Preventive zinc supplementation for children, and the effect of additional iron: a systematic review and meta-analysis

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Preventive zinc supplementation for children, and the effect of additional iron: a systematic review and meta-analysis

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ABSTRACT

Objective: Zinc deficiency is widespread, and preventive supplementation may have benefits in young children. Effects for children over 5 years of age, and effects when coadministered with other micronutrients are uncertain. These are obstacles to scale-up. This review seeks to determine if preventive supplementation reduces mortality and morbidity for children aged 6 months to 12 years.

Design: Systematic review conducted with the Cochrane Developmental, Psychosocial and Learning Problems Group. Two reviewers independently assessed studies. Meta-analyses were performed for mortality, illness and side effects.

Data sources: We searched multiple databases, including CENTRAL and MEDLINE in January 2013. Authors were contacted for missing information.

Eligibility criteria for selecting studies: Randomised trials of preventive zinc supplementation. Hospitalised children and children with chronic diseases were excluded.

Results: 80 randomised trials with 205 401 participants were included. There was a small but non-significant effect on all-cause mortality (risk ratio (RR) 0.95 (95% CI 0.86 to 1.05)). Supplementation may reduce incidence of all-cause diarrhoea (RR 0.87 (0.85 to 0.89)), but there was evidence of reporting bias. There was no evidence of an effect of incidence or prevalence of respiratory infections or malaria. There was moderate quality evidence of a very small effect on linear growth (standardised mean difference 0.09 (0.06 to 0.13)) and an increase in vomiting (RR 1.29 (1.14 to 1.46)). There was no evidence of an effect on iron status. Comparing zinc with and without iron cosupplementation and direct comparisons of zinc plus iron versus zinc administered alone favoured cointervention for some outcomes and zinc alone for other outcomes. Effects may be larger for children over 1 year of age, but most differences were not significant.

Conclusions: Benefits of preventive zinc supplementation may outweigh any potentially adverse effects in areas where risk of zinc deficiency is high. Further research should determine optimal intervention characteristics and delivery strategies.

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Strengths and limitations of this study

- This large review was conducted according to best practices and includes the highest quality current evidence about the effects of zinc supplementation.
- We investigated several outcomes and made multiple comparisons to explore the most important main effects and interactions.
- The analyses in this review could not identify the best way to deliver zinc supplements to children in need.

stored.[1] [2] In 2011, 116 000 deaths in children under 5 years were attributable to zinc deficiency (1.7% of mortalities in this group).[3]

Previous reviews have reached disparate conclusions about the benefits of zinc supplementation for young children,[4–12] and most have not examined evidence for children over 5 years of age. Zinc deficiency is prevalent in areas with other micronutrient deficiencies. Concerns about the administration of zinc with iron have been an obstacle to widespread delivery.[13] Understanding the effects of preventive zinc supplementation alone and with iron is crucially important to the future of global health policy.

To evaluate the effects of zinc with or without iron on illness and mortality, as well as growth, we analysed direct comparisons (ie, zinc plus iron vs zinc alone) as well as subgroups within an overall analysis.

METHODS
Selection criteria and search strategy
Following a published protocol,[14] we conducted a systematic review of randomised clinical trials of orally administered zinc compared with placebo and non-zinc cointerventions received by both groups (eg, vitamin A). We also compared zinc with and without iron cosupplementation. Participants were 6 months to 12 years of age. We excluded

INTRODUCTION
Regular dietary zinc intake is required because zinc cannot be produced or

studies of food fortification and children who were acutely ill.

We searched the African Index Medicus, CENTRAL, Conference Proceedings Citation Index, EMBASE, Global Health, ICTRP, IndMED, LILACS, MEDLINE, metaRegister of Controlled Trials, ProQuest Dissertations & Theses Database and WHOLIS in December 2012 and January 2013 (see online supplementary appendix 1). Reference lists from previous reviews and from included studies were examined, and trial authors were contacted for unpublished data. Two authors independently reviewed citations and extracted data, including participant demographics, details of the intervention, outcomes and risk of bias.[15]

Data synthesis
Relative risks and 95% CIs were calculated using Mantel-Haenszel methods. Standardised mean differences (SMDs) and 95% CIs were calculated for continuous measures using Hedges g and combined using inverse variance methods. When studies reported data in multiple formats, we calculated the SMD and its SE in Comprehensive Meta-Analysis (CMA) V.2 before entering data in Review Manager (RevMan) V.5.2. For incidence data, we combined risk ratios (RR) (events per child) and rate ratios (events per child year) because trials were relatively short and we did not anticipate interactions between the intervention and time at risk. For cluster-randomised trials, we used effects controlling for clustering, or we used a intracluster correlation coefficient to estimate robust SEs.[15] We used fixed-effect methods for all meta-analyses. Effects favour intervention when the relative risk is reduced (RR<1) or the standardised difference is positive (SMD>0).

When 10 or more studies reported an outcome, we conducted subgroup analyses to explore the effects of iron cosupplementation, national income (low-income countries compared with others), stunting, age (6–12, more than 12 months), dose (0–5, 5–10 mg, etc), duration (0–6, 6–12, more than 12 months) and formulation.

Quality of the evidence
Quality of the evidence was judged independently using GRADE.[16] The GRADE system rates evidence from each analysis (ie, pooled data where possible) as 'high', 'moderate', 'low' or 'very low'. A 'high' rating suggests that evidence is unlikely to be affected by further studies; a 'low' rating suggests that further research is required to confirm the direction and magnitude of the true effect. Ratings for meta-analyses of randomised controlled trials start at 'high' and may be downgraded for threats to internal validity (ie, within-study bias), inconsistency (ie, heterogeneity in results across studies), indirectness (eg, measures are proxies for the true outcome of interest), imprecision (eg, few participants, wide CIs) and reporting bias (ie, publication bias and selective outcome reporting). Because GRADE considers several domains in addition to internal validity, confidence in overall effects may be 'low' or 'very low' even when all studies are conducted rigorously. The following sections include significant and non-significant statistical results, and GRADE ratings in the text and tables provide further information about our confidence in these estimates.

RESULTS
Results of the search
From 6384 records, 80 studies were included (figure 1). Seventy-five studies were published in English, two each in Spanish and Portuguese and one in Chinese. Reasons for excluding 27 studies were enumerated (see online supplementary appendix 2); additionally, 11 ongoing studies were identified, and 5 studies could not be obtained. Seven included studies did not contribute to any meta-analysis because they did not report sufficient data (see online supplementary appendix 3).

Study characteristics
Included studies assigned 205 923 eligible participants (see online supplementary appendix 4). Twenty trials used factorial designs; there were 100 independent comparisons isolating zinc, and cointerventions were provided to both groups in 51 comparisons. There were eight independent comparisons of iron with zinc versus zinc alone including 1898 eligible participants. Sample sizes ranged from 21 to 72 438 eligible participants (median=200). Nine studies were cluster-randomised, including two randomising households. Three studies included 88% of participants.[17–19] Forty-six studies reported the mean baseline plasma or serum zinc concentration of their participants; the median of these mean concentrations was 72.5 µg/dL.

Thirty-two countries are represented; most studies were conducted in low-income or middle-income countries: 37 in Asia, 26 in Latin America and the Caribbean and 10 in sub-Saharan Africa. The median of mean age at baseline was 28 months, and 22 studies included children over 5 years of age. Stunted and non-stunted children were included in 42 studies; 5 included only stunted children, 5 included only non-stunted children and 28 did not specify if participants were stunted.

