



Cochrane
Library

Cochrane Database of Systematic Reviews

Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children (Review)

Takahashi R, Ota E, Hoshi K, Naito T, Toyoshima Y, Yuasa H, Mori R, Nango E

Takahashi R, Ota E, Hoshi K, Naito T, Toyoshima Y, Yuasa H, Mori R, Nango E.

Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children.

Cochrane Database of Systematic Reviews 2017, Issue 10. Art. No.: CD011850.

DOI: 10.1002/14651858.CD011850.pub2.

www.cochranelibrary.com

TABLE OF CONTENTS

HEADER	1
ABSTRACT	1
PLAIN LANGUAGE SUMMARY	2
SUMMARY OF FINDINGS FOR THE MAIN COMPARISON	4
BACKGROUND	6
OBJECTIVES	7
METHODS	7
RESULTS	10
Figure 1.	11
Figure 2.	13
DISCUSSION	14
AUTHORS' CONCLUSIONS	16
ACKNOWLEDGEMENTS	16
REFERENCES	16
CHARACTERISTICS OF STUDIES	19
DATA AND ANALYSES	22
Analysis 1.1. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 1 Children with caries in the primary teeth at 3 years.	22
Analysis 1.2. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 2 Children with caries in the primary teeth at 5 years.	23
Analysis 1.3. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 3 Decayed primary surfaces at 3 years.	24
Analysis 1.4. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 4 Filled primary surfaces at 3 years.	24
Analysis 1.5. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 5 Decayed or filled primary surfaces at 3 years.	25
Analysis 1.6. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 6 Decayed primary surfaces at 5 years.	25
Analysis 1.7. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 7 Filled primary surfaces at 5 years.	26
Analysis 1.8. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 8 Decayed or filled primary surfaces at 5 years.	26
Analysis 1.9. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 9 Fluorosis (maxillary teeth) at 5 years.	27
Analysis 1.10. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 10 Fluorosis (mandibular teeth) at 5 years.	27
APPENDICES	27
CONTRIBUTIONS OF AUTHORS	29
DECLARATIONS OF INTEREST	29
SOURCES OF SUPPORT	30
DIFFERENCES BETWEEN PROTOCOL AND REVIEW	30

[Intervention Review]

Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children

Rena Takahashi¹, Erika Ota², Keika Hoshi³, Toru Naito⁴, Yoshihiro Toyoshima⁵, Hidemichi Yuasa⁶, Rintaro Mori⁷, Eishu Nango⁸

¹Cariology and Operative Dentistry, Graduate School of Medical and Dental Sciences, Tokyo Medical and Dental University, Tokyo, Japan. ²Global Health Nursing, St. Luke's International University, Graduate School of Nursing Sciences, Tokyo, Japan. ³Department of Hygiene, Kitasato University, School of Medicine, Minami-ku, Sagamihara, Japan. ⁴Department of Geriatric Dentistry, Fukuoka Dental College, Fukuoka, Japan. ⁵Human Resource Department, Hibiya Employee Clinic, The Dai-ichi Life Insurance Company, Limited, Tokyo, Japan. ⁶Department of Oral and Maxillofacial Surgery, National Hospital Organization Toyohashi Medical Center, Toyohashi, Japan. ⁷Department of Health Policy, National Center for Child Health and Development, Tokyo, Japan. ⁸Department of General Medicine, Tokyo Kita Medical Center, Tokyo, Japan

Contact address: Rintaro Mori, Department of Health Policy, National Center for Child Health and Development, 2-10-1 Okura, Setagaya-ku, Tokyo, Tokyo, 157-0074, Japan. rintaromori@gmail.com.

Editorial group: Cochrane Oral Health Group.

Publication status and date: New, published in Issue 10, 2017.

Citation: Takahashi R, Ota E, Hoshi K, Naito T, Toyoshima Y, Yuasa H, Mori R, Nango E. Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children. *Cochrane Database of Systematic Reviews* 2017, Issue 10. Art. No.: CD011850. DOI: 10.1002/14651858.CD011850.pub2.

Copyright © 2017 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

ABSTRACT

Background

Dental caries (tooth decay) is one of the most common chronic childhood diseases. Caries prevalence in most industrialised countries has declined among children over the past few decades. The probable reasons for the decline are the widespread use of fluoride toothpaste, followed by artificial water fluoridation, oral health education and a slight decrease in sugar consumption overall. However, in regions without water fluoridation, fluoride supplementation for pregnant women may be an effective way to increase fluoride intake during pregnancy. If fluoride supplements taken by pregnant women improve neonatal outcomes, pregnant women with no access to a fluoridated drinking water supply can obtain the benefits of systemic fluoridation.

Objectives

To evaluate the effects of women taking fluoride supplements (tablets, drops, lozenges or chewing gum) compared with no fluoride supplementation during pregnancy to prevent caries in the primary teeth of their children.

Search methods

Cochrane Oral Health's Information Specialist searched the following databases: Cochrane Oral Health's Trials Register (to 25 January 2017); the Cochrane Central Register of Controlled Trials (CENTRAL; 2016, Issue 11) in the Cochrane Library (searched 25 January 2017); MEDLINE Ovid (1946 to 25 January 2017); Embase Ovid (1980 to 25 January 2017); LILACS BIREME Virtual Health Library (Latin American and Caribbean Health Science Information database; 1982 to 25 January 2017); and CINAHL EBSCO (Cumulative Index to Nursing and Allied Health Literature; 1937 to 25 January 2017). We searched the US National Institutes of Health Ongoing Trials Register (ClinicalTrials.gov) and the [World Health Organization International Clinical Trials Registry Platform](http://WorldHealthOrganization.org/clinical-trials-registry) for ongoing trials to 25 January 2017. No restrictions were placed on the language or date of publication when searching the electronic databases.

Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children (Review) |

Copyright © 2017 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

Selection criteria

Randomised controlled trials (RCTs) of fluoride supplements (tablets, drops, lozenges or chewing gum) administered to women during pregnancy with the aim of preventing caries in the primary teeth of their children.

Data collection and analysis

Two review authors independently screened the titles and abstracts (when available) of all reports identified through electronic searches. Two review authors independently extracted data and assessed risk of bias, as well as evaluating overall quality of the evidence utilising the GRADE approach. We could not conduct data synthesis as only one study was included in the analysis.

Main results

Only one RCT met the inclusion criteria for this review. This RCT showed no statistical difference on decayed or filled primary tooth surfaces (dfs) and the percentage of children with caries at 3 years (risk ratio (RR) 1.46, 95% confidence interval (CI) 0.75 to 2.85; participants = 938, very low quality of evidence) and 5 years old (RR 0.84, 95% CI 0.53 to 1.33; participants = 798, very low quality of evidence). The incidence of fluorosis at 5 years was similar between the group taking fluoride supplements (tablets) during the last 6 months of pregnancy and the placebo group.

Authors' conclusions

There is no evidence that fluoride supplements taken by women during pregnancy are effective in preventing dental caries in their offspring.

PLAIN LANGUAGE SUMMARY

Fluoride supplements taken by pregnant women for preventing dental caries in the primary teeth of their children

Review question

How effective and safe is the use of fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing tooth decay in the baby teeth of their children compared with placebo (tablets or other forms of supplements without fluoride) or no treatment?

Background

Tooth decay is one of the most common health problems among children. The condition has been decreasing among children in most parts of the world over the past few decades most likely due to the widespread use of fluoride toothpaste, followed by water fluoridation, oral health education and a slight decrease in sugar consumption. If fluoride supplements taken by pregnant women can prevent tooth decay in their children, pregnant women with no access to a fluoridated drinking water supply can obtain the benefits of systemic fluoridation. Fluoride tablets, drops, lozenges or chewing gums are sucked or chewed to provide topical fluoride and ingested to provide systemic fluoride.

Study characteristics

Authors from Cochrane Oral Health carried out this review of existing studies and the evidence is current up to 25 January 2017. It includes only one study in which 1400 pregnant women were randomly allocated to fluoride treatment or placebo. In this study, a daily dose of either 1 mg sodium fluoride tablets or placebo tablets were given to participants from the fourth month of pregnancy to delivery. Both groups were encouraged to use dietary fluoride supplements after delivery in the form of drops. A total of 1175 babies were born to participants in this study, and of this number, 938 children were followed up at 3 years (464 fluoride tablets versus 484 placebo tablets) and 798 children were followed up at 5 years (398 fluoride tablets versus 400 placebo tablets) of age. Published in 1997, this study took place in communities with unfluoridated drinking water in Southern Maine, USA.

Key results

Baby teeth decay measured in children aged 3 and 5 years old was very low in both the fluoride supplement group and the placebo group. At 5 years of age, 92% of children remained decay-free in the fluoride supplement group and 91% remained decay-free in the placebo group, showing no difference between the two groups. The incidence of fluorosis at 5 years was similar between the group taking fluoride supplements (tablets) during the last 6 months of pregnancy and the placebo group.

Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children (Review) 2

Copyright © 2017 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

There is no evidence that fluoride supplements taken by women during pregnancy are effective in preventing dental caries in their offspring.

Quality of the evidence

The included study was assessed as being at high risk of bias and the evidence was of very low quality.

SUMMARY OF FINDINGS FOR THE MAIN COMPARISON *[Explanation]*

Fluoride supplementation (tablets) compared to placebo for pregnant women for preventing dental caries in the primary teeth of their children						
Population: pregnant women for preventing dental caries in the primary teeth of their children Setting: USA Intervention: fluoride supplementation (tablets) Comparison: placebo						
Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	Number of participants (studies)	Quality of the evidence (GRADE)	Comments
	Risk with placebo	Risk with fluoride supplementation (tablets)				
Children with caries in the primary teeth at 3 years	Study population		RR 1.46 (0.75 to 2.85)	938 (1 RCT)	⊕○○○ VERY LOW ^{1,2}	At 5 years RR 0.84 (0.53 to 1.33)
	30 per 1000	43 per 1000 (22 to 84)				
Decayed or filled primary tooth surfaces at 3 years	The mean decayed or filled surfaces at 3 years was 0.30	MD 0.12 higher (0.05 lower to 0.29 higher)	-	938 (1 RCT)	⊕○○○ VERY LOW ^{2,3}	Not significant at 5 years
Fluorosis (maxillary teeth) at 5 years	Study population		RR 1.79 (0.95 to 3.40)	798 (1 RCT)	⊕○○○ VERY LOW ^{1,2}	Fluorosis in mandibular teeth at 5 years RR 0.89 (0.35 to 2.29)
	35 per 1000	63 per 1000 (33 to 119)				
<p>* The risk in the intervention group (and its 95% CI) is based on the assumed risk in the comparison group and the relative effect of the intervention (and its 95% CI)</p> <p>CI: confidence interval; MD: mean difference; RCT: randomised controlled trial; RR: risk ratio</p>						
<p>GRADE Working Group grades of evidence</p> <p>High quality: we are very confident that the true effect lies close to that of the estimate of the effect</p> <p>Moderate quality: we are moderately confident in the effect estimate: the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different</p> <p>Low quality: our confidence in the effect estimate is limited: the true effect may be substantially different from the estimate of the effect</p>						

Very low quality: we have very little confidence in the effect estimate: the true effect is likely to be substantially different from the estimate of effect

¹Downgraded by 2 for imprecision: few events, and CI included appreciable benefits and harms.

