 1				
Test for overall effect: $Z = 2.71$ (P = 0.007)					Favours 4mg FA/day Favours placebo	00		

Cl: confidence interval; df: degrees of freedom; FA: folic acid; mg: milligrams; MRC: Medical Research Council; NTDs: neural tube defects

Appendix F GRADE tables

GRADE tables for review question: Which groups of women should be advised to take high-dose folic acid supplements before and during the first 12 weeks of pregnancy?

Table 6: Evidence profile for comparison between folic acid supplementation 4mg/day and placebo in women with a history of births with neural tube defects (NTDs)

	Quality assessment						No of patients		Effect		Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	4mg folic acid/day	Placebo	Relative (95% CI)	Absolute	Quanty	Importance
Neural tube	defects (follow	w-up: at 12	weeks of pregnar	ncy and birth)				•			-	
21	randomised trials		no serious inconsistency		no serious imprecision	none	4/358 (1.1%)	17/351 (4.9%)	RR 0.22 (0.07 to 0.65)	38 fewer per 1000 (from 17 fewer to 46 fewer)	IMP. BENEFIT	CRITICAL
Red blood c	ell folate cond	entration	(µg/l red blood cel	lls) (Better ind	dicated by highe	r values) (follow-uj	o: at birth)					
1 Laurence 1981	randomised trials		no serious inconsistency		no serious imprecision	none	60	51	-	MD 331 higher (275.66 to 386.34 higher)	IMP. BENEFIT	IMPORTANT

CI: confidence interval; MD: mean difference; mg: milligrams; RR: risk ratio

Table 7: Evidence profile for comparison between folic acid supplementation 4mg/day + multivitamin and placebo for women with a history of births with neural tube defects (NTDs)

			o minimi modinar									
	Quality assessment					No of patients Effect			Effect	Quality	Importance	
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	4mg folic acid/day + multivitamin	Placebo	Relative (95% CI)	Absolute	Quality	importance
Neural tub	oe defects (fol	low-up: at	12 weeks of preg	nancy)								
1 MRC 1991	randomised trials		no serious inconsistency	serious ²	very serious²	none	4/295 (1.4%)	13/300 (4.3%)	RR 0.31 (0.10 to 0.95)	30 fewer per 1000 (from 2 fewer to 39 fewer)	VERY LOW	CRITICAL
			,						,	,	IMP. BENEFIT	

CI: confidence interval; mg: milligrams; RR: risk ratio

¹ See corresponding forest plot in appendix E for studies contributing to this outcome

² Serious risk of bias in the evidence contributing to the outcome as per RoB 2

³ Intervention is indirect as folic acid 4 mg/day dose was used in the study. Did not include of high dose folic acid supplements (≥5 mg daily) as per protocol.

¹ Serious risk of bias in the evidence contributing to the outcome as per RoB 2

² Intervention is indirect as folic acid 4 mg/day dose was used in the study. Did not include of high dose folic acid supplements (≥5 mg daily) as per protocol.

³ Statistical significance was used to assess clinical importance. Imprecision was based on total number of events. <150 events.

Table 8: Evidence profile for comparison between folic acid supplementation 4mg/day and placebo in women at high risk of preeclampsia

	procolar											
	Quality assessment					No of patients Effect			Ovelity I			
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	4mg folic acid/day + multivitamin	Placebo	Relative (95% CI)	Absolute	Quality	Importance
Severe pre	e-eclampsia (f	ollow-up: a	at 42 days postpar	tum)								
1 Wen	randomised	serious ¹	no serious	very serious ²	very	none	24/1144	16/1157	RR 1.52 (0.81	7 more per 1000 (from 3	VERY	CRITICAL
2018	trials		inconsistency		serious ³		(2.1%)	(1.4%)	to 2.84)	fewer to 25 more)	LOW	
											NO IMP. DIFF.	

CI: confidence interval; mg: milligrams; RR: risk ratio

¹ Serious risk of bias in the evidence contributing to the outcome as per RoB 2 ² Population and intervention are indirect due to >33% of population has a BMI>=35kg/m². Intervention is indirect as folic acid 4 mg/day dose was used in the study. Did not include of high dose folic acid supplements (≥5 mg daily) as per protocol.

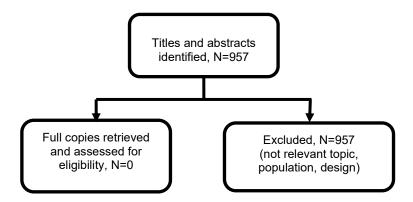
³ Statistical significance was used to assess clinical importance. Imprecision was based on total number of events. <150 events.

Appendix G Economic evidence study selection

Study selection for review question: Which groups of women should be advised to take high-dose folic acid supplements before and during the first 12 weeks of pregnancy?

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

Figure 3. Flow diagram of selection process for economic evaluations



Appendix H Economic evidence tables

Economic evidence tables for review question: Which groups of women should be advised to take high-dose folic acid supplements before and during the first 12 weeks of pregnancy?

No evidence was identified which was applicable to this review question.

Appendix I Economic model

Economic model for review question: Which groups of women should be advised to take high-dose folic acid supplements before and during the first 12 weeks of pregnancy?

No economic analysis was conducted for this review question.

Appendix J Excluded studies

Excluded studies for review question: Which groups of women should be advised to take high-dose folic acid supplements before and during the first 12 weeks of pregnancy?

Excluded effectiveness studies

Table 9: Excluded studies and reasons for their exclusion

lable 9: Excluded studies and reasons for	titeli exclusion				
Study	Code [Reason]				
Ala, Syed H; Husain, Samia; Husain, Saba (2021) Reasons for presenting to antenatal care clinics in a sample of Pakistani women and their knowledge of WHO antenatal care package. European journal of midwifery 5: 43	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in Pakistan.				
Ali Khan, R.E., Ahmad, T.I., Noreen, S. et al. (2020) Factors associated with the quality of Antenatal Care (ANC) services use in Bahawalpur. Journal of the Pakistan Medical Association 70(12b): 2457-2459	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in Pakistan.				
Alvestad, Silje, Nilsen Husebye, Elisabeth Synnove, Christensen, Jakob et al. (2022) Folic Acid and Risk of Preterm Birth, Preeclampsia and Fetal Growth Restriction Among Women With Epilepsy: A Prospective Cohort Study. Neurology	- Ineligible intervention Does not compare different doses of folic acid				
Avnon, Tomer, Anbar, Ronit, Lavie, Inbar et al. (2020) Does vegan diet influence umbilical cord vitamin B12, folate, and ferritin levels?. Archives of gynecology and obstetrics 301(6): 1417-1422	- Study population did not supplement diet with folic acid Participants were analysed according to diet (for example, vegan diet, vegetarian diet, and so on)				
Bakker, Rachel, Timmermans, Sarah, Steegers, Eric A P et al. (2011) Folic acid supplements modify the adverse effects of maternal smoking on fetal growth and neonatal complications. The Journal of nutrition 141(12): 2172-9	- Ineligible intervention Folic acid dosage of 0.4–0.5 mg/d. Prospective cohort study.				
Baumslag, N; Edelstein, T; Metz, J (1970) Reduction of incidence of prematurity by folic acid supplementation in pregnancy. British medical journal 1(5687): 16-7	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in South Africa.				
Baxter, JA.B., Wasan, Y., Soofi, S.B. et al. (2018) Effect of life skills building education and micronutrient supplements provided from preconception versus the standard of care on low birth weight births among adolescent and	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in Pakistan.				