Studies provided zinc for less than 6 (30), 6–12 (33) and 12 months or more (16). Of those reporting frequency of zinc supplementation, 48 provided zinc daily and 11 provided zinc weekly. Where reported, daily dose was 0–5 (5), 5–10 (19), 10–15 (30), 15–20 (8) and 20 mg or more (12). Studies reporting the chemical compound of their zinc supplements provided zinc as sulfate (45), gluconate (12), acetate (6) and other compounds (8). Studies comparing zinc with iron versus zinc alone provided daily dose equivalents of 3–36 mg of iron. Outcomes were observed for about 26 weeks (median) after randomisation, with follow-up from 2–80 weeks.

Figure 1 PRISMA flow chart.
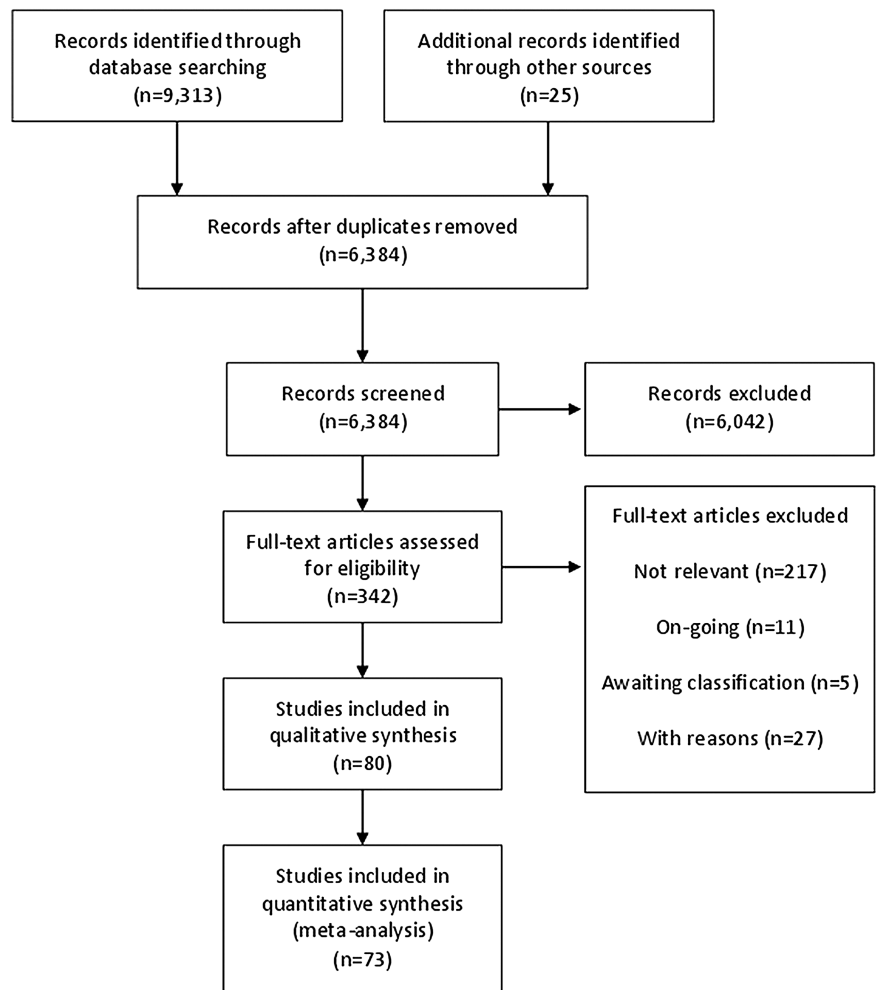
Risk of bias

Randomisation and allocation concealment were adequate in 34 and 32 studies; 46 and 48 studies were unclear (figure 2). For blinding of participants and personnel, 63 studies were at low risk of bias. For blinding of outcome assessment, 65 studies were at low risk of bias. For both types of blinding, 15 studies were unclear.

For all analyses, we attempted to include all randomised study participants; 47 studies were at low risk of bias for incomplete data, 31 were unclear and 2 were at high risk. For selective reporting, 3 studies were at low risk of bias, 44 were at unclear risk and 32 were at high risk (see online supplementary appendix 5).

Bias may affect secondary outcomes in this review, but it does not appear to be important for the primary outcome. For example, mortality and other objective measures are not vulnerable to bias related to blinding, and many missing outcomes were biomarkers or growth related.

Figure 2 Risk of bias summary.
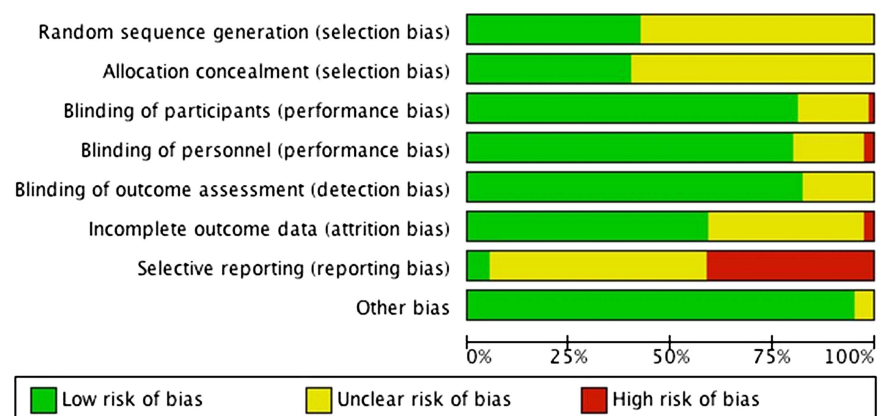
Table 1 Summary of findings