²Downgraded by 1 for risk of bias: high attrition bias, and potential limitations in blinding of outcome assessment are likely to lower confidence in the estimate of effect

³Downgraded by 2 for imprecision: CI included appreciable benefits and harms.

BACKGROUND

Description of the condition

Dental caries (tooth decay) is one of the most common chronic childhood diseases, and can have a negative impact on a child's growth, speech, self-confidence, general health and quality of life (Anderson 2004; Casamassimo 2009; Jankauskiene 2010). Dental caries occurs because of long-term exposure to a mixture of acid-producing bacteria and fermentable carbohydrates, and many other factors that include saliva secretion rate and buffering capacity (Rozier 2010; Selwitz 2007). In particular, acid production from bacteria, especially *mutans Streptococci* and *Lactobacilli*, and the subsequent decrease in local pH, cause the demineralization of tooth tissue (Featherstone 2004). If this process is not reversed, carious lesion progresses. Dissolved calcium and phosphate mineral ions can be redeposited on the tooth surface even though they are provided from saliva. This demineralization/remineralization process occurs continuously in oral fluids. Dental caries is generated when the demineralization/remineralization process lose the balance.

Dental caries is widespread in all countries (WHO 2009). Dental caries affects 30% to 50% of children aged 5 to 6 years (Armfield 2009; CDC 2007; Public Health England 2012), 60% to 90% of school-aged children and a large majority of adults (Petersen 2005). An increasing number of decayed, missing or filled teeth (dmft) is reported in some low-income and middle-income countries (Bagramian 2009). Although the mean dmft has declined in many high-income countries, this disparity remains and some population groups have high dmft (Bagramian 2009; Dawkins 2013; Jones 2017). Prevalence of dental caries is associated with increased consumption of sugar in the diet including high-sugar confectionery and sweet carbonated beverages (Ismail 1997), socioeconomic status, dental insurance coverage and residential locations (Campus 2009; Dawkins 2013; Hobdell 2003; Kolker 2007).

Several studies have reported that caries in primary teeth are correlated with caries in permanent teeth (Helfenstein 1991; Li 2002; Seppa 1989). 'Early childhood caries', which is the presence of at least one carious lesion on a primary tooth in a child under the age of 6 years, is a serious problem in the world (Anil 2017). If the fluoride supplementation in pregnant woman is effective to prevent dental caries in the primary teeth of their children, the prevalence of dental caries can be prevented.

Description of the intervention

The benefits of topical fluorides, such as toothpastes, gels, varnishes and mouthrinses, for preventing dental caries in children and adolescents are well established (Marinho 2003a; Marinho

2003b; Marinho 2013; Marinho 2015; Marinho 2016). Topical fluoride plays an important role in preventing dental caries (Marinho 2003b) including the inhibition of demineralization of the crystal surfaces, the enhancement of remineralization in demineralized lesions (Featherstone 1990) and inhibition of bacterial metabolism (Featherstone 1999; Featherstone 2000; Fejerskov 2004). The American Dental Association (ADA) and Centers for Disease Control and Prevention (CDC) state that an appropriate concentration of fluoride, which varies depending on age, is effective to prevent dental caries (CDC 2001; Rozier 2010).

The upper limit of fluoride intake from all sources (fluoridated water, food, beverages, fluoride dental products and dietary fluoride supplements) is set at 0.10 mg/kg/day for infants, toddlers, and children through to 8 years old. For older children and adults, who are no longer at risk for dental fluorosis, the upper limit of fluoride is set at 10 mg/day regardless of weight (Levy 1999). CDC reported water fluoridation is especially beneficial for communities of low socioeconomic status (CDC 1999).

It should be noted that separation of pre-eruptive, post-eruptive, systemic or topical effects of fluoride is impossible. Fluoridated water may have all effects. Topically applied fluorides are not intended for ingestion but can be swallowed unintentionally. Fluoride drops, lozenges or chewing gums are sucked or chewed to provide topical fluoride and ingested to provide systemic fluoride. Fluoride supplements taken during pregnancy have the effect of prenatal fluoride and possibly reduce dental caries in offspring (Stephen 1993). However, risks associated with exposure to fluoride during pregnancy including miscarriage, premature delivery and premature birth were also reported (Diouf 2012; Sastry 2011).

How the intervention might work

The enamel formation process is composed of two principal stages: secretory stage and maturation stage. In the former stage, ameloblasts produce protein matrix (predominantly amelogenins) and crystallites are deposited in the protein matrix. In the latter stage, ameloblasts transport the substances used in enamel formation out of the enamel. Excess water and organic materials are removed and mineral is transported into the tissue in order to achieve full mineralization of enamel (Hiller 1975; Termine 1980).

Fluoride can be transported from maternal serum to the fetus and prenatal deciduous enamel through the placenta (Toyama 2001). The mechanism of fluoride placental transfer is controversial; there are some hypotheses that the placenta allows passive diffusion of fluoride from mother to fetus, while others suggest that the placenta acts as a barrier. It has been reported that the placenta allows passive diffusion of fluoride from mother to fetus when fluoride intake is low and that the placenta acts as a selective barrier when fluoride intake exceeds a particular level (Gupta 1993; Toyama 2001). In the development of enamel formation, fluoride is incorporated into the crystal lattice and binds to calcium, which is contained within the protein matrix. As a result, fluoride alters crystal

formation in the enamel matrix and strengthens the properties of the enamel crystals (Tanimoto 2008).

All primary tooth enamel and some permanent tooth enamel start to grow and continue to develop in utero (Kraus 1965). Most of the permanent tooth enamel start to grow after birth. It has been reported that while the effect of fluoride supplements after birth was unclear on deciduous teeth, it was associated with a reduction in caries increment in permanent teeth (Tubert-Jeannin 2011). Namely, it might be suggested that postnatal systemic fluoride did not attribute to caries prevention in primary teeth. In regions without water fluoridation, fluoride supplementation for pregnant women may be an effective way to increase fluoride intake during pregnancy. If fluoride supplements taken by pregnant women improve neonatal outcomes, pregnant women with no access to a fluoridated drinking water supply can obtain the benefits of systemic fluoridation.

Why it is important to do this review

Recent systematic reviews and Cochrane Reviews have evaluated evidence regarding the effectiveness and safety of fluoride treatment and the adverse effects of high fluoride exposure in children and adults (Choi 2012; Iheozor-Ejiofor 2015; Ismail 2008; Marinho 2003a; Marinho 2003b; Marinho 2013; Marinho 2015; Marinho 2016; Tubert-Jeannin 2011; Wong 2010). However, it remains uncertain whether fluoride supplementation in pregnant women is effective in preventing dental caries in their offspring. Currently, no systematic reviews have investigated the effectiveness and safety of this intervention. This Cochrane Review aims to address this gap in the literature and assess the current available evidence on fluoride supplementation during pregnancy.

OBJECTIVES

To evaluate the effects of women taking fluoride supplements (tablets, drops, lozenges or chewing gum) compared with no fluoride supplementation during pregnancy to prevent caries in the primary teeth of their children.

METHODS

Criteria for considering studies for this review

Types of studies

We included all randomised controlled trials (RCTs), including quasi-randomised and cluster-randomised trials. We excluded

cross-over trials as they are an inappropriate study design (as the intervention may have a lasting effect that compromises entry to subsequent periods of the trial). We excluded observational studies (including cohort studies, case-control studies, etc.). We included both clinical and community-based trials.

Types of participants

We included pregnant women, regardless of their dental caries, exposure to fluorides, level of dental treatment, nationality or level of education. The women may or may not have had access to fluoridated water (naturally or artificially).

Types of interventions

We included studies of fluoride supplementation (tablets, drops, lozenges or chewing gum) of any dosage, frequency, duration and timing of delivery, which may or may not have included the use of topical fluorides such as fluoride dentifrice, fluoride rinse and topical fluoride application, compared with no fluoride supplementation.

Control group: no treatment or placebo.

Types of outcome measures

Primary outcomes

Primary outcomes (for deciduous teeth of children up to 6 years of age).

- Number of children with caries in the primary teeth.
- Decayed, missing and filled primary teeth (dmft) and components.
- Decayed, missing and filled primary tooth surfaces (dmfs) and components.
- Fluorosis.

Secondary outcomes

- Adverse effects (apart from fluorosis), e.g. miscarriage, premature delivery, or dental and any other possible negative effects. A full investigation of adverse effects was not possible as we did not include observational or retrospective epidemiological studies.

Search methods for identification of studies

Electronic searches

Cochrane Oral Health's Information Specialist conducted systematic searches in the following databases for randomised controlled trials and controlled clinical trials. There were no language, publication year or publication status restrictions:

- Cochrane Oral Health's Trials Register (searched 25 January 2017) ([Appendix 1](#));
- Cochrane Central Register of Controlled Trials (CENTRAL; 2016, Issue 11) in the Cochrane Library (searched 25 January 2017) ([Appendix 2](#));
- MEDLINE Ovid (1946 to 25 January 2017) ([Appendix 3](#));
- Embase Ovid (1980 to 25 January 2017) ([Appendix 4](#));
- LILACS BIREME Virtual Health Library (Latin American and Caribbean Health Science Information database; 1982 to 25 January 2017) ([Appendix 5](#));
- CINAHL EBSCO (Cumulative Index to Nursing and Allied Health Literature; 1937 to 25 January 2017) ([Appendix 6](#)).

Subject strategies were modelled on the search strategy designed for MEDLINE Ovid.

Searching other resources

We searched the following trial registries for ongoing studies:

- US National Institutes of Health Ongoing Trials Register ClinicalTrials.gov ([clinicaltrials.gov](#); searched 25 January 2017) ([Appendix 7](#));
- World Health Organization International Clinical Trials Registry Platform ([apps.who.int/trialsearch](#); searched 25 January 2017) ([Appendix 8](#)).

We searched the reference lists of included studies and relevant systematic reviews for further studies.

We did not perform a separate search for adverse effects of interventions used, and we considered adverse effects described in included studies only.

Data collection and analysis

Selection of studies

Two review authors (Rena Takahashi (RT) and Keika Hoshi (KH)) independently screened the titles and abstracts (when available) of all reports identified through the electronic searches and hand-searching that were entered into EndNote X7 software. We obtained the full text of potentially relevant studies or studies where it was difficult to make a clear decision from only the title and abstract. The full-text articles were assessed independently by two review authors to determine if they met the inclusion criteria. We resolved any disagreements by discussion. If this was not possible, we consulted a third review author. At the time of exclusion, we recorded reasons for exclusion in the 'Characteristics of excluded studies' tables.

Data extraction and management

Two review authors (RT and Erika Ota (EO)) independently extracted data using data extraction forms. We extracted data related to settings, participants (e.g. inclusion and exclusion criteria for pregnant women), interventions (e.g. type of intervention, comparison), outcomes (e.g. outcomes reported in the paper, duration and rates of follow-up, adverse effects), methods (e.g. study designs, randomisation methods) and other information (e.g. pharmaceutical sponsorship data). We recorded if clinical trials reported the presence of calcium in the fluoride supplement. We noted the topical fluoride exposure of children (up to 6 years of age) in the follow-up period of the studies. We included unpublished research data from recognised research groups and experts in the field, obtained via personal contact if the study was in the clinical register.

We resolved any disagreements by discussion and consultation with a third review author. In this review, we did not contact trial authors. In future updates, we will contact trial authors (if possible) asking for assistance with data transformation or raw data, if the data are not reported in a format suitable for analysis.