Study	Code [Reason]
young Pakistani women (15-24 years): A prospective, population-based cluster-randomized trial. Reproductive Health 15(1): 104	
Bermeo-Ovalle, A. (2020) Think Beyond Malformations: The Case for Periconceptional Folate in Women With Epilepsy. Currents 20(4): 202-204	- Ineligible study design Narrative commentary
Betts, T and Fox, C (1999) Proactive preconception counselling for women with epilepsyis it effective?. Seizure 8(6): 322-7	- Ineligible study design Retrospective study
Bezerra Espinola, Maria Salome; Bilotta, Gabriele; Aragona, Cesare (2021) Positive effect of a new supplementation of vitamin D3 with myo-inositol, folic acid and melatonin on IVF outcomes: a prospective randomized and controlled pilot study. Gynecological endocrinology: the official journal of the International Society of Gynecological Endocrinology 37(3): 251-254	- Ineligible intervention Study assesses effects of vitamin D3 on IVF outcomes
Bhutta, Zulfiqar A, Imdad, Aamer, Ramakrishnan, Usha et al. (2012) Is it time to replace iron folate supplements in pregnancy with multiple micronutrients?. Paediatric and perinatal epidemiology 26suppl1: 27-35	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Systematic review included studies from low-middle income countries only.
Bortolus, Renata, Blom, Fenneke, Filippini, Francesca et al. (2014) Prevention of congenital malformations and other adverse pregnancy outcomes with 4.0 mg of folic acid: community-based randomized clinical trial in Italy and the Netherlands. BMC pregnancy and childbirth 14: 166	- Trial protocol
Bortolus, Renata, Filippini, Francesca, Cipriani, Sonia et al. (2021) Efficacy of 4.0 mg versus 0.4 mg Folic Acid Supplementation on the Reproductive Outcomes: A Randomized Controlled Trial. Nutrients 13(12)	- Study conducted in general population and dose already established for this group Study investigated 4 milligrams of folic acid
Bower, C (1994) Epilepsy in pregnancy: neural tube defects and folate. The Medical journal of Australia 160(2): 56-7	- Ineligible study design Narrative. Published letter/comment on another article
Bramswig, Susanne, Prinz-Langenohl, Reinhild, Lamers, Yvonne et al. (2009) Supplementation	- Study conducted in general population and dose already established for this group

Study	Code [Reason]
with a multivitamin containing 800 microg of folic acid shortens the time to reach the preventive red blood cell folate concentration in healthy women. International journal for vitamin and nutrition research. Internationale Zeitschrift fur Vitamin- und Ernahrungsforschung. Journal international de vitaminologie et de nutrition 79(2): 61-70	Study investigated 800 micrograms of folic acid
Brieger, Katharine K, Bakulski, Kelly M, Pearce, Celeste L et al. (2022) The Association of Prenatal Vitamins and Folic Acid Supplement Intake with Odds of Autism Spectrum Disorder in a High-Risk Sibling Cohort, the Early Autism Risk Longitudinal Investigation (EARLI). Journal of autism and developmental disorders 52(6): 2801-2811	- Ineligible study design Prospective cohort study
Brough, L, Rees, G A, Crawford, M A et al. (2009) Social and ethnic differences in folic acid use preconception and during early pregnancy in the UK: effect on maternal folate status. Journal of human nutrition and dietetics: the official journal of the British Dietetic Association 22(2): 100-7	- Ineligible study outcomes Study reports on number of women taking folic acid supplements
Brough, Louise, Rees, Gail A, Crawford, Michael A et al. (2010) Effect of multiple-micronutrient supplementation on maternal nutrient status, infant birth weight and gestational age at birth in a low-income, multi-ethnic population. The British journal of nutrition 104(3): 437-45	- Ineligible comparison Compared low dose folic supplement (400µg folic acid in multivitamin "Pregnacare") vs placebo.
Bukowski, Radek, Malone, Fergal D, Porter, Flint T et al. (2009) Preconceptional folate supplementation and the risk of spontaneous preterm birth: a cohort study. PLoS medicine 6(5): e1000061	- Ineligible study design Prospective cohort study
Busby, A., Abramsky, L., Dolk, H. et al. (2005) Preventing neural tube defects in Europe: Population based study. British Medical Journal 330(7491): 574-575	- Ineligible study design Narrative commentary
Butler, E B (1968) Effect of iron and folic acid on red cell and plasma volume in pregnancy. The Journal of obstetrics and gynaecology of the British Commonwealth 75(5): 497-510	- Ineligible study outcomes Study assessed haemoglobin levels during pregnancy
Butler, EB (1968) The effect of iron and folic acid on red cell and plasma volume in pregnancy. Journal of obstetrics and	- Duplicate

Study	Code [Reason]
gynaecology of the british commonwealth 75: 497-510	
Carretti, N, Eremita, G A, Porcelli, B et al. (1994) Pattern of vitamin B12 and folic acid during pregnancy. Gynecologic and obstetric investigation 38(2): 78-81	- Ineligible intervention Study investigated folate/vitamin B12 concentrations and gestational age/levels of haemoglobin
Castren, O, Levanto, A, Rauramo, L et al. (1968) Preventive iron and folic acid therapy in pregnancy. Annales chirurgiae et gynaecologiae Fenniae 57(3): 382-6	- Ineligible intervention Study does not compare different doses of folic acid
Castro, Kamila, Klein, Luciana da Silveira, Baronio, Diego et al. (2016) Folic acid and autism: What do we know?. Nutritional neuroscience 19(7): 310-7	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not compare different doses of folic acid.
Catena, Andres, Munoz-Machicao, J Angela, Torres-Espinola, Francisco J et al. (2016) Folate and long-chain polyunsaturated fatty acid supplementation during pregnancy has long-term effects on the attention system of 8.5-y-old offspring: a randomized controlled trial. The American journal of clinical nutrition 103(1): 115-27	- Ineligible intervention Study assesses effect of fish oil and folate supplementation on children's neurocognitive development
Catov, J.M., Bodnar, L.M., Olsen, J. et al. (2011) Periconceptional multivitamin use and risk of preterm or small-for-gestational-age Births in the Danish National birth cohort. American Journal of Clinical Nutrition 94(3): 906-912	- Ineligible intervention Intervention compares timing of folic acid rather than dose
Catov, Janet M, Bodnar, Lisa M, Ness, Roberta B et al. (2007) Association of periconceptional multivitamin use and risk of preterm or small-forgestational-age births. American journal of epidemiology 166(3): 296-303	- Ineligible intervention No information on dose reported
Charles, Deborah H M, Ness, Andy R, Campbell, Doris et al. (2005) Folic acid supplements in pregnancy and birth outcome: re-analysis of a large randomised controlled trial and update of Cochrane review. Paediatric and perinatal epidemiology 19(2): 112-24	- Study conducted in general population and dose already established for this group Study investigated 200 micrograms and 5 milligrams of folic acid
Cheng, Zhengpei, Gu, Rui, Lian, Zenglin et al. (2022) Evaluation of the association between maternal folic acid supplementation and the risk of congenital heart disease: a systematic review and meta-analysis. Nutrition journal 21(1): 20	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not compare different doses of folic acid.