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	Number of participants (studies)	Quality of the evidence (GRADE)
	Assumed risk Control	Corresponding risk Zinc			
All-cause mortality Follow-up: 17–72 weeks	Low 2400/1 000 000 High 34 900/1 000 000	2280/1 000 000 (2064 to 2520) 33 155/1 000 000 (30 014 to 36 645)	RR 0.95 (0.86 to 1.05)	138 302 (13 studies)	⊕⊕⊕⊕ High
Mortality due to all-cause diarrhoea Follow-up: 52–69 weeks	Low 800/1 000 000 High 3000/1 000 000	760/1 000 000 (552 to 1048) 2850/1 000 000 (2070 to 3930)	RR 0.95 (0.69 to 1.31)	132 321 (4 studies)	⊕⊕⊕⊖ Moderate†
Mortality due to LRTI Follow-up: 52–69 weeks	Low 1200/1 000 000 High 3000/1 000 000	1032/1 000 000 (768 to 1380) 2580/1 000 000 (1920 to 3450)	RR 0.86 (0.64 to 1.15)	132 063 (3 studies)	⊕⊕⊕⊖ Moderate†
Mortality due to malaria Follow-up: 46–69 weeks	Low 7400/1 000 000 High 14 200/1 000 000	6660/1 000 000 (5698 to 7844) 12 780/1 000 000 (10 934 to 15 052)	RR 0.90 (0.77 to 1.06)	42 818 (2 studies)	⊕⊕⊕⊖ Moderate†
Incidence of all-cause diarrhoea Follow-up: 12–72 weeks	Low 20 000/1 000 000 High 1 770 000/1 000 000	17 400/1 000 000 (17 000 to 17 800) 1 539 900/1 000 000 (1 504 500 to 1 575 300)	RR 0.87 (0.85 to 0.89)	15 042 (35 studies)	⊕⊕⊖⊖ Low‡§
Incidence of LRTI Follow-up: 12–52 weeks	Low 30 000/1 000 000 High 370 000/1 000 000	30 000/1 000 000 (28 200 to 32 100) 370 000/1 000 000 (347 800 to 395 900)	RR 1.00 (0.94 to 1.07)	9610 (12 studies)	⊕⊕⊕⊕ High
Incidence of malaria Follow-up: 24–47 weeks	Low 140 000/1 000 000 High 2 950 000/1 000 000	147 000/1 000 000 (133 000 to 161 000) 3 097 500/1 000 000 (2 802 500 to 3 392 500)	RR 1.05 (0.95 to 1.15)	2407 (4 studies)	⊕⊕⊕⊖ Moderate¶
Height Follow-up: 10–60 weeks	The mean height in the control groups was −1 HAZ	The mean height in the intervention groups was 0.1 HAZ better (0 to 0.2 better)	SMD 0.09 (0.06 to 0.13)	13 669 (51 studies)	⊕⊕⊕⊖ Moderate**
Participants with one vomiting episode Follow-up: 24–52 weeks	Low 17 500/1 000 000 High 300 600/1 000 000	22 575/1 000 000 (19 950 to 25 550) 387 774/1 000 000 (342 684 to 438 876)	RR 1.29 (1.14 to 1.46)	35 192 (4 studies)	⊕⊕⊕⊕ High

GRADE working group grades of evidence.
High quality: further research is very unlikely to change our confidence in the estimate of effect.
Moderate quality: further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.
Low quality: further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.
Very low quality: we are very uncertain about the estimate.
*The basis for the assumed risk (eg, the median control group risk across studies) is provided in footnotes. The corresponding risk (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).
†Few deaths were observed overall.
‡I^2=88%.
§Trim-and-fill analysis suggests that the effect may be overestimated due to publication bias.
¶I^2=44%.
**I^2=86%.
LRTI, lower respiratory tract infection; SMD, standardised mean difference.

Effects of zinc supplementation

In addition to outcomes included in the summary of findings table (table 1), we analysed results for hospitalisation; prevalence of morbidities; additional measures of growth; as well as biological indicators of zinc, haemoglobin, iron and copper status (table 2). Subgroup analyses compared the effects of zinc supplementation with and without iron coadministration (table 3, see online supplementary appendix 6).

Fourteen studies including 138 302 participants were analysed for all-cause mortality, though other studies included no deaths in either group (figure 3), and there was high quality evidence of a small effect (risk ratio 0.95 (0.86 to 1.05)). There were similar effects for mortality due to diarrhoea (RR 0.95 (0.69 to 1.31)), mortality due to lower respiratory tract infection (LRTI; RR 0.86 (0.64 to 1.15)) and mortality due to malaria (RR 0.90 (0.77 to 1.06)), and the evidence for these outcomes was moderate quality.

In 25 studies including 15 042 participants, there was low quality evidence of a 13% reduction in incidence of all-cause diarrhoea (figure 4; RR 0.87 (0.85 to 0.89)). Other measures of diarrhoea were consistent with no difference or with a small reduction in morbidity, including prevalence of all-cause diarrhoea, hospitalisation due to all-cause diarrhoea, incidence of severe diarrhoea,

Table 2 Zinc compared with no zinc (all outcomes)

Outcomes	Trials	People	ES (95% CI), fixed effects	Heterogeneity I^2; χ^2 (p Value)
Zinc vs no zinc				
Mortality				
All-cause	13 (16%)	138 302 (67%)	Risk=0.95 (0.86 to 1.05)	0%; 10.57 (p=0.65)
Due to diarrhoea	4 (5%)	132 321 (64%)	Risk=0.95 (0.69 to 1.31)	0%; 0.82 (p=0.84)
Due to LRTI	3 (4%)	132 063 (64%)	Risk=0.86 (0.64 to 1.15)	0%; 0.07 (p=0.96)
Due to malaria	2 (3%)	42 818 (21%)	Risk=0.90 (0.77 to 1.06)	0%; 0.01 (p=0.94)
Hospitalisation				
All-cause	7 (9%)	92 872 (45%)	Risk=1.04 (0.97 to 1.11)	44%; 14.41 (p=0.07)
Due to diarrhoea	4 (5%)	74 039 (36%)	Risk=1.03 (0.87 to 1.22)	42%; 6.91 (p=0.14)
Due to LRTI	3 (4%)	74 743 (36%)	Risk=1.10 (0.93 to 1.30)	0%; 0.35 (p=0.95)
Diarrhoea				
Incidence (all-cause)	35 (44%)	15 042 (7%)	Risk=0.87 (0.85 to 0.89)	88%; 295.56 (p<0.00001)
Prevalence (all-cause)	13 (16%)	8519 (4%)	Rate=0.88 (0.86 to 0.90)	88%; 118.88 (p<0.00001)
Incidence (severe)	5 (6%)	4982 (2%)	Risk=0.89 (0.84 to 0.95)	56%; 13.54 (p=0.04)
Incidence (persistent)	7 (9%)	6216 (3%)	Risk=0.73 (0.62 to 0.85)	61%; 20.47 (p=0.009)
Prevalence (persistent)	1 (1%)	666 (0%)	Rate=0.70 (0.64 to 0.76)	91%; 11.76 (p=0.0006)
LRTI				
Incidence	12 (15%)	9610 (5%)	Risk=1.00 (0.94 to 1.07)	1%; 17.16 (p=0.44)
Prevalence	3 (4%)	1955 (1%)	Rate=1.20 (1.10 to 1.30)	97%; 89.87 (p<0.00001)
Malaria				
Incidence	4 (5%)	2407 (1%)	Risk=1.05 (0.95 to 1.15)	0%; 2.04 (p=0.84)
Prevalence	1 (1%)	661 (0%)	Rate=0.88 (0.47 to 1.64)	Not applicable
Growth				
Height	51 (64%)	13 669 (7%)	SMD=0.09 (0.06 to 0.13)	86%; 407.92 (p<0.00001)
Weight	44 (55%)	12 305 (6%)	SMD=0.10 (0.07 to 0.14)	76%; 216.64 (p<0.00001)
Weight-to-height ratio	24 (30%)	7901 (4%)	SMD=0.05 (0.01 to 0.10)	20%; 34.96 (p=0.17)
Prevalence of stunting	6 (8%)	3838 (2%)	Risk=0. 94 (0.86 to 1.02)	59%; 19.43 (p=0.01)
AEs				
Participants with one AE	2 (3%)	850 (0%)	SMD=1.13 (1.00 to 1.27)	0%; 0.49 (p=0.78)
Study withdrawal	6 (8%)	4263 (2%)	Risk=1.75 (0.93 to 3.32)	21%; 5.07 (p=0.28)
Vomiting (incidence)	5 (6%)	4095 (2%)	Risk=1.68 (1.61 to 1.75)	85%; 34.28 (p<0.00001)
Vomiting (prevalence)	4 (5%)	35 192 (17%)	Rate=1.29 (1.14 to 1.46)	37%; 6.31 (p=0.18)
Biological indicators				
Zn concentration	46 (58%)	9810 (5%)	SMD=0.62 (0.58 to 0.67)	91%; 582.45 (p<0.00001)
Zn deficiency (prevalence)	15 (19%)	5434 (3%)	Risk=0.49 (0.45 to 0.53)	86%; 144.77 (p<0.00001)
Haemoglobin concentration	27 (34%)	6024 (3%)	SMD=−0.05 (−0.10 to 0.00)	45%; 63.96 (p=0.002)
Anaemia (prevalence)	13 (16%)	4287 (2%)	Risk=1.00 (0.95 to 1.06)	37%; 28.52 (p=0.05)
Fe concentration	19 (24%)	4474 (2%)	SMD=0.07 (0.00 to 0.13)	95%; 480.50 (p<0.00001)
Fe deficiency (prevalence)	10 (13%)	3149 (2%)	Risk=0.99 (0.89 to 1.10)	15%; 16.44 (p=0.29)
Cu concentration	11 (14%)	3071 (1%)	SMD=−0.22 (−0.29 to 0.14)	68%; 37.47 (p=0.0002)
Cu deficiency (prevalence)	3 (4%)	1337 (1%)	Risk=2.64 (1.28 to 5.42)	59%; 4.94 (p=0.08)