Assessment of risk of bias in included studies

Two review authors (KH and Yoshihiro Toyoshima (YT)) assessed the risk of bias independently for each study using the criteria outlined in the *Cochrane Handbook for Systematic Reviews of Interventions* ([Higgins 2011](#)). We resolved any disagreements by discussion or by involving a third assessor (EO). We assessed the following domains of risk of bias: (1) random sequence generation (selection bias), (2) allocation concealment (selection bias), (3) blinding of participants and personnel (performance bias), (4) blinding of outcome assessors (detection bias), (5) incomplete outcome data (attrition bias), (6) selective reporting (reporting bias) and (7) other bias (recruitment bias, bias influenced by funding source, etc.).

Each domain was assessed as at either low, high or unclear risk of bias. We categorised the overall risk of bias of individual studies as follows:

- low risk of bias (plausible bias unlikely to seriously alter the results) if all domains were at low risk of bias;
- unclear risk of bias (plausible bias that raises some doubt about the results) if one or more domains had an unclear risk of bias, but none at high risk of bias;
- high risk of bias (plausible bias that seriously weakens confidence in the results) if one or more domains were at high risk of bias.

Measures of treatment effect

For dichotomous outcomes, we calculated risk ratios for differences between the intervention and comparison groups, along with 95% confidence intervals (CI). For continuous outcomes, we calculated the mean difference (MD) and 95% CIs where means and

standard deviations (SD) were presented or were calculable. We did not calculate the standardised mean difference (SMD) in this review. In future updates, where a continuous outcome is measured using different scales, we will calculate the SMD and SDs.

Unit of analysis issues

We did not include any cluster-randomised trials in the analyses. In future updates, if we include cluster-randomised trials, we will adjust their sample sizes using the methods described in Section 16 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011) using an estimate of the intracluster correlation coefficient (ICC) with either data from the trials (if possible), data from a similar trial or data from a study of a similar population. We will conduct sensitivity analyses to see the effect of variation in the ICC when we use ICCs from other sources. We plan to combine the relevant outcomes from both cluster-randomised and individually randomised trials if there is no substantial heterogeneity between the two study designs.

Dealing with missing data

In this review we did not contact trial authors. In future updates, where possible, we will contact trial authors to provide missing data. We will note levels of attrition for included studies. We will conduct sensitivity analysis to check the impact of including studies with substantial levels (more than 20%) of missing data in the overall assessment of the intervention effect.

If possible, we will conduct analysis on an intention-to-treat basis. If there are missing data, the denominator for each outcome in each trial will be the number randomised minus any participants whose outcomes are known to be missing in the analysis. If there are missing standard deviations of the continuous data, we will use the methods described in the *Cochrane Handbook of Systematic Reviews of Interventions* Section 7.7.3 (Higgins 2011) to estimate them.

Assessment of heterogeneity

We could not assess heterogeneity due to the inclusion of only one study. In future updates, we will assess heterogeneity by inspection of forest plots of the estimates and confidence intervals of treatment effects. We will assess statistical heterogeneity in each meta-analysis using the I^2 and Chi^2 statistics. We defined substantial heterogeneity as I^2 greater than 50%, or if there was a low P value (less than 0.10) in the Chi^2 test for heterogeneity.

Assessment of reporting biases

In future updates, if more than 10 trials are identified for any meta-analysis, we will assess publication bias using visual assessment of funnel plots according to recommendations in the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). Should asymmetry be identified in the contour-enhanced funnel

plots, we will investigate possible causes by performing exploratory analyses.

Data synthesis

We could not conduct data synthesis as only one study was included in the review. In future updates, we will conduct meta-analyses for studies with comparable analyses that report similar outcome measures using forest plots in Review Manager software (RevMan) (RevMan 2014). We will combine risk ratios for dichotomous data and mean differences for continuous data using random-effects models, assuming that the identified studies allow this procedure. If random-effects analyses are conducted, we will present the results as the average treatment effect with 95% CIs, and estimates of I^2 . If there are few studies or small sample sizes, it may be impossible to estimate between-study variance with any precision. In that case, a random-effects analysis would provide poor estimates of the distribution of intervention effects, therefore we would use a fixed-effect model (Higgins 2011).

In future updates, we will analyse cluster-RCTs at the individual level. We will meta-analyse results from appropriately analysed cluster-RCTs using the generic inverse variance method in RevMan. If original analyses do not account for clustering, we will conduct an adjusted analysis, provided that the necessary information (e.g. mean cluster size, proportion of individuals with events, ICCs) can be extracted (Higgins 2011). We will calculate the prevented fraction.

Subgroup analysis and investigation of heterogeneity

We could not assess heterogeneity by inspection of forest plots of the estimates and confidence intervals of treatment effects as only one study was included in the review. In future updates, if we identify substantial heterogeneity in the primary outcomes, we will conduct subgroup analyses for these relevant and clinically meaningful subgroups where sufficient data are available. In studies with more than one intervention group, such as those comparing different frequencies of application or different types of supplements, we will consider the results from all relevant experimental groups separately in the meta-analyses.

Moreover, we could not conduct two different subgroup analyses:

- low- and middle-income countries versus high-income countries (defined by World Bank criteria);
- region of high-level fluoride concentration in tap water or well water (1.5 mg/L or more, 0.3 mg/L or more) versus region of low- to medium-level fluoride concentration.

Sensitivity analysis

In future updates, we will undertake sensitivity analysis for primary outcomes based on the risk of bias assessment for allocation concealment and attrition rates. We will redo analyses and remove studies that are at high risk of bias for these domains in order to assess whether this makes any difference to the overall result.

Summary of findings

We created 'Summary of findings' tables using GRADE profiler (GRADEpro GDT 2015), with data imported from RevMan. We used the GRADE approach (Schünemann 2009) to assess the quality of the body of evidence relating to primary outcomes for the main comparisons. The quality of the body of evidence for each outcome was assessed under five domains (study limitations, consistency of effect, imprecision, indirectness and publication bias) and judged to be of high, moderate, low or very low quality.

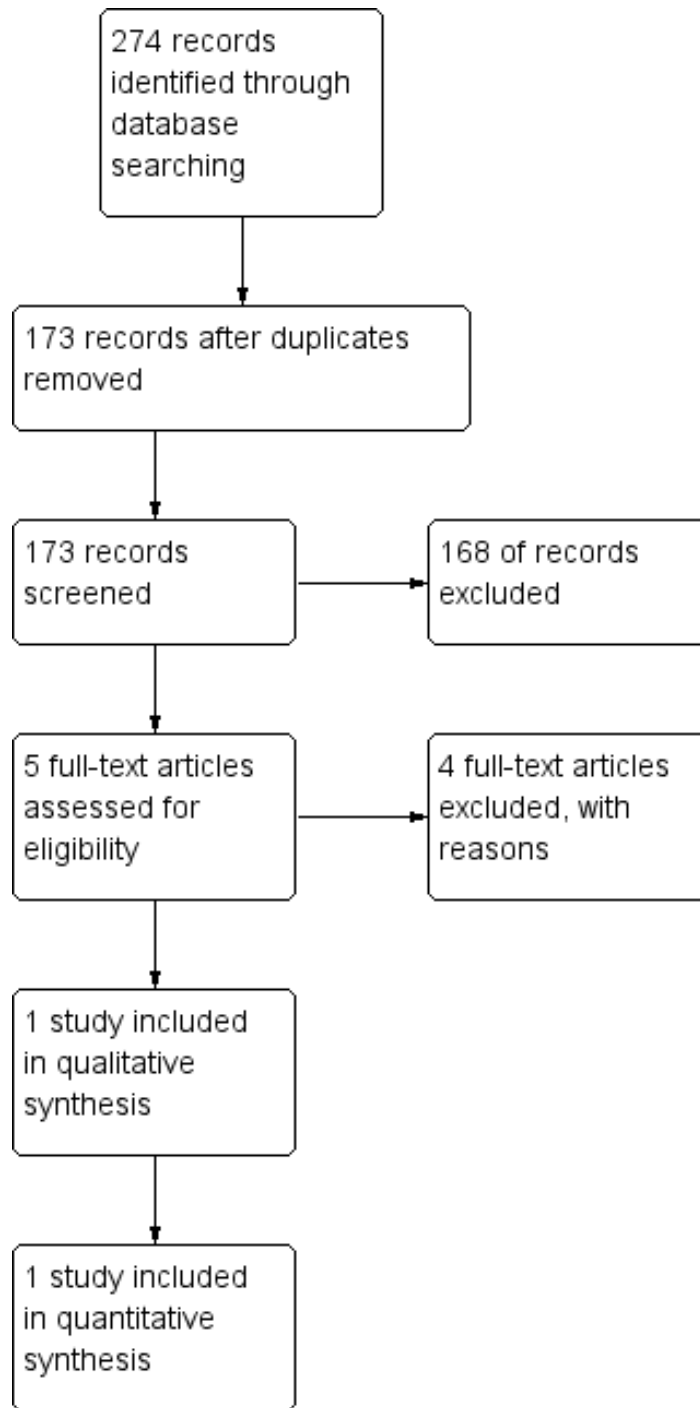
RESULTS

Description of studies

Results of the search

A total of 173 references were identified by the above search strategy after duplicates were removed. Assessment of the titles and abstracts, where available, resulted in five references of potential relevance; all of which were obtained in full. Four full-text articles were rejected since they were not randomised controlled trials (RCTs). We found one study suitable for inclusion in this review (Figure 1).

Figure 1. Study flow diagram.



Included studies

We included one individual RCT (Leverett 1997). See [Characteristics of included studies](#) table.

Participants

This trial recruited 1400 pregnant women in the first trimester residing in communities served by fluoride-deficient drinking water. There were 1175 babies born to participants and of these, 938 children were followed up at 3 years (intervention 464 versus control 484) and 798 children were followed up at 5 years (intervention 398 versus control 400).

Interventions and comparisons

The intervention group received one 2.2 mg dose of sodium fluoride (NaF) (1 mg active fluoride ion) in the form of one tablet to be taken daily from the fourth month of pregnancy. The control group received placebo tablets (no fluoride during pregnancy). Both the intervention and control groups received fluoride drops from birth to 2 years of age and one 0.5 mg tablet daily for children aged 2 to 3 years.

Participants were contacted in order to find out how many tablets remained in their supplied bottles. On the basis of that count, new supplies were mailed to coincide with the estimated date of exhaustion of the previous supply. Compliance during the prenatal

period was very good. Of 938 subjects that were examined at the 3-year interval, 88% had excellent adherence to the prenatal protocol with mean compliance scores (standard deviation (SD)) of 0.94 (0.10) and 0.95 (0.12) among the treatment and control groups respectively.

Outcomes

Caries experience between the two groups and the percentage of children with no caries in the primary teeth. Children's mean decayed, filled primary tooth surfaces (dfs) was taken at 3 and 5 years.

Settings

Communities with unfluoridated drinking water in Southern Maine, USA.