Study	Code [Reason]
Chiaffarino, F, Ascone, GB, Bortolus, R et al. (2010) [Effects of folic acid supplementation on pregnancy outcomes: a review of randomized clinical trials]. Minerva ginecologica 62(4): 293-301	- Not in English language
Chisholm, M (1966) A controlled clinical trial of prophylactic folic acid and iron in pregnancy. Journal of obstetrics and gynaecology of the British Commonwealth 73: 191-196	- Ineligible intervention Study assesses relationship between iron/folic acid and anaemia
Christian, P, Shahid, F, Rizvi, A et al. (2009) Treatment response to standard of care for severe anemia in pregnant women and effect of multivitamins and enhanced anthelminthics. American journal of clinical nutrition 89(3): 853-861	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in Pakistan.
Cochrane, K.M., Elango, R., Devlin, A.M. et al. (2023) Supplementation with (6 S)-5-methyltetrahydrofolic acid appears as effective as folic acid in maintaining maternal folate status while reducing unmetabolized folic acid in maternal plasma: A randomized trial of pregnant women in Canada. British Journal of Nutrition	- Study conducted in general population and dose already established for this group Study investigated 0.6 mg folic acid
Cochrane, Kelsey M, Elango, Rajavel, Devlin, Angela M et al. (2023) Human milk unmetabolized folic acid is increased following supplementation with synthetic folic acid as compared to (6S)-5-methyltetrahydrofolic acid. Scientific reports 13(1): 11298	- Study conducted in general population and dose already established for this group Study assessed 0.6 mg folic acid
Cochrane, Kelsey M, Elango, Rajavel, Devlin, Angela M et al. (2023) Supplementation with (6S)-5-methyltetrahydrofolic acid appears as effective as folic acid in maintaining maternal folate status while reducing unmetabolised folic acid in maternal plasma: a randomised trial of pregnant women in Canada. The British journal of nutrition: 1-11	- Duplicate
Czeizel, Andrew E (2009) Periconceptional folic acid and multivitamin supplementation for the prevention of neural tube defects and other congenital abnormalities. Birth defects research. Part A, Clinical and molecular teratology 85(4): 260-8	- Study conducted in general population and dose already established for this group Study investigated 0.8mg folic acid
Czeizel, Andrew E. (2011) Periconceptional Folic Acid-Containing Multivitamin Supplementation for the Prevention of Neural	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion

Study	Code [Reason]
<u>Tube Defects and Cardiovascular</u> <u>Malformations.</u> Annals of Nutrition & Metabolism 59(1): 38-40	because studies do not compare different doses of folic acid.
Davis, Deborah J (1997) Periconceptional folic acid and the prevention of neural tube defects: Some food for thought for Canadian paediatricians. Paediatrics & child health 2(2): 114-119	- Ineligible study design Questionnaire survey
Dawson, D W (1966) Microdoses of folic acid in pregnancy. The Journal of obstetrics and gynaecology of the British Commonwealth 73(1): 44-8	- Ineligible study design Prospective study
Dawson, LE; Pham, B; Hunter, AGW (2001) Low rate of adequate folic acid supplementation in well-educated women of high socioeconomic status attending a genetics clinic. Canadian Medical Association Journal (CMAJ) 164(8): 1149-1150	- Ineligible study design Research letter/comment
De Ocampo, Maria P G, Araneta, Maria Rosario G, Macera, Caroline A et al. (2018) Folic acid supplement use and the risk of gestational hypertension and preeclampsia. Women and birth: journal of the Australian College of Midwives 31(2): e77-e83	- Ineligible intervention Study does not compare two doses of folic acid
De-Regil, Luz Maria, Fernandez-Gaxiola, Ana C, Dowswell, Therese et al. (2010) Effects and safety of periconceptional folate supplementation for preventing birth defects. The Cochrane database of systematic reviews: cd007950	- Duplicate
De-Regil, Luz Maria, Pena-Rosas, Juan Pablo, Fernandez-Gaxiola, Ana C et al. (2015) Effects and safety of periconceptional oral folate supplementation for preventing birth defects. The Cochrane database of systematic reviews: cd007950	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion or have already been included through main search.
Desrosiers, T.A., Siega-Riz, A.M., Mosley, B.S. et al. (2018) Low carbohydrate diets may increase risk of neural tube defects. Birth Defects Research 110(11): 901-909	- Ineligible intervention Study investigated association between carbohydrate intake and neural tube defects
Dobo, M and Czeizel, A E (1998) Long-term somatic and mental development of children after periconceptional multivitamin	- Ineligible study outcomes Study reported long term effects of multivitamin supplementation in childhood

Study	Code [Reason]
supplementation. European journal of pediatrics 157(9): 719-23	
Dong, Jing, Yin, Lin-Liang, Deng, Xue-Dong et al. (2023) Initiation and duration of folic acid supplementation in preventing congenital malformations. BMC medicine 21(1): 292	- Ineligible study design Prospective cohort study.
Dwyer, Erin Rose, Filion, Kristian B, MacFarlane, Amanda J et al. (2022) Who should consume high-dose folic acid supplements before and during early pregnancy for the prevention of neural tube defects? BMJ (Clinical research ed.) 377: e067728	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion or have already been included through main search.
Edelstein, T, Stevens, K, Baumslag, N et al. (1968) Folic acid and vitamin B12 supplementation during pregnancy in a population subsisting on a sub-optimal diet. Journal of obstetrics and gynaecology of the British Commonwealth 75: 133-137	- Ineligible intervention Study does not compare different doses of folic acid (both arms recieved 5mg folic acid)
Einarson, A; Parshuram, C; Koren, G (2000) Periconceptional use of folic acid to reduce the rates of neural tube defects: is it working?. Reproductive toxicology (Elmsford, N.Y.) 14(4): 291-292	- Ineligible study design Narrative commentary
Fernandez-Roig, Silvia, Cavalle-Busquets, Pere, Fernandez-Ballart, Joan D et al. (2013) Low folate status enhances pregnancy changes in plasma betaine and dimethylglycine concentrations and the association between betaine and homocysteine. The American journal of clinical nutrition 97(6): 1252-9	- Ineligible study design Prospective cohort study
Fleming, A F, Martin, J D, Hahnel, R et al. (1974) Effects of iron and folic acid antenatal supplements on maternal haematology and fetal wellbeing. The Medical journal of Australia 2(12): 429-36	- Ineligible intervention Study investigates the association between iron/folic acid on anaemia
Furness, Denise, Fenech, Michael, Dekker, Gustaaf et al. (2013) Folate, vitamin B12, vitamin B6 and homocysteine: impact on pregnancy outcome. Maternal & child nutrition 9(2): 155-66	- Ineligible study design Prospective cohort study
Gao, Yunfei, Sheng, Chao, Xie, Ri-Hua et al. (2016) New Perspective on Impact of Folic Acid Supplementation during Pregnancy on Neurodevelopment/Autism in the Offspring	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not have eligible outcomes.