AE, adverse event; ES, effect size; LRTI, lower respiratory tract infection; SMD, standardised mean difference.

Table 3 Subgroup analyses

Subgroup Mortality	Trials 13	People 138 302	Risk ratio (95% CI), fixed 0.95 (0.86 to 1.05)	I^2; χ^2 (p Value) 0%; 10.57 (p=0.65)
Iron cosupplementation (I^2=23%; χ^2=1.30, p=0.25)				
With iron	4	99 242	0.99 (0.86 to 1.15)	0%; 0.76 (p=0.86)
Without iron	11	64 985	0.89 (0.79 to 1.00)	0%; 9.99 (p=0.44)
Age (I^2=59.8%; χ^2=2.48, p=0.11)				
6 months to 1 year	6	29 879	1.06 (0.88 to 1.27)	0%; 2.56 (p=0.77)
1–5 years	8	125 903	0.89 (0.80 to 0.99)	12%; 10.28 (p=0.33)
Dose, mg (I^2=0%; χ^2=2.64, p=0.45)				
0–5	2	717	0.72 (0.08 to 6.47)	29%; 1.41 (p=0.23)
5–10	1	274	3.04 (0.32 to 28.90)	Not applicable
10–15	11	152 062	0.93 (0.84 to 1.02)	0%; 8.16 (p=0.61)
20 or more	1	2464	0.14 (0.01 to 2.78)	Not applicable
Duration, months (I^2=0%; χ^2=1.20, p=0.55)				
0–6	2	2817	0.59 (0.07 to 5.15)	47%; 1.88. (p=0.17.)
6–12	7	3898	0.68 (0.37 to 1.25)	4%; 6.23 (p=0.40)
12 or more	6	148 802	0.93 (0.85 to 1.03)	0%; 2.91 (p=0.71)
Formulation (I^2=0%; χ^2=0.54, p=0.91)				
Solution	5	3639	0.99 (0.25 to 3.91)	15%; 4.68 (p=0.32)
Pill/tablet	8	149 854	0.93 (0.85 to 1.02)	0%; 6.99 (p=0.43)
Capsule	1	306	0.51 (0.05 to 5.60)	Not applicable
Powder	1	1718	0.71 (0.27 to 1.86)	Not applicable
Incidence of diarrhoea	35	15 042	0.87 (0.85 to 0.89)	88%; 295.56 (p<0.00001)
Iron cosupplementation (I^2=99%; χ^2=65.11, p<0.00001)				
With iron	10	4299	1.00 (0.96 to 1.05)	76%; 37.33 (p<0.00001)
Without iron	22	11 344	0.82 (0.80 to 0.84)	87%; 196.27 (p<0.00001)
Age (I^2=0%; χ^2=0.32, p=0.85)				
6 months to 1 year	10	5576	0.88 (0.85 to 0.90)	95%; 252.46 (p<0.00001)
1–5 years	15	8370	0.87 (0.84 to 0.90)	43%; 31.48 (p=0.03)
5–13 years	1	842	0.90 (0.81 to 0.98)	Not applicable
Dose, mg (I^2=98%; χ^2=195.69, p<0.00001)				
0–5	4	1784	0.95 (0.89 to 1.01)	73%; 22.46 (p=0.001)
5–10	6	2630	0.73 (0.64 to 0.83)	67%; 15.32 (p=0.009)
10–15	11	5452	0.96 (0.92 to 0.99)	69%; 38.39 (p=0.0001)
15–20	2	477	0.61 (0.58 to 0.65)	0%; 0.21 (p<0.00001)
20 or more	6	4931	0.90 (0.87 to 0.94)	75%; 28.17 (p<0.00001)
Duration, months (I^2=0%; χ^2=1.15, p=0.56)				
0–6	7	4190	0.89 (0.85 to 0.93)	57%; 16.42 (p=0.02)
6–12	14	8971	0.86 (0.84 to 0.89)	93%; 250.92 (p<0.00001)
12 or more	5	1881	0.88 (0.82 to 0.95)	73%; 29.82 (p=0.0002)
Formulation (I^2=94%; χ^2=51.34, p<0.00001)				
Solution	19	10 768	0.84 (0.82 to 0.86)	90%; 236.48 (p<0.00001)
Pill/tablet	3	1696	0.90 (0.81 to 0.99)	5%; 3.15 (p=0.37)
Capsule	1	612	0.78 (0.60 to 1.01)	Not applicable
Powder	2	1861	1.04 (0.98 to 1.09)	0%; 0.65 (p=0.42)

prevalence of severe diarrhoea, incidence of persistent diarrhoea and prevalence of persistent diarrhoea.

In 12 trials (9610 participants), there was high-quality evidence of no effect on LRTI incidence (see online supplementary appendix 7; RR 1.00 (0.94 to 1.07)). One trial reported no LRTI in either group.[20] Results for prevalence were consistent with no difference in respiratory morbidity.

Four trials (2407 participants) found moderate quality evidence that would be consistent with no effect or a harmful effect on malaria incidence (RR 1.04 (0.94, 1.14)). One study reported no significant effect on malaria prevalence.

Fifty studies reported height for 13 669 participants (figure 5). There was moderate quality evidence of a very small but statistically significant increase in linear growth (SMD 0.09 (0.06 to 0.13)). Results for weight, weight-to-height ratio and prevalence of stunting were consistent with no difference or a small effect on growth.