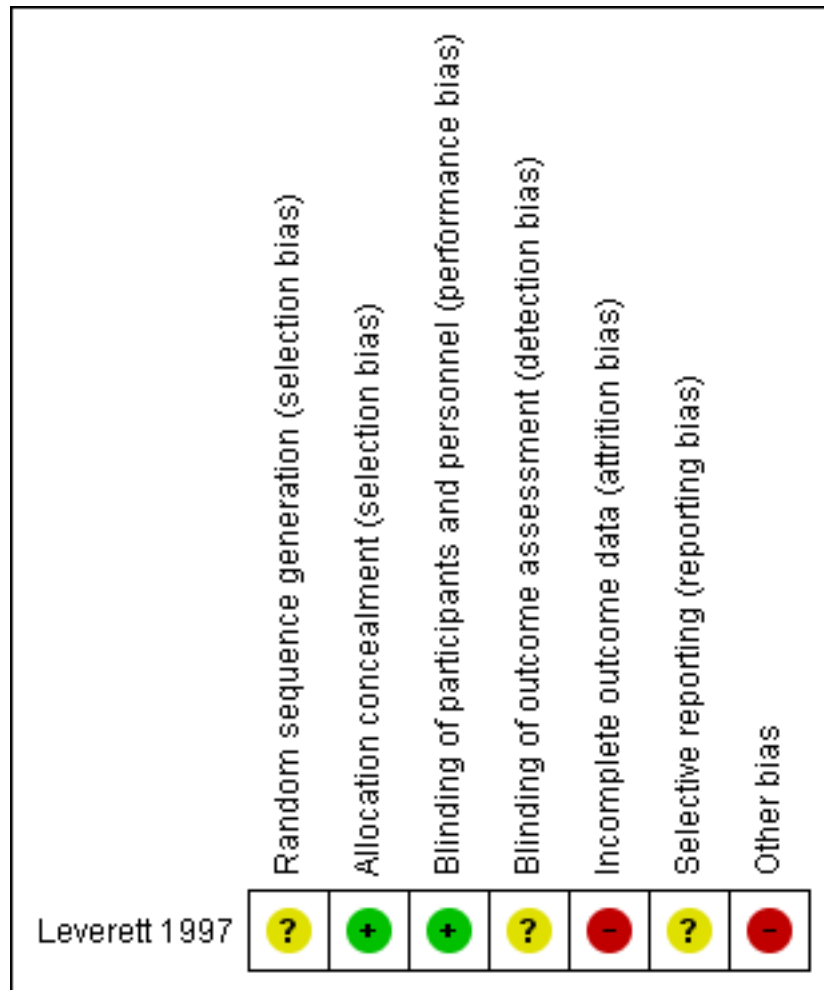
Excluded studies

We excluded four studies (Glenn 1979; Glenn 1982; Glenn 1984; Restrepo 1993) after full-text assessment because these studies were not RCTs. See [Characteristics of excluded studies](#) table.

Risk of bias in included studies

We summarised risk of bias graphically using the plots available in RevMan (RevMan 2014). See [Figure 2](#). The included study was at high overall risk of bias.

Figure 2. Risk of bias summary: review authors' judgements about each risk of bias item for each included study.



Allocation

We assessed random sequence generation (selection bias) as being at unclear risk of bias (the method of randomisation was not described in detail), and allocation concealment (selection bias) as at low risk of bias.

Blinding

We assessed blinding of participants and personnel (performance bias) as at low risk of bias and blinding of outcome assessment (detection bias) as being at unclear risk of bias (no description provided).

Incomplete outcome data

We assessed incomplete outcome data (attrition bias) as at high risk of bias.

Selective reporting

We assessed selective reporting (reporting bias) as at unclear risk of bias (no protocol or registry was available for the study).

Other potential sources of bias

We assessed the other potential sources of bias as at high risk of bias, as administering fluoride supplementation to both groups after delivery might affect the results.

Effects of interventions

See: [Summary of findings for the main comparison](#) Fluoride supplementation (tablets) compared to placebo for pregnant women for preventing dental caries in the primary teeth of their children

Fluoride supplementation (tablets) versus placebo

One trial involving 938 women and their children was included (Leverett 1997).

Primary outcomes

Number of children with caries in the primary teeth

For this primary outcome, there was no difference in effect for children with caries in the primary teeth at 3 years in the fluoride supplementation group compared to the control group (risk ratio (RR) 1.46, 95% confidence interval (CI) 0.75 to 2.85; participants = 938; studies = 1) (Analysis 1.1).

There was no difference in effect at 5 years (RR 0.84, 95% CI 0.53 to 1.33; participants = 798; studies = 1) (Analysis 1.2).

Decayed, missing and filled primary teeth (dmft) and components

Outcome not assessed.

Decayed, missing and filled primary tooth surfaces (dmfs) and components

There was no difference in effect on decayed surfaces at 3 years (mean difference (MD) 0.05, 95% CI -0.02 to 0.12; participants = 938; studies = 1) (Analysis 1.3); filled surfaces at 3 years (MD 0.07, 95% CI -0.07 to 0.21; participants = 938; studies = 1) (Analysis 1.4); decayed or filled surfaces at 3 years (MD 0.12, 95% CI -0.05 to 0.29; participants = 938; studies = 1) (Analysis 1.5); decayed surfaces at 5 years (MD -0.06, 95% CI -0.17 to 0.05; participants = 798; studies = 1) (Analysis 1.6); filled surfaces at 5 years (MD 0.03, 95% CI -0.28 to 0.34; participants = 798; studies = 1) (Analysis 1.7); and decayed or filled surfaces at 5 years (MD -0.05, 95% CI -0.42 to 0.32; participants = 798; studies = 1) (Analysis 1.8). Leverett 1997 did not assess missing surfaces.

Fluorosis

Regarding side effects, there was no difference in effect for fluorosis (maxillary teeth) at 5 years (RR 1.79, 95% CI 0.95 to 3.40; participants = 798; studies = 1) (Analysis 1.9); and fluorosis (mandibular teeth) at 5 years (RR 0.89, 95% CI 0.35 to 2.29; participants = 798; studies = 1) (Analysis 1.10).

Secondary outcomes

Adverse effects (apart from fluorosis)

There were no other adverse events of interest for the review reported in this trial.

DISCUSSION

Summary of main results

The main question addressed by this review was the efficacy of fluoride supplementation (with tablets, drops, lozenges or chewing gum) in women during pregnancy in preventing caries in the primary teeth of their children. Only one randomised controlled trial (RCT) (Leverett 1997) met the inclusion criteria for this review. This RCT had some limitations.

This RCT showed no statistical difference on decayed, filled primary tooth surfaces (dfs) and percentage of children with caries at 3 and 5 years. The incidence of fluorosis at 5 years was similar between the group taking fluoride supplements during the last 6 months of pregnancy and the placebo group. See [Summary of findings for the main comparison](#) for the summary of the main results.

The trial authors stated that fluoride carried over into the postnatal period might well have contributed to the low level of dental caries in both groups.

Fluorosis was scored as very mild using Dean's score (Dean 1942). The number of all types of tooth fluorosis was counted. Fluorosis event rate was too low to detect a difference between the groups.

Overall completeness and applicability of evidence

No evidence was found on decayed, missing and filled primary teeth (dmft) or missing surfaces (ms). Limited information was available on adverse events, and only dental fluorosis at 5 years was reported. In addition, systemic side effects were not examined in the included trial. The different dosages of fluoride supplements (tablets, drops, lozenges or chewing gum) were not explored in this review. Furthermore in the included study, fluoride supplements were routinely given to both the intervention and control groups after birth. At the time that this RCT was conducted, ingestion of fluoride supplements until the age of 3 was common practice. Therefore, fluoride supplements were given to both the intervention and control groups. Whilst the effect of fluoride supplements on the primary teeth after birth is unclear (Tubert-Jeannin 2011), children could obtain the benefits of fluoride supplements after

birth (Limeback 1999). Thus, the effect of prenatal fluoride supplementation is unclear. In future research, a postnatal preventive approach would be needed equally between the groups. Such ethical consideration is necessary in a study to investigate the effect of prenatal fluoride supplementation in pregnant women for preventing dental caries in the primary teeth of their children.

Quality of the evidence

Overall, the included trial was at high risk of bias due mainly to high attrition bias. Due to the long-term follow-up, the rate of losses to follow-up was quite high (20% at 3 years and 38% at 5 years). The study was affected by attrition bias, however most of the other biases such as selection bias (random sequence generation), blinding of outcome assessment (detection bias), and reporting bias were not evaluated or no information was available and were assessed as unclear. Participant compliance could also influence the result of the study.

The quality of the evidence as assessed using GRADE (Summary of findings for the main comparison) was very low for children with caries in the primary teeth at 3 years. This was downgraded due to potential limitations in the blinding of outcome assessment, which are likely to lower confidence in the estimate of effect, as well as imprecision due to few events, and the 95% confidence interval (CI) included appreciable benefits and harms. The quality of the evidence for children with caries in the primary teeth at 5 years was graded as very low due to high attrition bias and imprecision (few events and the CI included appreciable benefits and harms). The evidence for the outcomes of decayed or filled primary tooth surfaces at 3 years and decayed or filled surfaces at 5 years was considered to be of very low quality due to high attrition bias, potential limitations in the blinding of outcome assessment, which were likely to lower confidence in the estimate of effect, and imprecision (CI included appreciable benefits and harms). The evidence for fluorosis (maxillary teeth) at 5 years and fluorosis (mandibular teeth) at 5 years was also of very low quality, downgraded due to imprecision (few events and CI included appreciable benefits and harms), and risk of bias (high attrition bias and potential limitations in blinding of outcome assessment).

Potential biases in the review process

We tried to minimize the potential biases in this review following the guidance provided in the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). For example, the search was conducted by the Cochrane Oral Health's Information Specialist. Two review authors screened and assessed studies for eligibility for inclusion and risk of bias independently. We described the reasons for each judgement of risk of bias. Data entry was checked by two review authors.

Agreements and disagreements with other studies or reviews

A previous review (Fassman 1993) of prenatal fluoridation also mentioned that the effect was inconclusive. However, the quality of this previous review is questionable. First, inclusion criteria were not clearly shown. Secondly, quality assessment for included studies was not implemented. Those drawbacks are possibly due to the fact that at the time of publication, the systematic review process was not well established. Our study examined relevant literature thoroughly and found that the effect of prenatal fluoride supplement was inconclusive.

We only included one RCT in this review, therefore, we conducted further searches of the literature for observational studies for discussion to obtain more information below.

Kailis et al (Kailis 1968) enrolled 374 children in Perth, Australia, at the age of 4 to 6 years old and implemented a questionnaire on prenatal fluoridation. Due to the study method, the dose, duration, and gestational age of ingestion were not reported. Participants were divided into three groups (non-fluoride, postnatal fluoride, and prenatal and postnatal fluoride). Because our question was to evaluate the effect of prenatal fluoride supplementation, we extracted the results of postnatal fluoride, and prenatal and postnatal fluoride groups. The decayed, missing and filled primary teeth (dmft) rates and the percentage of caries-free children among the prenatal and postnatal fluoride group were significantly lower than that of the postnatal fluoride group.

Glenn et al (Glenn 1982) examined 375 children in Miami, USA without a prenatal fluoride supplement (without PNF) and 117 children with a prenatal 2.2 mg sodium fluoride tablet (with PNF) in a fluoridated water area. Because of the study method, duration and gestational age of ingestion were not reported. The results were as follows: the mean birth weight of the group with PNF was heavier than that of the group without PNF. The mean birth length of the group with PNF was longer than that of the group without PNF group. The group without PNF included two children with Down's syndrome, two children with an intellectual disability, one child with an intraventricular septal defect, one child with minimal brain dysfunction syndrome, one child with clubfoot, one child with congenital hip dysplasia, one child with congenital epidermolysis bullosa, four children with congenitally missing teeth, and six children with supernumerary teeth. No medical and dental defects were found in the group with PNF, while no dental fluorosis were found in either group.