Study	Code [Reason]
Children - A Systematic Review. PloS one 11(11): e0165626	
Gildestad, Trude, Bjorge, Tone, Haaland, Oystein A et al. (2020) Maternal use of folic acid and multivitamin supplements and infant risk of birth defects in Norway, 1999-2013. The British journal of nutrition 124(3): 316-329	- Ineligible study design Prospective cohort study
Gildestad, Trude, Bjorge, Tone, Vollset, Stein Emil et al. (2015) Folic acid supplements and risk for oral clefts in the newborn: a population-based study. The British journal of nutrition 114(9): 1456-63	- No information on folic acid dose available
Giles, P F; Harcourt, A G; Whiteside, M G (1971) The effect of prescribing folic acid during pregnancy on birth-weight and duration of pregnancy. A double-blind trial. The Medical journal of Australia 2(1): 17-21	- Study conducted in general population and dose already established for this group Study investigated 5mg folic acid
Giunta, G., Cardea, C., Matarazzo, M.G. et al. (2018) Is calcium levofolinate pentahydrate more effective than folic acid in young healthy women before conception?. Clinical and Experimental Obstetrics and Gynecology 45(5): 687-691	- Ineligible intervention Study assesses whether levofolinic acid is better than folic acid to increase serum folate levels
Gogou, Maria and Kolios, George (2020) Nutritional Supplements During Gestation and Autism Spectrum Disorder: What Do We Really Know and How Far Have We Gone?. Journal of the American College of Nutrition 39(3): 261-271	- Ineligible intervention Study assesses impact of dietary supplements during pregnancy on the risk of autism spectrum disorder in the child
Gomes, Filomena, Askari, Sufia, Black, Robert E et al. (2023) Antenatal multiple micronutrient supplements versus iron-folic acid supplements and birth outcomes: Analysis by gestational age assessment method. Maternal & child nutrition 19(3): e13509	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because none of the papers were from high income countries (as defined by the OECD).
Gunabalasingam, Sowmiya, De Almeida Lima Slizys, Daniele, Quotah, Ola et al. (2023) Micronutrient supplementation interventions in preconception and pregnant women at increased risk of developing pre-eclampsia: a systematic review and meta-analysis. European journal of clinical nutrition 77(7): 710-730	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion or have already been included through main search. Relevant Wen 2018 paper included via primary sift.
Guo, BQ., Li, HB., Zhai, DS. et al. (2019) Maternal multivitamin supplementation is associated with a reduced risk of autism	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion

Study	Code [Reason]
spectrum disorder in children: a systematic review and meta-analysis. Nutrition Research 65: 4-16	because studies do not compare different doses of folic acid.
Guo, Bao-Qiang, Li, Hong-Bin, Zhai, De-Sheng et al. (2019) Association of maternal prenatal folic acid intake with subsequent risk of autism spectrum disorder in children: A systematic review and meta-analysis. Progress in neuro-psychopharmacology & biological psychiatry 94: 109650	- Ineligible intervention Study assesses association between folic acid and autism spectrum disorder
Halicioglu, Oya, Sutcuoglu, Sumer, Koc, Feyza et al. (2012) Vitamin B12 and folate statuses are associated with diet in pregnant women, but not with anthropometric measurements in term newborns. The journal of maternal-fetal & neonatal medicine: the official journal of the European Association of Perinatal Medicine, the Federation of Asia and Oceania Perinatal Societies, the International Society of Perinatal Obstetricians 25(9): 1618-21	- Ineligible study design Cross-sectional study design
Hampel, K and Roetz, R (1974) Influence of a long-time substitution with a folate-iron combination in pregnancy on serum folate and serum iron and on hematological parameters. Geburtshilfe und Frauenheilkunde 34: 409-417	- Not in English language
Hashemi, Mohammad, Heshmat-Ghahdarijani, Kiyan, Zarean, Elahe et al. (2016) Evaluation of the effect of high-dose folic acid on endothelial dysfunction in pre-eclamptic patients: A randomized clinical trial. Journal of research in medical sciences: the official journal of Isfahan University of Medical Sciences 21: 114	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in Iran.
Hodgetts, V A, Morris, R K, Francis, A et al. (2015) Effectiveness of folic acid supplementation in pregnancy on reducing the risk of small-for-gestational age neonates: a population study, systematic review and metanalysis. BJOG: an international journal of obstetrics and gynaecology 122(4): 478-90	- Ineligible intervention Study assesses timing of folic acid, not dose
Holmes-Siedle, M., Dennis, J., Lindenbaum, R.H. et al. (1992) Long term effects of periconceptional multivitamin supplements for prevention of neural tube defects: A seven to 10 year follow up. Archives of Disease in Childhood 67(12): 1436-1441	- Ineligible study design Prospective cohort study

Study	Code [Reason]
Hossain, N, Kanani, FH, Ramzan, S et al. (2014) Obstetric and neonatal outcomes of maternal vitamin D supplementation: results of an open- label, randomized controlled trial of antenatal vitamin D supplementation in Pakistani women. Journal of clinical endocrinology and metabolism 99(7): 2448-2455	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in Pakistan.
Hunter, Patricia J, Muthiani, Yvonne, Nasanen-Gilmore, Pieta K et al. (2023) A modular systematic review of antenatal interventions to address undernutrition during pregnancy in the prevention of low birth weight. The American journal of clinical nutrition 117suppl2: 134-s147	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not compare different doses of folic acid.
Hursthouse, Nicola A, Gray, Andrew R, Miller, Jody C et al. (2011) Folate status of reproductive age women and neural tube defect risk: the effect of long-term folic acid supplementation at doses of 140 microg and 400 microg per day. Nutrients 3(1): 49-62	- Study conducted in general population and dose already established for this group Study investigated 140mg and 400ug of folic acid
Iglesias Vazquez, Lucia; Canals, Josefa; Arija, Victoria (2019) Review and meta-analysis found that prenatal folic acid was associated with a 58% reduction in autism but had no effect on mental and motor development. Acta paediatrica (Oslo, Norway: 1992) 108(4): 600-610	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not have eligible outcomes.
Jarvenpaa, J, Schwab, U, Lappalainen, T et al. (2008) Mineral water fortified with folic acid and vitamins B6, B12, D and calcium improves folate status and decreases plasma homocysteine concentration in pregnant women. 35th nordic congress of obstetrics and gynecology; 2006 may 23-25; goteburg, sweden: 55	- Conference abstract
Kancherla, Vijaya, Botto, Lorenzo D, Rowe, Laura A et al. (2022) Preventing birth defects, saving lives, and promoting health equity: an urgent call to action for universal mandatory food fortification with folic acid. The Lancet. Global health 10(7): e1053-e1057	- Ineligible study design Narrative summary
Kannan, S, Menotti, E, Scherer, HK et al. (2007) Folic acid and the prevention of neural tube defects: a survey of awareness among Latina women of childbearing age residing in southeast Michigan. Health Promotion Practice 8(1): 60-68	- Ineligible study design Survey
Kelly, Dervla; O'Dowd, Tom; Reulbach, Udo (2012) Use of folic acid supplements and risk of	- Ineligible study design