Forty-six studies reported serum zinc for 9810 participants. There was evidence of a medium effect (SMD
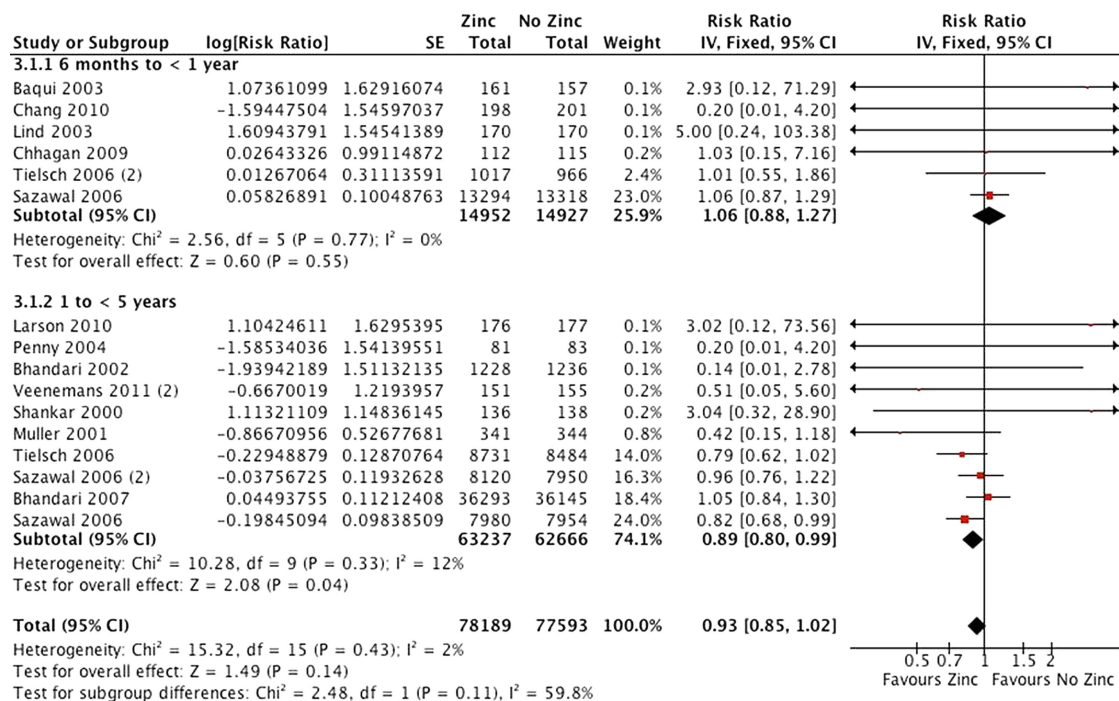
Figure 3 All-cause mortality by age.

0.62 (0.58 to 0.67)) on zinc concentration. Results consistently favoured zinc rather than no-intervention, but they were extremely inconsistent in magnitude, possibly due to differences in participants and settings (χ^2=582.45, df=47 (p<0.00001); I^2=91%). Eleven studies reported serum copper for 3071 participants (1% of participants in this review). There was very low-quality evidence of a small reduction in copper (SMD −0.22 (−0.29 to 0.14)); as above, the results were inconsistent (χ^2=37.47, df=10 (p<0.0002); I^2=68%). There was no evidence of an effect on haemoglobin, prevalence of anaemia or iron status.

In five trials (35 192participants), there was high-quality evidence of increased vomiting (RR 1.29 (1.14 to 1.46)). Two trials reported no adverse events in either group (ie, supplemented or non-supplemented).[21 22] Results for study withdrawal, participants with one or more side effects and number of vomiting episodes indicate some short-term side effects; there was no evidence of serious adverse events.

Effects of zinc plus iron compared with zinc alone

Effects on mortality were not significantly different between subgroups with and without iron (χ^2=1.30, p=0.25); however, there was no mortality effect in groups receiving iron (RR 0.99 (0.86 to 1.15)) while the effect for groups that did not receive iron was nearly significant (RR 0.89 (0.79 to 1.00)). Effects on the incidence of diarrhoea differed between groups (figure 4; χ^2=65.11, p<0.00001), with no benefit for the group that received iron (RR 1.00 (0.96 to 1.05)) and a significant benefit for the group that did not receive iron (RR 0.82 (0.80 to 0.84)). There were significant effects with and

without iron cosupplementation on zinc status; these were greater in studies without iron for serum zinc (χ^2=27.07, p<0.00001) and prevalence of zinc deficiency (χ^2=34.27, p<0.00001). There were also differences between these groups of studies for serum ferritin and serum copper; zinc had no effect in studies with iron cointervention, but zinc without iron cointervention reduced ferritin and copper. Overall effects on growth were small; there was a significant difference between subgroups for height but not weight, and the difference for weight-to-height ratio favoured the group that received iron (ie, the opposite of the other results). There were no significant effects in either subgroup for LRTIs, serum haemoglobin, prevalence of anaemia or prevalence of iron deficiency.

Several trials compared zinc coadministered with iron versus zinc given alone (see online supplementary appendix 6). One trial reported no significant difference in all-cause mortality (323 participants; RR 0.33 (0.01 to 8.39)). In five trials (1530 participants), effects on incidence of all-cause diarrhoea favoured zinc alone (RR 1.10 (1.03 to 1.18)). In one trial (399 participants), effects on prevalence of all-cause diarrhoea favoured zinc with iron, but this was not significant (RR 0.90 (0.79 to 1.06)). Five trials (1329 participants) reported no difference in height (SMD 0.06 (−0.04 to 0.16)). Similarly, there was low-quality evidence and mixed results for other outcomes (table 4).

Additional subgroup analyses

Studies in high-income countries did not evaluate most outcomes, so we were unable to explore differences in effect by national income. Effects on weight and