Accumulated fluoride to immature matrix proteins onto the apatite crystal surface of calcifying deciduous dentition seems to be the essential mechanism for preventing caries. Prichard examined the prenatal and postnatal effects of fluoride supplementation on Australian children. There were three groups: a non-fluoride group, a prenatal and postnatal fluoride supplementation group, and a postnatal fluoride supplementation group. He reported caries reduction in primary teeth of 70% after prenatal and postnatal fluoride supplementation and 40% after postnatal

fluoride supplementation only (Prichard 1969). From this result, prenatal fluoride supplementation might work if the placenta does not act as a barrier.

As we mentioned in the introduction, fluoride placental transfer is questionable and no new investigation has been undertaken on this topic in 20 years. This theoretical weakness discourages investigators from administering prenatal fluoride supplementation. In contrast, topical fluoride administration after birth has a growing evidence base of efficacy, and topical fluoride is currently considered to have a caries-inhibiting effect (Marinho 2003b). As Glenn 1982 suggested, the probability of side effects of prenatal fluoride ingestion is another point to consider. A lack of evidence for prenatal fluoride treatment, the establishment of an alternative method, and the probability of side effects means that limited resources are available for RCTs as well as observational studies. Overall, only poor quality studies were published.

AUTHORS' CONCLUSIONS

Implications for practice

There is no evidence that fluoride supplements taken by women during pregnancy are effective in preventing dental caries in their offspring.

Implications for research

Careful consideration should be given to whether or not this is a priority topic for future research. The caries levels in the included study are far below those which cause concern. If future studies are undertaken they should focus on areas where young children are at high risk of developing caries.

ACKNOWLEDGEMENTS

We would like to thank Cochrane Oral Health for their help in developing this review. We thank Valeria Marinho, Helen Worthington, Annetta Tsang and Derek Richards for their comments on the draft. We would also like to thank Emma Barber for her editorial assistance in preparing the protocol for this review, and Anne Littlewood for developing the search strategy.

REFERENCES

References to studies included in this review

Leverett 1997 *{published data only}*

Leverett DH, Adair SM, Vaughan BW, Proskin HM, Moss ME. Randomized clinical trial of the effect of prenatal fluoride supplements in preventing dental caries. *Caries Research* 1997;**31**(3):174–9. PUBMED: 9165186]

References to studies excluded from this review

Glenn 1979 *{published data only}*

Glenn FB. Immunity conveyed by sodium-fluoride supplement during pregnancy: part II. *ASDC Journal of Dentistry for Children* 1979;**46**(1):17–24. PUBMED: 283075]

Glenn 1982 *{published data only}*

Glenn FB, Glenn WD 3rd, Duncan RC. Fluoride tablet supplementation during pregnancy for caries immunity: a study of the offspring produced. *American Journal of Obstetrics and Gynecology* 1982;**143**(5):560–4. PUBMED: 7091227]

Glenn 1984 *{published data only}*

Glenn FB, Glenn WD 3rd, Duncan RC. Prenatal fluoride tablet supplementation and the fluoride content of teeth: Part VII. *ASDC Journal of Dentistry for Children* 1984;**51**(5):344–51. PUBMED: 6592186]

Restrepo 1993 *{published data only}*

Restrepo Baena EA. Vitamins and minerals supplements during pregnancy [Suplemento de vitaminas y minerales durante el periódico de gestión]. *Iatreia* 1993;**6**(3):144–9.

Additional references

Anderson 2004

Anderson HK, Drummond BK, Thomson WM. Changes in aspects of children's oral-health-related quality of life following dental treatment under general anaesthesia. *International Journal of Paediatric Dentistry* 2004;**14**(5): 317–25.

Anil 2017

Anil S, Anand PS. Early childhood caries: prevalence, risk factors, and prevention. *Frontiers in Pediatrics* 2017;**5**:157. [PUBMED: 28770188]

Armfield 2009

Armfield JM, Spencer AJ, Brennan DS. Dental health of Australia's teenagers and pre-teen children: the Child Dental Health Survey, Australia 2003-04. Canberra: Australian Institute of Health and Welfare; 2009. Dental Statistics and Research Series No. 52 Cat. No. DEN 199.

Bagramian 2009

Bagramian RA, Garcia-Godoy F, Volpe AR. The global increase in dental caries. A pending public health crisis. *American Journal of Dentistry* 2009;**22**(1):3–8. [PUBMED: 19281105]

Campus 2009

Campus G, Solinas G, Strohmenger L, Cagetti MG, Senna A, Minelli L, et al. National pathfinder survey on children's oral health in Italy: pattern and severity of caries disease in 4-year-olds. *Caries Research* 2009;**43**(2):155–62.

Casamassimo 2009

Casamassimo PS, Thikkurissy S, Edelstein BL, Maiorini E. Beyond the dmft: the human and economic cost of early childhood caries. *Journal of the American Dental Association* 2009;**140**(6):650–7.

CDC 1999

CDC. Achievements in public health, 1900-1999: fluoridation of drinking water to prevent dental caries. *MMWR Weekly* 1999;**48**(41):933–40.

CDC 2001

CDC. Recommendations for using fluoride to prevent and control dental caries in the United States. *MMWR Recommendations and Reports* 2001;**50**(RR14):1–42.

CDC 2007

Dye BA, Tan S, Smith V, Lewis BG, Barker LK, Thornton-Evans G, et al. Trends in oral health status: United States, 1988-1994 and 1999-2004. *Vital and Health Statistics. Series 11, Data from the National Health Survey* 2007;**(248)**:1–92.

Choi 2012

Choi AL, Sun G, Zhang Y, Grandjean P. Developmental fluoride neurotoxicity: a systematic review and meta-analysis. *Environmental Health Perspectives* 2012;**120**(10):1362–8.

Dawkins 2013

Dawkins E, Michimi A, Ellis-Griffith G, Peterson T, Carter D, English G. Dental caries among children visiting a mobile dental clinic in South Central Kentucky: a pooled cross-sectional study. *BMC Oral Health* 2013;**13**:19. [PUBMED: 23639250]

Dean 1942

Dean HT. The investigation of physiological effects by the epidemiological method. In: Moulton FR editor(s). *Fluorine and Dental Health*. Washington DC: American Association for the Advancement of Science, 1942:23–31.

Diouf 2012

Diouf M, Cisse D, Lo CM, Ly M, Faye D, Ndiaye O. Pregnant women living in areas of endemic fluorosis in Senegal and low birthweight newborns: case-control study [Femme enceinte vivant en zone de fluorose endémique au Sénégal et faible poids du nouveau-né à la naissance: étude cas-témoins]. *Revue d'Epidémiologie et de Santé Publique* 2012;**60**(2):103–8.

Fassman 1993

Fassman DK. Prenatal fluoridation. A literature review. *New York State Dental Journal* 1993;**59**(6):47–51.

Featherstone 1990

Featherstone JD, Glena R, Shariati M, Shields CP. Dependence of in vitro demineralization of apatite and remineralization of dental enamel on fluoride concentration.

Journal of Dental Research 1990;**69** Spec No:620-5; discussion 634-6. [PUBMED: 2312892]

Featherstone 1999

Featherstone JD. Prevention and reversal of dental caries: role of low level fluoride. *Community Dentistry and Oral Epidemiology* 1999;**27**(1):31–40. [PUBMED: 10086924]

Featherstone 2000

Featherstone JD. The science and practice of caries prevention. *Journal of the American Dental Association* 2000;**131**(7):887–99. [PUBMED: 10916327]

Featherstone 2004

Featherstone JD. The continuum of dental caries - evidence for a dynamic disease process. *Journal of Dental Research* 2004;**83** Spec No C:C39–42.

Fejerskov 2004

Fejerskov O. Changing paradigms in concepts on dental caries: consequences for oral health care. *Caries Research* 2004;**38**(3):182–91. [PUBMED: 15153687]

GRADEpro GDT 2015 [Computer program]

McMaster University (developed by Evidence Prime). GRADEpro GDT. Version accessed 11 May 2017. Hamilton (ON): McMaster University (developed by Evidence Prime), 2015.

Gupta 1993

Gupta S, Seth AK, Gupta A, Gavane AG. Transplacental passage of fluorides. *Journal of Pediatrics* 1993;**123**(1):139–41.

Helfenstein 1991

Helfenstein U, Steiner M, Marthaler TM. Caries prediction on the basis of past caries including precavity lesions. *Caries Research* 1991;**25**(5):372–6.

Higgins 2011

Higgins JP, Green S, editor(s). Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0 (updated March 2011). The Cochrane Collaboration, 2011. Available from handbook.cochrane.org.

Hiller 1975

Hiller CR, Robinson C, Weatherell JA. Variations in the composition of developing rat incisor enamel. *Calcified Tissue Research* 1975;**18**(1):1–12. [PUBMED: 1148889]

Hobdell 2003

Hobdell MH, Oliveira ER, Bautista R, Myburgh NG, Lalloo R, Narendran S, et al. Oral diseases and socio-economic status (SES). *British Dental Journal* 2003;**194**(2):91–6; discussion 88.

Iheozor-Ejiofor 2015

Iheozor-Ejiofor Z, Worthington HV, Walsh T, O'Malley L, Clarkson JE, Macey R, et al. Water fluoridation for the prevention of dental caries. *Cochrane Database of Systematic Reviews* 2015, Issue 6. [DOI: 10.1002/14651858.CD010856.pub2]

Ismail 1997

Ismail AI, Tanzer JM, Dingle JL. Current trends of sugar consumption in developing societies. *Community Dentistry and Oral Epidemiology* 1997;**25**(6):438–43.

Ismail 2008

Ismail AI, Hasson H. Fluoride supplements, dental caries and fluorosis: a systematic review. *Journal of the American Dental Association* 2008;**139**(11):1457–68.

Jankauskiene 2010

Jankauskiene B, Narbutaite J. Changes in oral health-related quality of life among children following dental treatment under general anaesthesia. A systematic review. *Stomatologija* 2010;**12**(2):60–4.

Jones 2017

Jones CM, Davies GM, Monaghan N, Morgan MZ, Neville JS, Pitts NB. The caries experience of 5 year-old children in Scotland in 2013–2014, and in England and Wales in 2014–2015. Reports of cross-sectional dental surveys using BASCD criteria. *Community Dental Health* 2017;**34**(3): 157–62. [PUBMED: 28872810]

Kailis 1968

Kailis DG, Taylor SR, Davis GB, Bartlett LG, Fitzgerald DJ, Grose IJ, et al. Fluoride and caries: observations on the effects of prenatal and postnatal fluoride on some Perth pre-school children. *Medical Journal of Australia* 1968;**2**(23): 1037–1040.

Kolker 2007

Kolker JL, Yuan Y, Burt BA, Sandretto AM, Sohn W, Lang SW, et al. Dental caries and dietary patterns in low-income African American children. *Pediatric Dentistry* 2007;**29**(6): 457–64.

Kraus 1965

Kraus BS, Jordan RE. *The Human Dentition Before Birth*. 1st Edition. Philadelphia: Lea & Febiger, 1965.