Study	Code [Reason]
cleft lip and palate in infants: a population-based cohort study. The British journal of general practice: the journal of the Royal College of General Practitioners 62(600): e466-72	Cross-sectional study design
Kennedy, D; Pastuszak, A; Koren, G (1997) Taking folic acid during pregnancy. Don't leave it too late. Canadian family physician Medecin de famille canadien 43: 2113-4	- Ineligible study design Information or leaflet for women in the pre- conception stage
Kerr, Stephen M, Parker, Samantha E, Mitchell, Allen A et al. (2017) Periconceptional maternal fever, folic acid intake, and the risk for neural tube defects. Annals of epidemiology 27(12): 777-782e1	- Ineligible study design Case-control study
Kinnunen, Tarja I, Sletner, Line, Sommer, Christine et al. (2017) Ethnic differences in folic acid supplement use in a population-based cohort of pregnant women in Norway. BMC pregnancy and childbirth 17(1): 143	- No information on folic acid dose available
Kirke, P N; Daly, L E; Elwood, J H (1992) A randomised trial of low dose folic acid to prevent neural tube defects. The Irish Vitamin Study Group. Archives of disease in childhood 67(12): 1442-6	- Ineligible comparison Compared low dose (0.36mg) folic acid supplement vs multivitamins with no folic acid.
Koren, G (1993) Preconceptional folate and neural tube defects: time for rethinking. Canadian journal of public health = Revue canadienne de sante publique 84(3): 207-8	- Ineligible study design Narrative commentary
Lassi, Zohra S, Salam, Rehana A, Haider, Batool A et al. (2013) Folic acid supplementation during pregnancy for maternal health and pregnancy outcomes. The Cochrane database of systematic reviews: cd006896	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion or have already been included through main search.
Laurence, K.M.; James, N.; Miller, M. (1982) A double randomised controlled trial for preconceptional folic acid supplementation for the prevention of recurrence of neural tube defects in high risk pregnancies. Journal of Medical Genetics 19(1): 66	- Conference abstract
Laurence, KM (1982) The role of maternal nutrition and folic acid therapy before conception to prevent neural tube defects. Monatsschrift fur Kinderheilkunde 130: 646	- Conference abstract

Study	Code [Reason]
Lira, P, Barrena, N, Foradori, A et al. (1989) Deficiency of folates in pregnancy: effect of supplementary folic acid. Sangre 34(1): 24-27	- Not in English language
Lira, P, Barrena, N, Foradori, A et al. (1989) Folate deficiency in pregnancy: effect of supplemental folate. Sangre 34: 24-27	- Not in English language
Liu, Cheng, Liu, Chongdong, Wang, Qiushi et al. (2018) Supplementation of folic acid in pregnancy and the risk of preeclampsia and gestational hypertension: a meta-analysis. Archives of gynecology and obstetrics 298(4): 697-704	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion or have already been included through main search.
Meador, Kimford J, Pennell, Page B, May, Ryan C et al. (2020) Effects of periconceptional folate on cognition in children of women with epilepsy: NEAD study. Neurology 94(7): e729-e740	- Ineligible study design Prospective cohort study
Millar, Wayne J (2004) Folic acid supplementation. Health reports 15(3): 49-52	- No information on folic acid dose available
Mills, J L and Raymond, E (1993) Effects of recent research on recommendations for periconceptional folate supplement use. Annals of the New York Academy of Sciences 678: 137-45	- Ineligible study design Cost-Benefit Analysis
Monoarfa, Y, Gumilar, E, Widasari, L et al. (2020) The effect of selenium and multiple micronutrient administration during periconception period on the level of malondialdehyde. Enfermeria clinica 30suppl4: 114-118	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in Indonesia.
Morrow, J I, Hunt, S J, Russell, A J et al. (2009) Folic acid use and major congenital malformations in offspring of women with epilepsy: a prospective study from the UK Epilepsy and Pregnancy Register. Journal of neurology, neurosurgery, and psychiatry 80(5): 506-11	- Ineligible intervention Unclear what dose of folic acid was given to women in intervention group (some had high dose of 5mg and some had 400mcg)
Muchowski, Karen and Paladine, Heather (2004) An ounce of prevention: the evidence supporting periconception health care. The Journal of family practice 53(2): 126-33	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion or have already been included through main search.

Study	Code [Reason]
Murphy, Malia S Q, Muldoon, Katherine A, Sheyholislami, Hauna et al. (2021) Impact of high-dose folic acid supplementation in pregnancy on biomarkers of folate status and 1-carbon metabolism: An ancillary study of the Folic Acid Clinical Trial (FACT). The American journal of clinical nutrition 113(5): 1361-1371	- Ineligible intervention Study assesses folate and 1-carbon metabolism biomarkers in pregnant women
Murphy, P A (1992) Periconceptional supplementation with folic acid. Does it prevent neural tube defects?. Journal of nurse-midwifery 37(1): 25-32	- Ineligible study design Literature review
Nilsen, Roy M, Vollset, Stein E, Gjessing, Hakon K et al. (2006) Patterns and predictors of folic acid supplement use among pregnant women: the Norwegian Mother and Child Cohort Study. The American journal of clinical nutrition 84(5): 1134-41	- Ineligible study design Prospective cohort study
Nkrumah, Isaac, North, Madelon, Kothe, Emily et al. (2020) The Relationship Between Pregnancy Intentions and Diet or Physical Activity Behaviors in the Preconception and Antenatal Periods: A Systematic Review and Meta-Analysis. Journal of midwifery & women's health 65(5): 660-680	- Ineligible intervention Study assesses the association between women's pregnancy intentions and diet or physical activity behaviors in the preconception and antenatal periods
O'Malley, Eimer G, Reynolds, Ciara M E, Cawley, Shona et al. (2018) Folate and vitamin B12 levels in early pregnancy and maternal obesity. European journal of obstetrics, gynecology, and reproductive biology 231: 80-84	- Ineligible population Study based on women with a BMI ≥25kg/m²
Obeid, Rima, Schon, Christiane, Wilhelm, Manfred et al. (2018) The effectiveness of daily supplementation with 400 or 800 microg/day folate in reaching protective red blood folate concentrations in non-pregnant women: a randomized trial. European journal of nutrition 57(5): 1771-1780	- Study conducted in general population and dose already established for this group Study investigated 400ug or 800ug folic acid
Oulhote, Youssef, Lanphear, Bruce, Braun, Joseph M. et al. (2020) Gestational Exposures to Phthalates and Folic Acid, and Autistic Traits in Canadian Children. Environmental Health Perspectives 128(2): 027004-1	- Ineligible intervention Study assesses the relationship between gestational phthalates and autistic traits in 3 to 4 year old children
Ozerol, Elif, Ozerol, Ibrahim, Gokdeniz, Remzi et al. (2004) Effect of smoking on serum concentrations of total homocysteine, folate,	- Ineligible intervention