Study or Subgroup	log[Risk Ratio]	SE	Zinc Total	No Zinc Total	Weight	Risk Ratio IV, Fixed, 95% CI	Risk Ratio IV, Fixed, 95% CI
3.9.1 Iron co-intervention							
Meeks Gardner 2005	-4.48367263	1.42163731	55	59	0.0%	0.01 [0.00, 0.18]	
Cole 2012	-0.26503449	0.37388896	75	68	0.1%	0.77 [0.37, 1.60]	
Alarcon 2004	-0.33963305	0.30181636	112	111	0.1%	0.71 [0.39, 1.29]	
Rosado 1997 (2)	-0.52044108	0.18680745	55	54	0.4%	0.59 [0.41, 0.86]	
Veenemans 2011 (2)	-0.27010424	0.17803183	151	155	0.4%	0.76 [0.54, 1.08]	
Lind 2003 (2)	-0.07145896	0.09311961	170	170	1.4%	0.93 [0.78, 1.12]	
Chang 2010 (2)	-0.25350318	0.0794688	400	201	2.0%	0.78 [0.66, 0.91]	
Richard 2006 (2)	0.03619935	0.06581383	210	208	2.9%	1.04 [0.91, 1.18]	
Baqui 2003 (2)	0.08849308	0.05540904	162	165	4.0%	1.09 [0.98, 1.22]	
Soofi 2013	0.03774033	0.02689562	853	865	17.1%	1.04 [0.99, 1.09]	
Subtotal (95% CI)			**2243**	**2056**	**28.4%**	**1.00 [0.96, 1.05]**	
Heterogeneity: Chi² = 37.33, df = 9 (P < 0.0001); I² = 76%							
Test for overall effect: Z = 0.20 (P = 0.84)							
3.9.2 No iron co-intervention							
Chang 2010 (2)	-0.27537422	0.36843806	201	201	0.1%	0.76 [0.37, 1.56]	
Chang 2010	-0.58978851	0.33149677	198	201	0.1%	0.55 [0.29, 1.06]	
Han 2002	-0.75662302	0.28075433	33	22	0.2%	0.47 [0.27, 0.81]	
Han 2002 (2)	-0.17923468	0.27919905	24	26	0.2%	0.84 [0.48, 1.44]	
Meeks Gardner 1998	0.00421645	0.25610818	31	30	0.2%	1.00 [0.61, 1.66]	
Umeta 2000	-0.78170058	0.23234928	100	100	0.2%	0.46 [0.29, 0.72]	
Gupta 2003	-0.89381788	0.22473329	186	94	0.2%	0.41 [0.26, 0.64]	
Rosado 1997	-0.40188729	0.20280294	54	56	0.3%	0.67 [0.45, 1.00]	
Veenemans 2011	-0.22342699	0.19743482	153	153	0.3%	0.80 [0.54, 1.18]	
Penny 2004	-0.11778304	0.10486269	80	79	1.1%	0.89 [0.72, 1.09]	
Ruel 1997	-0.25131443	0.10417938	55	53	1.1%	0.78 [0.63, 0.95]	
Gupta 2007	-0.14660347	0.097381	854	858	1.3%	0.86 [0.71, 1.05]	
Wuehler 2008	-0.30010459	0.09601332	353	116	1.3%	0.74 [0.61, 0.89]	
Lind 2003	0.06899287	0.09065627	170	170	1.5%	1.07 [0.90, 1.28]	
Larson 2010	-0.23381808	0.08123116	176	177	1.9%	0.79 [0.68, 0.93]	
Muller 2001	-0.14077255	0.07595902	342	344	2.1%	0.87 [0.75, 1.01]	
Chhagan 2009	-0.04913269	0.07335707	104	105	2.3%	0.95 [0.82, 1.10]	
Richard 2006	-0.28106642	0.07078211	209	215	2.5%	0.75 [0.66, 0.87]	
Long 2006	0.12218449	0.07061765	181	183	2.5%	1.13 [0.98, 1.30]	
Long 2006 (2)	-0.26514098	0.06888881	192	180	2.6%	0.77 [0.67, 0.88]	
Rahman 2001 (2)	-0.05001042	0.06042171	175	160	3.4%	0.95 [0.84, 1.07]	
Rahman 2001	-0.13353139	0.0579042	170	161	3.7%	0.88 [0.78, 0.98]	
Baqui 2003	-0.02439145	0.05739701	161	157	3.8%	0.98 [0.87, 1.09]	
Sazawal 1996	-0.08461665	0.04515217	286	293	6.1%	0.92 [0.84, 1.00]	
Malik 2013	-0.48917615	0.02918358	134	124	14.5%	0.61 [0.58, 0.65]	
Bhandari 2002	-0.10583655	0.02613411	1228	1236	18.1%	0.90 [0.85, 0.95]	
Subtotal (95% CI)			**5850**	**5494**	**71.6%**	**0.82 [0.80, 0.84]**	
Heterogeneity: Chi² = 196.27, df = 25 (P < 0.00001); I² = 87%							
Test for overall effect: Z = 14.83 (P < 0.00001)							
Total (95% CI)			**8093**	**7550**	**100.0%**	**0.87 [0.85, 0.89]**	
Heterogeneity: Chi² = 298.70, df = 35 (P < 0.00001); I² = 88%							
Test for overall effect: Z = 12.45 (P < 0.00001)							
Test for subgroup differences: Chi² = 65.11, df = 1 (P < 0.00001), I² = 98.5%							

0.5 0.7 1 1.5 2
Favours Zinc Favours No Zinc

Figure 4 Incidence of all-cause diarrhoea with and without iron cosupplementation.

weight-to-height ratio were not statistically different, and there was no evidence of consistent differences in biological outcomes (see online supplementary appendix 6).
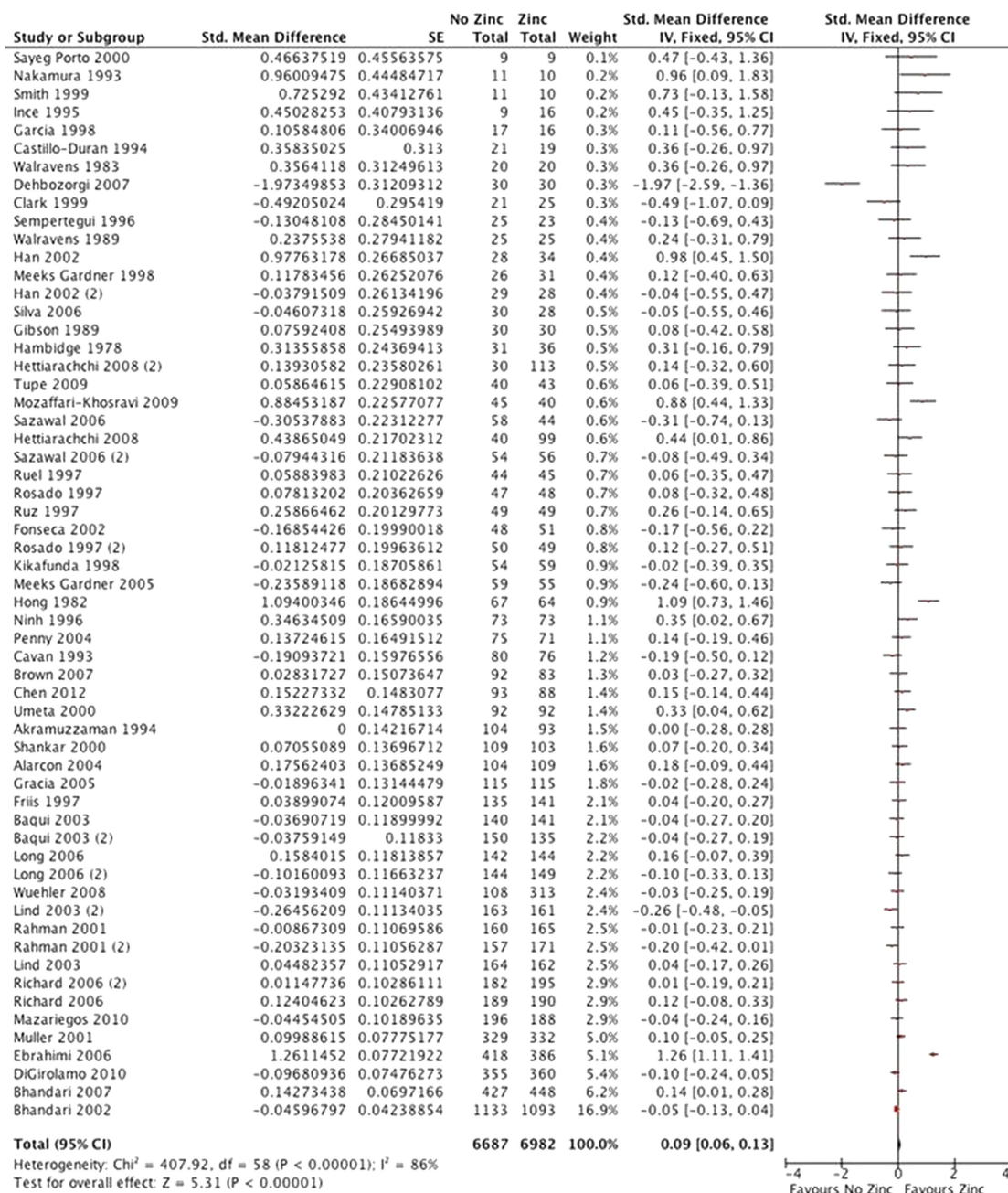
Most studies included stunted and non-stunted children, and it was not possible to compare effects between studies for most outcomes. Differences between groups were not significant for growth, but these would be consistent with larger effects in studies of stunted children.