Levy 1999

Levy SM, Guha-Chowdhury N. Total fluoride intake and implications for dietary fluoride supplementation. *Journal of Public Health Dentistry* 1999;**59**(4):211–23. [PUBMED: 10682326]

Li 2002

Li Y, Wang W. Predicting caries in permanent teeth from caries in primary teeth: an eight-year cohort study. *Journal of Dental Research* 2002;**81**(8):561–6.

Limeback 1999

Limeback H. A re-examination of the pre-eruptive and post-eruptive mechanism of the anti-caries effects of fluoride: is there any anti-caries benefit from swallowing fluoride?. *Community Dentistry and Oral Epidemiology* 1999;**27**(1): 62–71.

Marinho 2003a

Marinho VCC, Higgins J, Logan S, Sheiham A (deceased). Fluoride toothpastes for preventing dental caries in children and adolescents. *Cochrane Database of Systematic Reviews* 2003, Issue 1. [DOI: 10.1002/14651858.CD002278]

Marinho 2003b

Marinho VCC, Higgins JPT, Logan S, Sheiham A (deceased). Topical fluoride (toothpastes, mouthrinses, gels or varnishes) for preventing dental caries in children and

adolescents. *Cochrane Database of Systematic Reviews* 2003, Issue 4. [DOI: 10.1002/14651858.CD002782]

Marinho 2013

Marinho VCC, Worthington HV, Walsh T, Clarkson JE. Fluoride varnishes for preventing dental caries in children and adolescents. *Cochrane Database of Systematic Reviews* 2013, Issue 7. [DOI: 10.1002/14651858.CD002279.pub2]

Marinho 2015

Marinho VCC, Worthington HV, Walsh T, Chong LY. Fluoride gels for preventing dental caries in children and adolescents. *Cochrane Database of Systematic Reviews* 2015, Issue 6. [DOI: 10.1002/14651858.CD002280.pub2]

Marinho 2016

Marinho VCC, Chong LY, Worthington HV, Walsh T. Fluoride mouthrinses for preventing dental caries in children and adolescents. *Cochrane Database of Systematic Reviews* 2016, Issue 7. [DOI: 10.1002/14651858.CD002284.pub2]

Petersen 2005

Petersen PE, Bourgeois D, Ogawa H, Estupinan-Day S, Ndiaye C. The global burden of oral diseases and risks to oral health. *Bulletin of the World Health Organization* 2005; **83**(9):661–9.

Prichard 1969

Prichard JL. The pre-natal and post-natal effects of fluoride supplements on West Australian schoolchildren, aged 6, 7, and 8, Perth, 1967. *Australian Dental Journal* 1969;**14**(5): 335–8. [PUBMED: 5264547]

Public Health England 2012

Public Health England. National Dental Epidemiology Programme for England: oral health survey of five-year-old children 2012. A report on the prevalence and severity of dental decay. www.nwph.net/dentalhealth/Oral%20Health%205yr%20old%20children%202012%20final%20report%20gateway (accessed 11 May 2017).

RevMan 2014 [Computer program]

Nordic Cochrane Centre, The Cochrane Collaboration. Review Manager 5 (RevMan 5). Version 5.3. Copenhagen: Nordic Cochrane Centre, The Cochrane Collaboration, 2014.

Rozier 2010

Rozier RG, Adair S, Graham F, Iafolla T, Kingman A, Kohn W, et al. Evidence-based clinical recommendations on the prescription of dietary fluoride supplements for caries prevention: a report of the American Dental Association Council on Scientific Affairs. *Journal of the American Dental Association* 2010;**141**(12):1480–9.

Sastry 2011

Sastry GM, Mohanty S, Bhongir AV, Mishra AK, Rao P. Association of higher maternal serum fluoride with adverse fetal outcomes. *International Journal of Medicine and Public Health* 2011;**1**(2):13–7.

Schünemann 2009

Schünemann HJ. GRADE: from grading the evidence to developing recommendations. A description of the system and a proposal regarding the transferability of the results of

- clinical research to clinical practice. *Zeitschrift für Evidenz, Fortbildung und Qualität im Gesundheitswesen* 2009;**103**(6): 391–400. [PUBMED: 19839216]
- Selwitz 2007**
Selwitz RH, Ismail AI, Pitts NB. Dental caries. *Lancet* 2007;**369**(9555):51–9.
- Seppa 1989**
Seppa L, Hausen H, Pöllänen L, Helasharju K, Kärkkäinen S. Past caries recordings made in Public Dental Clinics as predictors of caries prevalence in early adolescence. *Community Dentistry and Oral Epidemiology* 1989;**17**(6): 277–81.
- Stephen 1993**
Stephen KW. Systemic fluorides: drops and tablets. *Caries Research* 1993;**27 Suppl 1**:9–15.
- Tanimoto 2008**
Tanimoto K, Le T, Zhu L, Chen J, Featherstone JD, Li W, et al. Effects of fluoride on the interactions between amelogenin and apatite crystals. *Journal of Dental Research* 2008;**87**(1):39–44.
- Termine 1980**
Termine JD, Belcourt AB, Miyamoto MS, Conn KM. Properties of dissociatively extracted fetal tooth matrix proteins. II. Separation and purification of fetal bovine dentin phosphoprotein. *Journal of Biological Chemistry* 1980;**255**(20):9769–72. [PUBMED: 7430100]
- Toyama 2001**
Toyama Y, Nakagaki H, Kato S, Huang S, Mizutani Y, Kojima S, et al. Fluoride concentrations at and near the neonatal line in human deciduous tooth enamel obtained from a naturally fluoridated and a non-fluoridated area. *Archives of Oral Biology* 2001;**46**(2):147–53.
- Tubert-Jeannin 2011**
Tubert-Jeannin S, Auclair C, Amsallem E, Tramini P, Gerbaud L, Ruffieux C, et al. Fluoride supplements (tablets, drops, lozenges or chewing gums) for preventing dental caries in children. *Cochrane Database of Systematic Reviews* 2011, Issue 12. [DOI: 10.1002/14651858.CD007592.pub2]
- WHO 2009**
World Health Organization. Milk fluoridation for the prevention of dental caries. [www.who.int/oral/health/publications/milk fluoridation '2009' en.pdf?ua=1](http://www.who.int/oral/health/publications/milk%20fluoridation%2009/en.pdf?ua=1) (accessed 11 May 2017).
- Wong 2010**
Wong MCM, Glenny AM, Tsang BWK, Lo ECM, Worthington HV, Marinho VCC. Topical fluoride as a cause of dental fluorosis in children. *Cochrane Database of Systematic Reviews* 2010, Issue 1. [DOI: 10.1002/14651858.CD007693.pub2]
- * Indicates the major publication for the study

CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Leverett 1997

Methods	Trial design: RCT Location: Southern Maine, USA Duration: September 1983 to May 1985
Participants	1400 pregnant women in the first trimester recruited from communities served by fluoride-deficient drinking water 1175 born babies 938 children followed up at 3 years (intervention 464 versus control 484) 798 children followed up at 5 years (Intervention 398 versus control 400)
Interventions	Fluoride supplementation (tablets) versus placebo Group A: 1 dose of 2.2 mg sodium fluoride (NaF) (1 mg active fluoride ion), 1 tablet to be taken daily from 4th month of pregnancy Group B: placebo tablets (no fluoride) Both the intervention and control groups received fluoride drops from birth to 2 years of age and 1 0.5 mg tablet daily for children aged 2 to 3 years Follow-up: 5.5 years (6 months prenatal - 5 years after birth)
Outcomes	Caries experience between 2 groups, prevalence of "caries-free" children, mean dfs
Notes	Funding: National Institutes of Health/National Institute of Dental Research (NIH/NIDR) grants

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Quote: "...randomized on ID number into one-half containing fluoride and one-half containing placebo. The bottle labels did not distinguish between active drug and the placebo" Comment: the method of randomisation is not described in detail
Allocation concealment (selection bias)	Low risk	Quote: "The manufacturer of the fluoride products provided..." Comment: similar to central allocation, participants and personnel did not know their allocation before the intervention started

Leverett 1997 (Continued)

Blinding of participants and personnel (performance bias) All outcomes	Low risk	Quote: “The bottle labels did not distinguish between active drug and the placebo” Comment: participants and personnel did not know their allocation during the intervention
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	No description provided
Incomplete outcome data (attrition bias) All outcomes	High risk	3 years follow-up: 464/585 (79%) for the intervention group, 474/590 (80%) for placebo group 5 years follow-up: 398/585 (68%) for the intervention group, 400/590 (68%) for placebo group High attrition bias existed due to long-term follow-up
Selective reporting (reporting bias)	Unclear risk	No protocol or registry available
Other bias	High risk	Giving fluoride supplementation to both groups after delivery might affect the results

dfs: decayed, filled primary tooth surfaces; RCT: randomised controlled trial.

Characteristics of excluded studies [ordered by study ID]

Study	Reason for exclusion
Glenn 1979	771 participants were divided into 3 groups (control, 1.0 mg tablet per day, and 2.2 mg tablet per day). Allocation method was not described
Glenn 1982	33% of participants were randomly selected from 1374 private practice patients. Group 1: control, Group 2a: sibling of Group 1: no tablet, Group 2b: sibling of Group 1: with prenatal fluoride tablet, Group 3a: no prenatal intervention and no postnatal fluoride tablet, Group 3b: no prenatal intervention with postnatal fluoride tablet, Group 4a: twins with no prenatal intervention and no postnatal fluoride tablet, Group 4b: twins with prenatal fluoride tablet. Groups 3a and 3b are families with only 1 child. All participated patients were observed. It is cohort design
Glenn 1984	Participants were divided into 3 groups (control, postnatal fluoride tablet, and prenatal fluoride tablet). Allocation method was not described
Restrepo 1993	Narrative review

DATA AND ANALYSES

Comparison 1. Fluoride supplementation (tablets) versus placebo

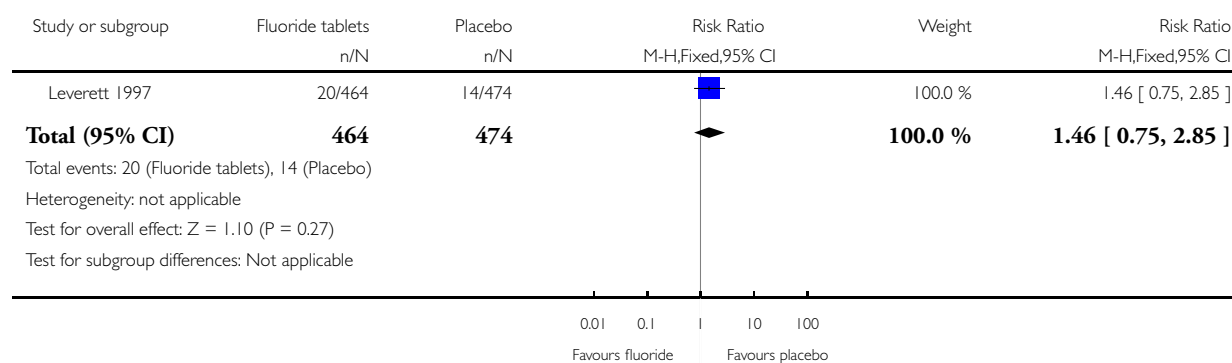
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Children with caries in the primary teeth at 3 years	1	938	Risk Ratio (M-H, Fixed, 95% CI)	1.46 [0.75, 2.85]
2 Children with caries in the primary teeth at 5 years	1	798	Risk Ratio (M-H, Fixed, 95% CI)	0.84 [0.53, 1.33]
3 Decayed primary surfaces at 3 years	1	938	Mean Difference (IV, Fixed, 95% CI)	0.05 [-0.02, 0.12]
4 Filled primary surfaces at 3 years	1	938	Mean Difference (IV, Fixed, 95% CI)	0.07 [-0.07, 0.21]
5 Decayed or filled primary surfaces at 3 years	1	938	Mean Difference (IV, Fixed, 95% CI)	0.12 [-0.05, 0.29]
6 Decayed primary surfaces at 5 years	1	798	Mean Difference (IV, Fixed, 95% CI)	-0.06 [-0.17, 0.05]
7 Filled primary surfaces at 5 years	1	798	Mean Difference (IV, Fixed, 95% CI)	0.03 [-0.28, 0.34]
8 Decayed or filled primary surfaces at 5 years	1	798	Mean Difference (IV, Fixed, 95% CI)	-0.05 [-0.42, 0.32]
9 Fluorosis (maxillary teeth) at 5 years	1	798	Risk Ratio (M-H, Fixed, 95% CI)	1.79 [0.95, 3.40]
10 Fluorosis (mandibular teeth) at 5 years	1	798	Risk Ratio (M-H, Fixed, 95% CI)	0.89 [0.35, 2.29]

Analysis 1.1. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 1 Children with caries in the primary teeth at 3 years.