Study	Code [Reason]
vitamin B12, and nitric oxide in pregnancy: a preliminary study. Fetal diagnosis and therapy 19(2): 145-8	Study assesses differences in the serum levels between smoking andd non-smoking pregnant women
Park, Hyesook, Kim, Young Ju, Ha, Eun Hee et al. (2004) The risk of folate and vitamin B(12) deficiencies associated with hyperhomocysteinemia among pregnant women. American journal of perinatology 21(8): 469-75	- Ineligible intervention Study compares folate and vitamin B12 levels in pregnant and nonpregnant women to evaluate the risk for hyperhomocysteinemia and for folate and vitamin B12 deficiencies during pregnancy
Partap, Uttara, Chowdhury, Ranadip, Taneja, Sunita et al. (2022) Preconception and periconception interventions to prevent low birth weight, small for gestational age and preterm birth: a systematic review and meta-analysis. BMJ global health 7(8)	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not have eligible outcomes.
Pereira, Abilio and Keating, Elisa (2023) Maternal folate and metabolic programming of the offspring: A systematic review of the literature. Reproductive toxicology (Elmsford, N.Y.) 120: 108439	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion or have already been included through main search.
Pritchard, J A; Scott, D E; Whalley, P J (1971) Maternal folate deficiency and pregnancy wastage. IV. Effects of folic acid supplements, anticonvulsants, and oral contraceptives. American journal of obstetrics and gynecology 109(3): 341-6	- Ineligible intervention Study assesses the role of folate supplements, pregnancy and oral contraceptives in megaloblastic anaemia
Ramakrishnan, Usha, Grant, Frederick Kobina, Goldenberg, Tamar et al. (2012) Effect of multiple micronutrient supplementation on pregnancy and infant outcomes: a systematic review. Paediatric and perinatal epidemiology 26suppl1: 153-67	- Ineligible intervention Systematic review does not compare different doses of folic acid
Ramakrishnan, Usha, Grant, Frederick Kobina, Imdad, Aamer et al. (2013) Effect of multiple micronutrient versus iron-folate supplementation during pregnancy on intrauterine growth. Nestle Nutrition Institute workshop series 74: 53-62	- No information on folic acid dose available
Ray, Joel G and Mamdani, Muhammad M (2002) Association between folic acid food fortification and hypertension or preeclampsia in pregnancy. Archives of internal medicine 162(15): 1776-7	- Ineligible study design Retrospective longitudinal study

Study	Code [Reason]
Ray, Joel G, Wyatt, Philip R, Vermeulen, Marian J et al. (2005) Greater maternal weight and the ongoing risk of neural tube defects after folic acid flour fortification. Obstetrics and gynecology 105(2): 261-5	- Ineligible study design Retrospective cohort study
Renard, Emeline, Leheup, Bruno, Gueant-Rodriguez, Rosa-Maria et al. (2020) Folinic acid improves the score of Autism in the EFFET placebo-controlled randomized trial. Biochimie 173: 57-61	- Ineligible population Study included children with autistic spectrum disorder
Rhoads, G G and Mills, J L (1986) Can vitamin supplements prevent neural tube defects? Current evidence and ongoing investigations. Clinical obstetrics and gynecology 29(3): 569-79	- Ineligible study design Literature review
Riaz, Musarrat, Shaikh, Fareeha, Fawwad, Asher et al. (2018) Maternal Nutrition during Early Pregnancy and Cardiometabolic Status of Neonates at Birth. Journal of diabetes research 2018: 7382946	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in Pakistan.
Rolschau, J; Date, J; Kristoffersen, K (1979) Folic acid supplement and intrauterine growth. Acta obstetricia et gynecologica Scandinavica 58(4): 343-6	- Ineligible study design Prospective cohort study
Rose, Elaine G, Murphy, Malia S Q, Erwin, Erica et al. (2021) Gestational Folate and Folic Acid Intake among Women in Canada at Higher Risk of Pre-Eclampsia. The Journal of nutrition 151(7): 1976-1982	- Ineligible study design Secondary analysis of randomised controlled trial
Rucklidge, Julia J, Eggleston, Matthew J F, Darling, Kathryn A et al. (2019) Can we predict treatment response in children with ADHD to a vitamin-mineral supplement? An investigation into pre-treatment nutrient serum levels, MTHFR status, clinical correlates and demographic variables. Progress in neuro-psychopharmacology & biological psychiatry 89: 181-192	- Ineligible population Study investigated children with ADHD
Saccone, Gabriele and Berghella, Vincenzo (2016) Folic acid supplementation in pregnancy to prevent preterm birth: a systematic review and meta-analysis of randomized controlled trials. European journal of obstetrics, gynecology, and reproductive biology 199: 76-81	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion or have already been included through main search.

Study	Code [Reason]
Sadeeqa, S., Aslam, F., Zafar, M. et al. (2019) Relationship of medications and pregnancy complications. Journal of Medical Sciences (Peshawar) 27(3): 213-219	- Ineligible study design Cross-sectional study design
Sayyah-Melli, Manizheh, Ghorbanihaghjo, Amir, Alizadeh, Mahasti et al. (2016) The Effect of High Dose Folic Acid throughout Pregnancy on Homocysteine (Hcy) Concentration and Pre-Eclampsia: A Randomized Clinical Trial. PloS one 11(5): e0154400	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in Iran.
Shaw, M T and Hoffbrand, A V (1970) The use and abuse of folic acid. The Practitioner 204(224): 795-804	- Ineligible study design Literature review
Shere, Mahvash, Nguyen, Patricia, Tam, Carolyn et al. (2015) Pregnancy-induced changes in the long-term pharmacokinetics of 1.1 mg vs. 5 mg folic acid: a randomized clinical trial. Journal of clinical pharmacology 55(2): 159-67	- Study data presented graphically and cannot be used in analysis
Shim, Sang-Min; Yun, Yeo-Ul; Kim, Yun Sook (2016) Folic acid alone or multivitamin containing folic acid intake during pregnancy and the risk of gestational hypertension and preeclampsia through meta-analyses. Obstetrics & gynecology science 59(2): 110-5	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion or have already been included through main search.
Shiralizadeh, Javad, Barmaki, Haleh, Haiaty, Sanya et al. (2017) The effects of high and low doses of folic acid on oxidation of protein levels during pregnancy: a randomized double-blind clinical trial. Hormone molecular biology and clinical investigation 33(3)	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in Dubai.
Siddiqui, M.; Khatri, I.A.; Ahmad, A. (2014) Maternal and fetal complications in pregnant epileptic women. Rawal Medical Journal 39(2): 174-177	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in Pakistan.
Stern, S.J., Matok, I., Kapur, B. et al. (2011) A comparison of folic acid pharmacokinetics in obese and nonobese women of childbearing age. Therapeutic Drug Monitoring 33(3): 336-340	- Ineligible population Study population is women with with a BMI ≥25kg/m²
Suzuki, T., Nishigori, T., Obara, T. et al. (2022) Maternal folic acid supplement use/dietary folate	- Ineligible study design