Age was not significantly associated with effects on mortality or incidence diarrhoea, but results would be consistent with greater benefits in children over 1 year of age (figure 3). Effects on weight were greatest in studies of older children, and there was a similar pattern for height, though the largest study of children over 5 years of age included only 804 participants. The effect of supplementation on zinc deficiency was greater in studies of older children, as was the negative effect on

copper. There was no evidence of consistent differences in other biological outcomes.

Dose was not significantly associated with effects on mortality, incidence of LRTI, haemoglobin or weight-to-height ratio. The pattern of results was inconsistent for incidence and prevalence of diarrhoea, height, weight and plasma ferritin (see online supplementary appendix 6). Subgroups were significantly different for serum zinc, prevalence of zinc deficiency, prevalence of iron deficiency and plasma copper; only these results are consistent with a dose–response relationship.

Duration of supplementation was not significantly associated with effects on mortality, incidence of diarrhoea, incidence of LRTI, weight-to-height ratio or prevalence of iron deficiency (see online supplementary appendix 6). There was a significant difference for

Study or Subgroup	Std. Mean Difference	SE	No Zinc Total	Zinc Total	Weight	Std. Mean Difference IV, Fixed, 95% CI	Std. Mean Difference IV, Fixed, 95% CI
Sayeg Porto 2000	0.46637519	0.45563575	9	9	0.1%	0.47 [-0.43, 1.36]	
Nakamura 1993	0.96009475	0.44484717	11	10	0.2%	0.96 [0.09, 1.83]	
Smith 1999	0.725292	0.43412761	11	10	0.2%	0.73 [-0.13, 1.58]	
Ince 1995	0.45028253	0.40793136	9	16	0.2%	0.45 [-0.35, 1.25]	
Garcia 1998	0.10584806	0.34006946	17	16	0.3%	0.11 [-0.56, 0.77]	
Castillo-Duran 1994	0.35835025	0.313	21	19	0.3%	0.36 [-0.26, 0.97]	
Walravens 1983	0.3564118	0.31249613	20	20	0.3%	0.36 [-0.26, 0.97]	
Dehbozorgi 2007	-1.97349853	0.31209312	30	30	0.3%	-1.97 [-2.59, -1.36]	
Clark 1999	-0.49205024	0.295419	21	25	0.3%	-0.49 [-1.07, 0.09]	
Sempertegui 1996	-0.13048108	0.28450141	25	23	0.4%	-0.13 [-0.69, 0.43]	
Walravens 1989	0.2375538	0.27941182	25	25	0.4%	0.24 [-0.31, 0.79]	
Han 2002	0.97763178	0.26685037	28	34	0.4%	0.98 [0.45, 1.50]	
Meeks Gardner 1998	0.11783456	0.26252076	26	31	0.4%	0.12 [-0.40, 0.63]	
Han 2002 (2)	-0.03791509	0.26134196	29	28	0.4%	-0.04 [-0.55, 0.47]	
Silva 2006	-0.04607318	0.25926942	30	28	0.5%	-0.05 [-0.55, 0.46]	
Gibson 1989	0.07592408	0.25493989	30	30	0.5%	0.08 [-0.42, 0.58]	
Hambidge 1978	0.31355858	0.24369413	31	36	0.5%	0.31 [-0.16, 0.79]	
Hettiarachchi 2008 (2)	0.13930582	0.23580261	30	113	0.5%	0.14 [-0.32, 0.60]	
Tupe 2009	0.05864615	0.22908102	40	43	0.6%	0.06 [-0.39, 0.51]	
Mozaffari-Khosravi 2009	0.88453187	0.22577077	45	40	0.6%	0.88 [0.44, 1.33]	
Sazawal 2006	-0.30537883	0.22312277	58	44	0.6%	-0.31 [-0.74, 0.13]	
Hettiarachchi 2008	0.43865049	0.21702312	40	99	0.6%	0.44 [0.01, 0.86]	
Sazawal 2006 (2)	-0.07944316	0.21183638	54	56	0.7%	-0.08 [-0.49, 0.34]	
Ruel 1997	0.05883983	0.21022626	44	45	0.7%	0.06 [-0.35, 0.47]	
Rosado 1997	0.07813202	0.20362659	47	48	0.7%	0.08 [-0.32, 0.48]	
Ruz 1997	0.25866462	0.20129773	49	49	0.7%	0.26 [-0.14, 0.65]	
Fonseca 2002	-0.16854426	0.19990018	48	51	0.8%	-0.17 [-0.56, 0.22]	
Rosado 1997 (2)	0.11812477	0.19963612	50	49	0.8%	0.12 [-0.27, 0.51]	
Kikafunda 1998	-0.02125815	0.18705861	54	59	0.9%	-0.02 [-0.39, 0.35]	
Meeks Gardner 2005	-0.23589118	0.18682894	59	55	0.9%	-0.24 [-0.60, 0.13]	
Hong 1982	1.09400346	0.18644996	67	64	0.9%	1.09 [0.73, 1.46]	
Ninh 1996	0.34634509	0.16590035	73	73	1.1%	0.35 [0.02, 0.67]	
Penny 2004	0.13724615	0.16491512	75	71	1.1%	0.14 [-0.19, 0.46]	
Cavan 1993	-0.19093721	0.15976556	80	76	1.2%	-0.19 [-0.50, 0.12]	
Brown 2007	0.02831727	0.15073647	92	83	1.3%	0.03 [-0.27, 0.32]	
Chen 2012	0.15227332	0.1483077	93	88	1.4%	0.15 [-0.14, 0.44]	
Umeta 2000	0.33222629	0.14785133	92	92	1.4%	0.33 [0.04, 0.62]	
Akramuzzaman 1994	0	0.14216714	104	93	1.5%	0.00 [-0.28, 0.28]	
Shankar 2000	0.07055089	0.13696712	109	103	1.6%	0.07 [-0.20, 0.34]	
Alarcon 2004	0.17562403	0.13685249	104	109	1.6%	0.18 [-0.09, 0.44]	
Gracia 2005	-0.01896341	0.13144479	115	115	1.8%	-0.02 [-0.28, 0.24]	
Friis 1997	0.03899074	0.12009587	135	141	2.1%	0.04 [-0.20, 0.27]	
Baqui 2003	-0.03690719	0.11899992	140	141	2.1%	-0.04 [-0.27, 0.20]	
Baqui 2003 (2)	-0.03759149	0.11833	150	135	2.2%	-0.04 [-0.27, 0.19]	
Long 2006	0.1584015	0.11813857	142	144	2.2%	0.16 [-0.07, 0.39]	
Long 2006 (2)	-0.10160093	0.11663237	144	149	2.2%	-0.10 [-0.33, 0.13]	
Wuehler 2008	-0.03193409	0.11140371	108	313	2.4%	-0.03 [-0.25, 0.19]	
Lind 2003 (2)	-0.26456209	0.11134035	163	161	2.4%	-0.26 [-0.48, -0.05]	
Rahman 2001	-0.00867309	0.11069586	160	165	2.5%	-0.01 [-0.23, 0.21]	
Rahman 2001 (2)	-0.20323135	0.11056287	157	171	2.5%	-0.20 [-0.42, 0.01]	
Lind 2003	0.04482357	0.11052917	164	162	2.5%	0.04 [-0.17, 0.26]	
Richard 2006 (2)	0.01147736	0.10286111	182	195	2.9%	0.01 [-0.19, 0.21]	
Richard 2006	0.12404623	0.10262789	189	190	2.9%	0.12 [-0.08, 0.33]	
Mazariegos 2010	-0.04454505	0.10189635	196	188	2.9%	-0.04 [-0.24, 0.16]	
Muller 2001	0.09988615	0.07775177	329	332	5.0%	0.10 [-0.05, 0.25]	
Ebrahimi 2006	1.2611452	0.07721922	418	386	5.1%	1.26 [1.11, 1.41]	
DiGirolamo 2010	-0.09680936	0.07476273	355	360	5.4%	-0.10 [-0.24, 0.05]	
Bhandari 2007	0.14273438	0.0697166	427	448	6.2%	0.14 [0.01, 0.28]	
Bhandari 2002	-0.04596797	0.04238854	1133	1093	16.9%	-0.05 [-0.13, 0.04]	
Total (95% CI)			**6687**	**6982**	**100.0%**	**0.09 [0.06, 0.13]**	