Review: Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children

Comparison: 1 Fluoride supplementation (tablets) versus placebo

Outcome: 1 Children with caries in the primary teeth at 3 years



Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children (Review) 22

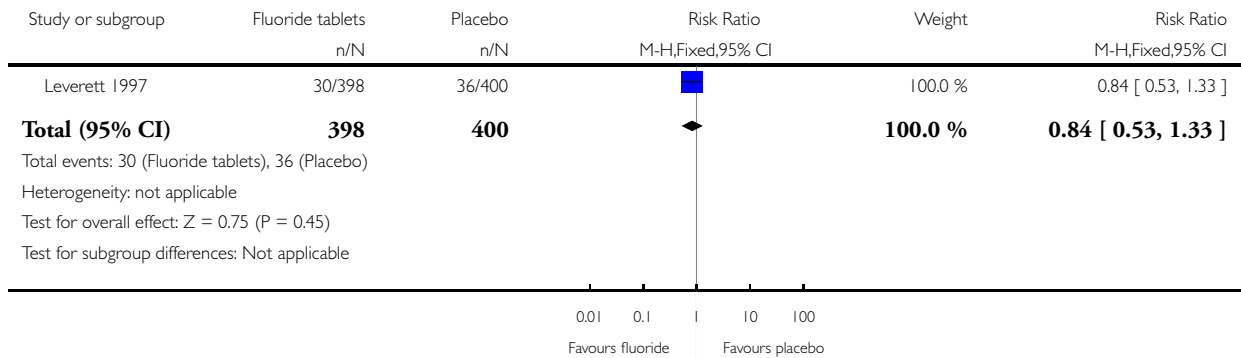
Copyright © 2017 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

Analysis 1.2. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 2 Children with caries in the primary teeth at 5 years.

Review: Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children

Comparison: 1 Fluoride supplementation (tablets) versus placebo

Outcome: 2 Children with caries in the primary teeth at 5 years

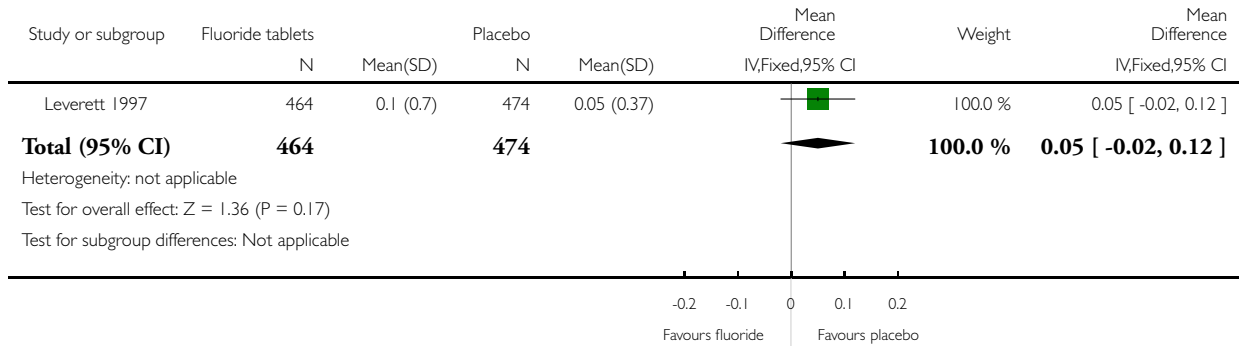


Analysis I.3. Comparison I Fluoride supplementation (tablets) versus placebo, Outcome 3 Decayed primary surfaces at 3 years.

Review: Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children

Comparison: I Fluoride supplementation (tablets) versus placebo

Outcome: 3 Decayed primary surfaces at 3 years

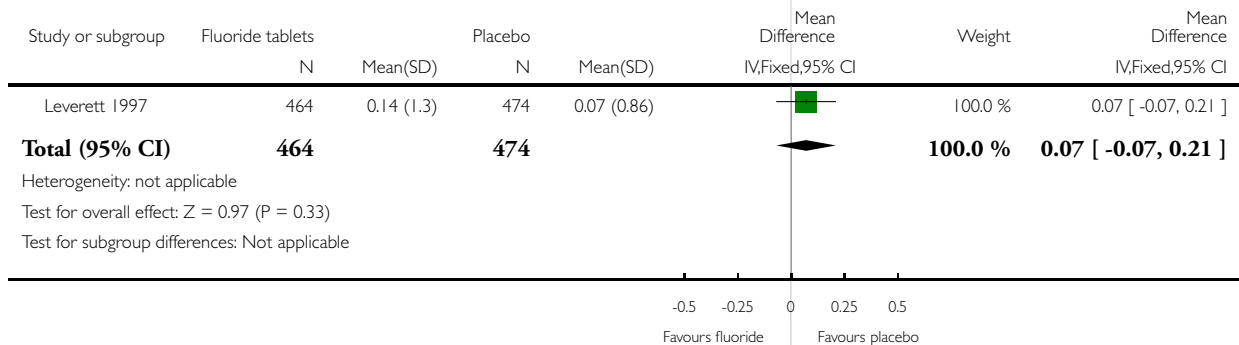


Analysis I.4. Comparison I Fluoride supplementation (tablets) versus placebo, Outcome 4 Filled primary surfaces at 3 years.

Review: Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children

Comparison: I Fluoride supplementation (tablets) versus placebo

Outcome: 4 Filled primary surfaces at 3 years

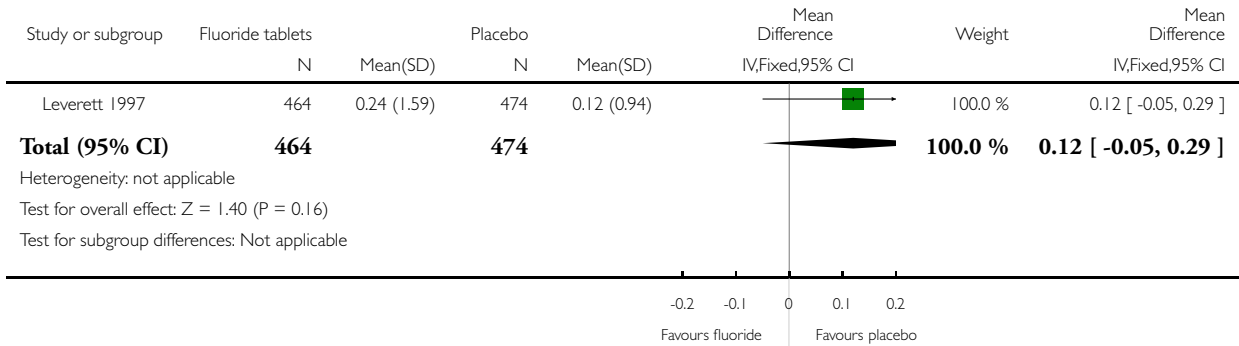


Analysis I.5. Comparison I Fluoride supplementation (tablets) versus placebo, Outcome 5 Decayed or filled primary surfaces at 3 years.

Review: Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children

Comparison: I Fluoride supplementation (tablets) versus placebo

Outcome: 5 Decayed or filled primary surfaces at 3 years

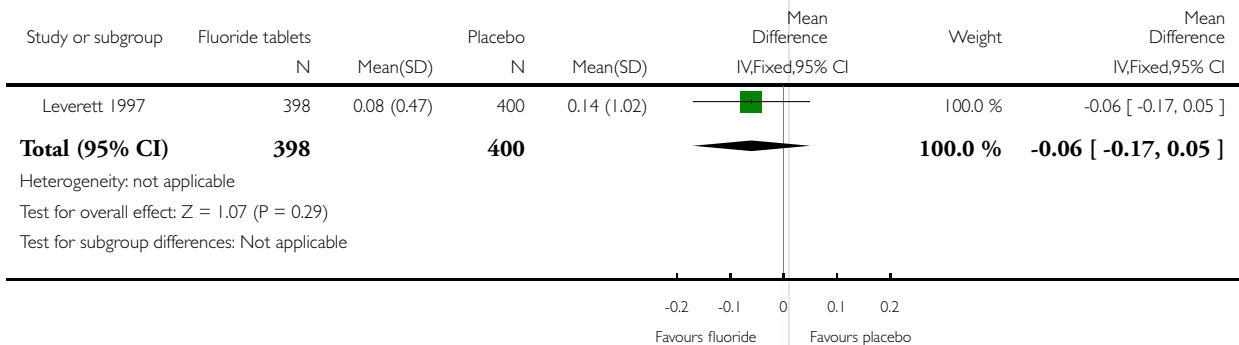


Analysis I.6. Comparison I Fluoride supplementation (tablets) versus placebo, Outcome 6 Decayed primary surfaces at 5 years.

Review: Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children

Comparison: I Fluoride supplementation (tablets) versus placebo

Outcome: 6 Decayed primary surfaces at 5 years

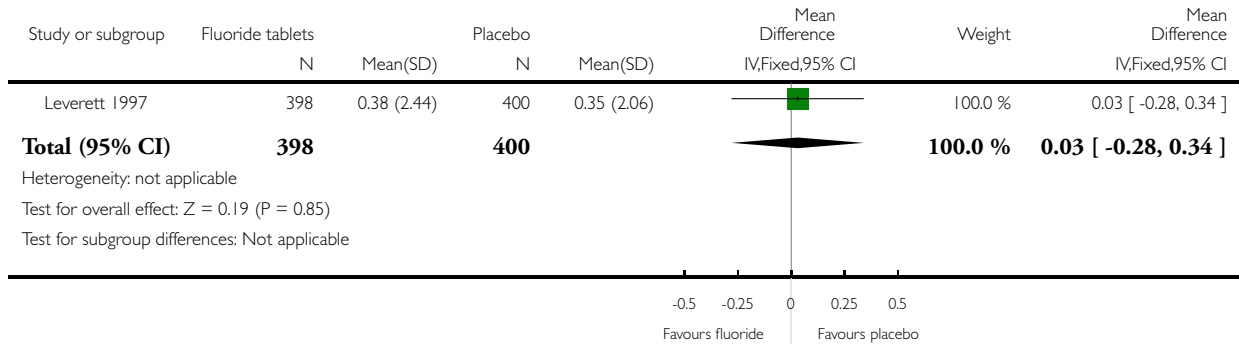


Analysis I.7. Comparison I Fluoride supplementation (tablets) versus placebo, Outcome 7 Filled primary surfaces at 5 years.