Study	Code [Reason]
intake from preconception to early pregnancy and neurodevelopment in 2-year-old offspring: the Japan Environment and Children's Study. British Journal of Nutrition 128(12): 2480-2489	Prospective cohort study.
Takacs, P and Rodriguez, L (2005) High folic acid levels and failure of single-dose methotrexate treatment in ectopic pregnancy. International journal of gynaecology and obstetrics: the official organ of the International Federation of Gynaecology and Obstetrics 89(3): 301-2	- Conference abstract
Tamura, Tsunenobu, Goldenberg, Robert L, Chapman, Victoria R et al. (2005) Folate status of mothers during pregnancy and mental and psychomotor development of their children at five years of age. Pediatrics 116(3): 703-8	- Ineligible study design Retrospetive cohort study
Tchernia, G, Blot, I, Rey, A et al. (1982) Maternal folate status, birthweight and gestational age. Developmental Pharmacology Therapeutics 4: 58-65	- Article unavailable
Ter Borg, Sovianne; Koopman, Nynke; Verkaik- Kloosterman, Janneke (2023) An Evaluation of Food and Nutrient Intake among Pregnant Women in The Netherlands: A Systematic Review. Nutrients 15(13)	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion or have already been included through main search.
Trigg, K H, Rendall, E J, Johnson, A et al. (1976) Folate supplements during pregnancy. The Journal of the Royal College of General Practitioners 26(164): 228-30	- Ineligible intervention Study does not compare different doses of folic acid. Study compares two doses of ferrous sulphate
Tuenter, Annelies, Bautista Nino, Paula K, Vitezova, Anna et al. (2019) Folate, vitamin B12, and homocysteine in smoking-exposed pregnant women: A systematic review. Maternal & child nutrition 15(1): e12675	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not compare different doses of folic acid.
Turck, Dominique, Bohn, Torsten, Castenmiller, Jacqueline et al. (2023) Scientific opinion on the tolerable upper intake level for folate. EFSA journal. European Food Safety Authority 21(11): e08353	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not have eligible outcomes.
Ulrich, Cornelia M and Potter, John D (2006) Folate supplementation: too much of a good thing?. Cancer epidemiology, biomarkers & prevention: a publication of the American	- Ineligible study design Narrative commentary

Study	Code [Reason]
Association for Cancer Research, cosponsored by the American Society of Preventive Oncology 15(2): 189-93	
Ulrich, M, Kristoffersen, K, Rolschau, J et al. (1999) The influence of folic acid supplement on the outcome of pregnancies in the county of Funen in Denmark. Part II. Congenital anomalies. A randomised study. European journal of obstetrics, gynecology, and reproductive biology 87(2): 111-4	- Unclear how many participants randomised to each arm
Ulrich, M, Kristoffersen, K, Rolschau, J et al. (1999) The influence of folic acid supplement on the outcome of pregnancies in the county of Funen in Denmark. Part III. Congenital anomalies. An observational study. European journal of obstetrics, gynecology, and reproductive biology 87(2): 115-4	- Ineligible study design Retrospective cohort study
US Preventive Services Task, Force, Bibbins- Domingo, Kirsten, Grossman, David C et al. (2017) Folic Acid Supplementation for the Prevention of Neural Tube Defects: US Preventive Services Task Force Recommendation Statement. JAMA 317(2): 183-189	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not have eligible outcomes.
van der Windt, Melissa, van der Kleij, Rianne Maria, Snoek, Katinka Marianne et al. (2020) Impact of a Blended Periconception Lifestyle Care Approach on Lifestyle Behaviors: Before- and-After Study. Journal of medical Internet research 22(9): e19378	- Ineligible intervention Study assesses the effectiveness of a periconceptional lifestyle intervention on the improvement of lifestyle components
Van Dyke, Don C, Stumbo, Phyllis J, Mary, J Berg et al. (2002) Folic acid and prevention of birth defects. Developmental Medicine & Child Neurology 44(6): 426-429	- Ineligible study design Literature review
van Gellekom, S.A., Lindauer-van der Werf, G., Hague, W.M. et al. (2008) Anaemia and haemolysis in pregnancy due to rapid folic acid and vitamin B12 depletion. Netherlands Journal of Medicine 66(5): 216-217	- Ineligible study design Case report
Virk, Jasveer, Liew, Zeyan, Olsen, Jorn et al. (2016) Preconceptional and prenatal supplementary folic acid and multivitamin intake and autism spectrum disorders. Autism: the international journal of research and practice 20(6): 710-8	- Ineligible study outcomes Study reported on austism spectrum disorders in children

Study	Code [Reason]
Virk, Jasveer, Liew, Zeyan, Olsen, Jorn et al. (2018) Pre-conceptual and prenatal supplementary folic acid and multivitamin intake, behavioral problems, and hyperkinetic disorders: A study based on the Danish National Birth Cohort (DNBC). Nutritional neuroscience 21(5): 352-360	- Ineligible study design Prospective cohort study
Viswanathan, M, Urrutia, RP, Hudson, KN et al. (2023) Folic Acid Supplementation to Prevent Neural Tube Defects: A Limited Systematic Review Update for the U.S. Preventive Services Task Force. U.S. Preventive Services Task Force Evidence Syntheses, formerly Systematic Evidence Reviews	- Duplicate
Viswanathan, Meera, Urrutia, Rachel Peragallo, Hudson, Kesha N et al. (2023) Folic Acid Supplementation to Prevent Neural Tube Defects: Updated Evidence Report and Systematic Review for the US Preventive Services Task Force. JAMA 330(5): 460-466	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not have eligible outcomes. 1 RCT was included in this SR (Bortolus 2021), which was screened out via primary sift.
Wald, N (1993) Folic acid and the prevention of neural tube defects. Annals of the New York Academy of Sciences 678: 112-29	- Ineligible study design Narrative comment/letter
Wald, N J and Bower, C (1994) Folic acid, pernicious anaemia, and prevention of neural tube defects. Lancet (London, England) 343(8893): 307	- Conference abstract
Wald, N J, Hackshaw, A, Stone, R et al. (1993) Serum alpha-fetoprotein and neural tube defects in the first trimester of pregnancy. MRC Vitamin Study Research Group. Prenatal diagnosis 13(11): 1047-50	- No information on folic acid dose available
Wald, N.J. (2004) Folic Acid and the Prevention of Neural-Tube Defects. New England Journal of Medicine 350(2): 101-103	- Ineligible study design Case-control study
Wen, S W, Champagne, J, Rennicks White, R et al. (2012) OS040. Effect of folic acid supplementation in pregnancy on preeclampsia-Folic acid clinical trial (FACT). Pregnancy hypertension 2(3): 198	- Trial protocol
Wen, S.W., White, R.R., Rybak, N. et al. (2019) Effect of high dose folic acid supplementation in pregnancy on pre-eclampsia (fact): Double	- Duplicate

Study	Code [Reason]
blind, Phase III, randomised controlled, international, multicentre trial. Obstetrical and Gynecological Survey 74(2): 68-70	
Wen, Shi Wu, Guo, Yanfang, Rodger, Marc et al. (2016) Folic Acid Supplementation in Pregnancy and the Risk of Pre-Eclampsia-A Cohort Study. PloS one 11(2): e0149818	- Ineligible study design Prospective cohort study
Wen, SW, White, RR, Rybak, N et al. (2019) Effect of high dose folic acid supplementation in pregnancy on pre-eclampsia (fact): double blind, Phase III, randomised controlled, international, multicentre trial. Obstetrical & gynecological survey 74(2): 68-70	- Duplicate
Willoughby, M L and Jewell, F G (1968) Folate status throughout pregnancy and in postpartum period. British medical journal 4(5627): 356-60	- Ineligible study design Prospective cohort study
Willoughby, M L and Jewell, F J (1966) Investigation of folic acid requirements in pregnancy. British medical journal 2(5529): 1568-71	- Ineligible study outcomes Study reports on incidence of megaloblastic anaemia of pregnancy
Wilson, R Douglas and O'Connor, Deborah L (2022) Guideline No. 427: Folic Acid and Multivitamin Supplementation for Prevention of Folic Acid-Sensitive Congenital Anomalies. Journal of obstetrics and gynaecology Canada: JOGC = Journal d'obstetrique et gynecologie du Canada: JOGC 44(6): 707-719e1	- Ineligible study design Guideline recommendations
Withanage, Nishadi N, Botfield, Jessica R, Srinivasan, Sonia et al. (2022) Effectiveness of preconception interventions in primary care: a systematic review. The British journal of general practice: the journal of the Royal College of General Practitioners 72(725): e865-e872	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not have eligible outcomes.
Yang, Yan; Cai, Zixin; Zhang, Jingjing (2021) Association between maternal folate status and gestational diabetes mellitus. Food science & nutrition 9(4): 2042-2052	- Ineligible study design Study reports on gestational diabetes mellitus
Yang, Yan; Cai, Zixin; Zhang, Jingjing (2021) The effect of prepregnancy body mass index on maternal micronutrient status: a meta-analysis. Scientific reports 11(1): 18100	- Ineligible intervention Study assesses the effect between prepregnancy body mass index and maternal micronutrient status