Review: Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children

Comparison: I Fluoride supplementation (tablets) versus placebo

Outcome: 7 Filled primary surfaces at 5 years

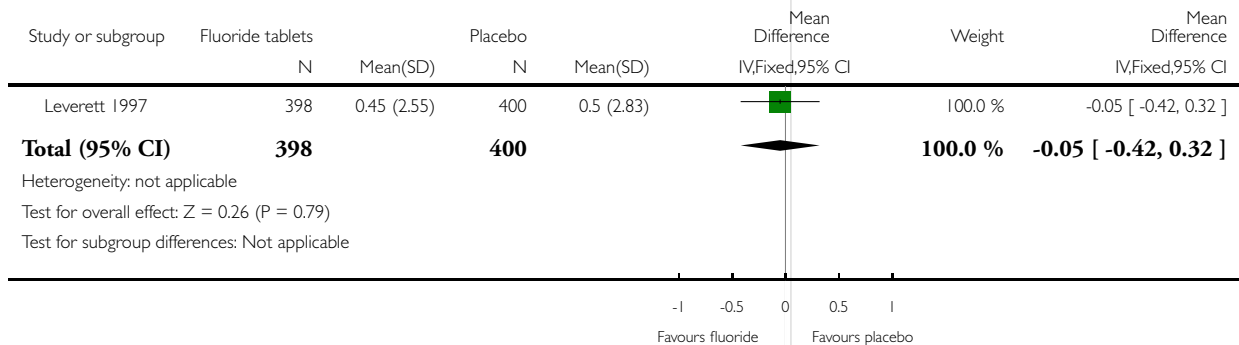


Analysis I.8. Comparison I Fluoride supplementation (tablets) versus placebo, Outcome 8 Decayed or filled primary surfaces at 5 years.

Review: Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children

Comparison: I Fluoride supplementation (tablets) versus placebo

Outcome: 8 Decayed or filled primary surfaces at 5 years

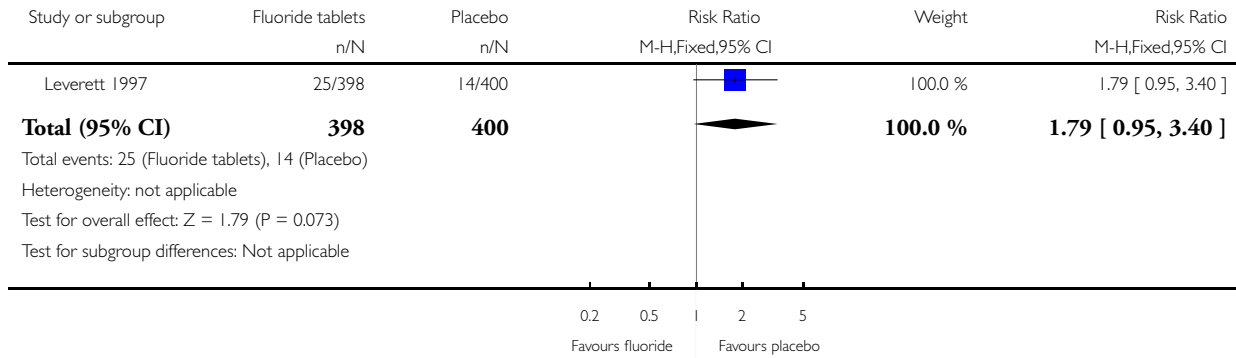


Analysis 1.9. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 9 Fluorosis (maxillary teeth) at 5 years.

Review: Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children

Comparison: 1 Fluoride supplementation (tablets) versus placebo

Outcome: 9 Fluorosis (maxillary teeth) at 5 years

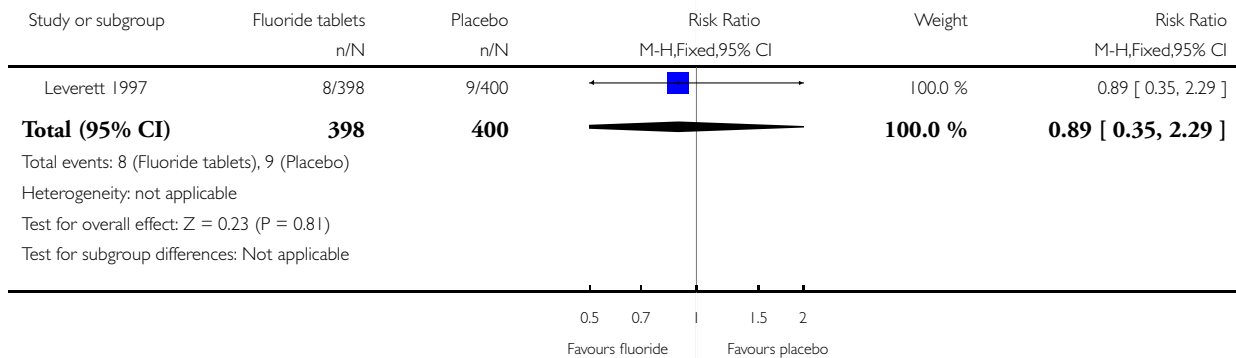


Analysis 1.10. Comparison 1 Fluoride supplementation (tablets) versus placebo, Outcome 10 Fluorosis (mandibular teeth) at 5 years.

Review: Fluoride supplementation (with tablets, drops, lozenges or chewing gum) in pregnant women for preventing dental caries in the primary teeth of their children

Comparison: 1 Fluoride supplementation (tablets) versus placebo

Outcome: 10 Fluorosis (mandibular teeth) at 5 years



APPENDICES

Appendix 1. Cochrane Oral Health's Trials Register search strategy

- 1 (pregnan* or prenatal:ti,ab) AND (INREGISTER)
- 2 ((expect* and mother*):ti,ab) AND (INREGISTER)
- 3 (#1 or #2) AND (INREGISTER)
- 4 ((fluorid* and (tablet* or drop* or lozenge* or pill* or gum* or supplement*))) AND (INREGISTER)
- 5 (#3 and #4) AND (INREGISTER)

Appendix 2. Cochrane Central Register of Controlled Clinical Trials (CENTRAL) search strategy

- #1 [mh Pregnancy]
- #2 (pregnan* or prenatal)
- #3 (expect* near/3 mother*)
- #4 #1 or #2 or #3
- #5 [mh Fluorides]
- #6 fluorid*
- #7 #5 or #6
- #8 (tablet* or drop* or lozenge* or pill* or gum* or supplement*)
- #9 #7 and #8
- #10 #4 and #9

Appendix 3. MEDLINE Ovid search strategy

- 1 exp Pregnancy/
- 2 (pregnan\$ or prenatal).ti,ab.
- 3 (expect\$ adj3 mother\$).ti,ab.
- 4 or/1-3
- 5 exp Fluorides/
- 6 fluorid\$.ti,ab.
- 7 or/5-6
- 8 (tablet\$ or drop\$ or lozenge\$ or pill\$ or gum\$ or supplement\$).ti,ab.
- 9 7 and 8
- 10 4 and 9

Appendix 4. Embase Ovid search strategy

- 1 exp Pregnancy/
- 2 (pregnan\$ or prenatal).ti,ab.
- 3 (expect\$ adj3 mother\$).ti,ab.
- 4 or/1-3
- 5 exp Fluorides/
- 6 fluorid\$.ti,ab.
- 7 or/5-6
- 8 (tablet\$ or drop\$ or lozenge\$ or pill\$ or gum\$ or supplement\$).ti,ab.
- 9 7 and 8
- 10 4 and 9

Appendix 5. LILACS BIREME Virtual Health Library (Latin American and Caribbean Health Science Information database) search strategy

((Mh pregnancy or pregnan\$ or embarazo or gravidez) AND (Mh Fluorides or fluor\$))
AND

(tablet\$ or pill\$ or drop\$ or supplement\$ or gum\$ or lozenge\$ or comprimido\$ or pilula\$ or suplemento\$ or goma\$ or pastilha\$ or pildora\$ or pastilla\$ or chicle\$)

Appendix 6. CINAHL EBSCO (Cumulative Index to Nursing and Allied Health Literature)

S10 S4 and S9

S9 S7 and S8

S8 (tablet* or drop* or lozenge* or pill* or gum* or supplement*)

S7 S5 or S6

S6 fluorid*

S5 (MH "Fluorides+")

S4 S1 or S2 or S3

S3 (expect* N3 mother*)

S2 (pregnan* or prenatal)

S1 (MH "Pregnancy+")

Appendix 7. US National Institutes of Health Ongoing Trials Register (ClinicalTrials.gov) search strategy

pregnancy and fluoride

Appendix 8. World Health Organization International Clinical Trials Registry Platform search strategy

pregnancy and fluoride

CONTRIBUTIONS OF AUTHORS

Erika Ota (EO) conceived and drafted the full review. Rena Takahashi (RT) and Keika Hoshi (KH) were responsible for the selection of studies. RT and EO were responsible for data extraction and management. KH, Yoshihiro Toyoshima (YT) and EO were responsible for the risk of bias assessment. KH, Hidemichi Yuasa (HY), YT and Rintaro Mori (RM) commented on and supervised the protocol and review. All review authors read and approved the final version.

DECLARATIONS OF INTEREST

Rena Takahashi: none known.

Erika Ota: none known.

Keika Hoshi: none known.

Toru Naito: none known.

Yoshihiro Toyoshima: none known.

Hidemichi Yuasa: none known.

Rintaro Mori: none known.

Eishu Nango: none known.

SOURCES OF SUPPORT

Internal sources

- National Center for Child Health and Development, Japan.
Grant 26A-5

External sources

- National Institute for Health Research (NIHR), UK.
This project was supported by the NIHR, via Cochrane Infrastructure funding to Cochrane Oral Health. The views and opinions expressed herein are those of the review authors and do not necessarily reflect those of the Systematic Reviews Programme, the NIHR, the NHS or the Department of Health

- Cochrane Oral Health Global Alliance, Other.
The production of Cochrane Oral Health reviews has been supported financially by our Global Alliance since 2011 (oralhealth.cochrane.org/partnerships-alliances). Contributors over the past year have been the American Association of Public Health Dentistry, USA; the British Association for the Study of Community Dentistry, UK; the British Society of Paediatric Dentistry, UK; the Canadian Dental Hygienists Association, Canada; the Centre for Dental Education and Research at All India Institute of Medical Sciences, India; the National Center for Dental Hygiene Research & Practice, USA; New York University College of Dentistry, USA; NHS Education for Scotland, UK; and the Swiss Society for Endodontology, Switzerland

- Ministry of Health Labour and Welfare, Japan.
Health Labour Sciences Research Grant (No. 13800128)
- School of Dentistry, The University of Manchester, UK.
- The Clinical Research Program for Child Health and Development from the Japan Agency for Medical Research and Development (AMED), Japan.

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

Fluorosis added as a primary outcome of the review. Adverse effects other than fluorosis added as secondary outcomes in the review.