Study	Code [Reason]
Yerby, Mark S (2003) Clinical care of pregnant women with epilepsy: neural tube defects and folic acid supplementation. Epilepsia 44suppl3: 33-40	- Ineligible study design Literature review
Yi, Yunni, Lindemann, Marion, Colligs, Antje et al. (2011) Economic burden of neural tube defects and impact of prevention with folic acid: a literature review. European journal of pediatrics 170(11): 1391-400	- Ineligible study design Literature review
Yusuf, Korede K, Salihu, Hamisu M, Wilson, Ronee et al. (2019) Folic Acid Intake, Fetal Brain Growth, and Maternal Smoking in Pregnancy: A Randomized Controlled Trial. Current developments in nutrition 3(6): nzz025	- Ineligible study outcomes Study reports on prenatal fetal brain growth, measured by head circumference, brain weight, and brain-body weight ratio
Zheng, Lili, Huang, Jing, Kong, Hongfang et al. (2020) The effect of folic acid throughout pregnancy among pregnant women at high risk of pre-eclampsia: A randomized clinical trial. Pregnancy hypertension 19: 253-258	- Ineligible country Not high-income country (as defined by the OECD) as specified in the protocol. Study was conducted in China.
Zhong, Caichen, Tessing, Jillian, Lee, Brian K et al. (2020) Maternal Dietary Factors and the Risk of Autism Spectrum Disorders: A Systematic Review of Existing Evidence. Autism research: official journal of the International Society for Autism Research 13(10): 1634-1658	- Systematic review. Included studies checked for eligibility. No studies identified for inclusion because studies do not compare different doses of folic acid.
Zia, S., Rafique, M., Khan, M.A. et al. (2015) Demographic and clinical characteristics of children with sickle cell disease. Pakistan Paediatric Journal 39(2): 85-92	- Ineligible study design Cross-sectional study

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: Which groups of women should be advised to take high-dose folic acid supplements before and during the first 12 weeks of pregnancy?

Research recommendation

What is the most effective and safe dose for folic acid supplementation before and during the first 12 weeks of pregnancy for people at a high risk of conceiving a child with a neural tube defect?

Why this is important

There was no evidence to support the use of the high-dose (5 mg) folic acid during preconception and pregnancy for people at greater risk of delivering a baby with a neural tube defect. Research in this topic will add additional clarity to what is the optimal supplemental dose of folate to help prevent neural tube defects in high-risk pregnancies.

Rationale for research recommendation

Table 10: Research recommendation rationale

evidence to support the use of the 5 mg higher of folate recommended for people at greater risk of h a neural tube defect. Most other middle to high commend a supplement of between 1-4 mg for s. It is important to determine whether we are ementing, and this is even more relevant given the fortification of flour with folate.
mmends high dose (5 mg folic acid /day) when y or in early stages of pregnancy for personal or ural tube defect (NTD), previous baby with NTD, amily history of NTDs, type 1 or type 2 diabetes, antiepileptic drug. For low risk people, a dose of is recommended for all people planning re pregnant (PH11, CG137). NICE currently does dose folic acid for pregnant people with obesity. Inch will add additional clarity to what is the optimal of folate to help prevent neural tube defects in s.
dity evidence that periconceptional high dose folic th adverse outcomes but possible adverse effects. In addition to NICE recommendations, the Royal cans recommends that people with obesity take folic acid daily, based on greater risk of neural ver folate status, but without supporting evidence tudies or trials.
entified on high-dose (5 mg) folic acid during regnancy, although 5 mg is the current for those with an increased risk of conceiving a e defects or other congenital malformations, so the esearch recommendation.
economic factors

mg: milligrams; NICE: National Institute for Health and Care Excellence; NHS: National Health Service; NTD: neural tube defect

Modified PICO table

Table 11: Research recommendation modified PICO table

i able 11. Nesealcii	recommendation modified PICO table
Population	People at risk of neural tube defect (NTDs) who are trying to become pregnant or people during the first 12 weeks of a single or multiple pregnancy.
Intervention	High-dose folic acid supplementation (5 mg daily with no additional vitamins/minerals) provided ≥12 weeks prior to conception and for at least the first trimester.
Comparator	 Low-dose folic acid supplementation (<1 mg daily with no additional vitamins/minerals) provided ≥12 weeks prior to conception and for at least the first trimester.
	 Medium-dose folic acid supplementation (≥1 to <5 mg daily with no additional vitamins/minerals) provided ≥12 weeks prior to conception and for at least the first trimester.
Outcome	Primary outcomes
	Neural tube defects (NTDs) in the baby
	Birthweight centile
	Compliance
	Secondary outcomes
	Congenital heart defects
	Midline facial defects such as cleft lip or cleft palate Compared to the compared to t
	 Red blood cell (RBC) (or serum if RBC is not reported) folate concentrations in the mother
	Infant neurocognitive outcomes e.g. ASQ
	 Childhood neurodevelopmental outcomes (dichotomous outcome, not continuous outcomes such as mean change in score):
	 Severe (score of >2SD below normal on validated assessment scales, or Bayley assessment scale of mental development index [MDI] or psychomotor developmental index [PDI])
	-Moderate (score of 1-2 SD below normal on validated assessment scales, or Bayley assessment scale MDI or PDI 70-84)
	 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	Short- term follow-up (for diagnosis of congenital malformation during pregnancy or at delivery) and medium-term follow-up (for neurodevelopment outcomes which is 2- 5 years post-delivery)
Additional information	Sub-group analysis:
inomation	 Single versus multiple pregnancies Women with a previous history of a pregnancy affected by a neural tube
	defect versus not 3. Women with diabetes (type 1 and type 2) versus not
	Socio economic status and deprivation (measured using IMD)
	5. BMI ≤ 25 kg/m² versus BMI ≥ 25 kg/m² or more
	6. Age <40 years versus ≥40 years
	7. Ethnicity 8. Women who smoke versus not
	C. Tromon who office volue flot

ASQ: The Ages and Stages Questionnaire; EQ-5D: European Quality of Life Five Dimension; MDI: mental development index; mg: milligrams; NTDs: neural tube defects; PDI: psychomotor developmental index; RCT: randomized controlled trials; SF-6D: Short-Form Six-Dimension; SD: standard deviation; QALY: The quality-adjusted life-year