Heterogeneity: Chi² = 407.92, df = 58 (P < 0.00001); I² = 86%
Test for overall effect: Z = 5.31 (P < 0.00001)

Favours No Zinc   Favours Zinc

Figure 5 Height.

prevalence of diarrhoea, but the magnitude of this difference may not be important. Studies of longer supplementation were associated with greater effects on height; the pattern of results was not consistent for weight. By contrast, the largest benefits for biological markers (serum zinc and prevalence of zinc deficiency) were reported in the shortest studies.

Formulation was associated with differences among subgroups, though few studies included capsules or powder. When comparing solution and tablets, differences were not significant for mortality, incidence and prevalence of diarrhoea; incidence of LRTI; blood haemoglobin; prevalence of anaemia or prevalence of iron deficiency. There were significant differences in the

effects of serum ferritin and serum copper, but only three studies of each outcome used tablets, and they were highly heterogeneous. The effects on height, weight and serum zinc were greater in studies using solution compared with tablet, but all effects were small (see online supplementary appendix 6).

Reporting bias
For outcomes included in the summary of findings table with 10 or more studies, we also conducted a trim-and-fill analysis to investigate reporting bias (see online supplementary appendix 8).[23] There was some evidence of small study bias—studies were trimmed for all-cause mortality (1 trimmed) and incidence of

Table 4 Zinc with iron compared with zinc alone (all outcomes)

Outcomes	Trials	People	ES (95% CI), fixed effects	Heterogeneity I^2; χ^2 (p Value)
All-cause mortality	1 (13%)	323 (17%)	Risk=0.33 (0.01 to 8.31)	Not applicable
Hospitalisation				
All-cause	1 (13%)	399 (21%)	Risk=0.92 (0.45 to 1.89)	Not applicable
Due to diarrhoea	1 (13%)	399 (21%)	Risk=0.99 (0.25 to 3.88)	Not applicable
Diarrhoea				
Incidence (all-cause)	5 (63%)	1530 (81%)	Risk=1.10 (1.03 to 1.18)	76%; 16.92 (p=0.002)
Prevalence (all-cause)	1 (13%)	399 (21%)	Rate=0.90 (0.79 to 1.06)	Not applicable
Incidence (severe)	1 (13%)	323 (17%)	Rate=0.78 (0.59 to 1.04)	Not applicable
Lower respiratory tract infection				
Incidence	3 (38%)	1065 (56%)	Risk=0.93 (0.83 to 1.04)	21%; 2.52 (p=0.28)
Malaria				
Incidence	1 (13%)	410 (22%)	Rate=0.86 (0.59 to 1.24)	Not applicable
Growth				
Height	5 (63%)	1517 (80%)	SMD=0.06 (−0.04 to 0.16)	0%; 3.54 (p=0.47)
Weight	4 (50%)	910 (48%)	SMD=0.12 (−0.01 to 0.25)	0%; 2.29 (p=0.51)
Weight-to-height ratio	4 (50%)	514 (27%)	SMD=−0.06 (−0.07 to 0.19)	0%; 1.36 (p=0.71)
Prevalence of stunting	2 (25%)	462 (24%)	Risk=0.92 (0.85 to 0.99)	45%; 1.82 (p=0.18)
Adverse events				
Study withdrawal	2 (25%)	557 (29%)	Risk=1.41 (0.91 to 2.18)	0%; 0.08 (p=0.78)
Biological indicators				
Zn concentration	8 (100%)	1337 (70%)	SMD=0.16 (0.05 to 0.27)	61%; 17.84 (p=0.01)
Zn deficiency (prevalence)	3 (38%)	350 (18%)	Risk=1.42 (0.75 to 2.68)	5%; 2.10 (p=0.35)
Haemoglobin concentration	8 (100%)	1341 (71%)	SMD=−0.23 (−0.34 to −0.12)	79%; 33.53 (p<0.0001)
Anaemia (prevalence)	3 (38%)	482 (25%)	Risk=0.78 (0.67 to 0.92)	0%; 1.25 (p=0.54)
Fe concentration	6 (75%)	945 (50%)	SMD=−1.79 (−1.99 to −1.56)	99%; 927.92 (p<0.00001)
Fe deficiency (prevalence)	2 (25%)	248 (13%)	Risk=0.12 (0.04 to 0.32)	87%; 8.00 (p=0.005)
Cu concentration	2 (25%)	353 (19%)	SMD=−0.06 (−0.27 to 0.15)	0%; 0.11 (p=0.74)

Effects favour intervention (ie, zinc rather than iron; zinc plus iron rather than zinc alone) when the relative risk is reduced (RR<1) or the standardised difference is positive (SMD>0).
ES, effect size; Rate, rate ratio; Risk, risk ratio; SMD, standardised mean difference.

all-cause diarrhoea (13 trimmed; figure 6). None were trimmed for incidence of LRTI; nor were any trimmed for height. The adjusted effect for mortality was not importantly different from the observed effect, but the observed effect for diarrhoea (RR 0.87 (0.85 to 0.89)) was larger than the adjusted value (RR 0.95 (0.93 to 0.97)).

DISCUSSION
Consistent with previous reviews, this review finds high-quality evidence from several large, well-conducted trials.[5 7 10] We believe that these results suggest zinc supplementation is probably associated with a small reduction in all-cause mortality for children at risk of deficiency. In interpreting these results, we considered that the results of this meta-analysis are drawn from 13 trials including almost 140 000 participants. The results of those studies are statistically consistent, the overall CIs are relatively small and the balance of probability favours zinc supplementation rather than placebo. Small reductions in cause-specific mortality were consistent with effects on illness and cause-specific mortality, and the results were biologically plausible. Benefits in any specific area may be related to the level of deficiency;

countries with very high levels of deficiency could expect the largest reductions in mortality as a result of supplementation.[24] This review also suggests that benefits may not be restricted to young children; there is some evidence of benefits on secondary outcomes in trials including children over 5 years of age, but there is a lack of evidence about the effects on mortality in this group.

Results for secondary outcomes suggest modest benefits. The main results for diarrhoea morbidity were consistent with previous reviews,[4 5 7 10] but an asymmetrical funnel plot was indicative of small-study bias. After adjustment, the effect for diarrhoea was halved, and the reduced estimate was consistent with other critical outcomes in this review. Previous reviews have suggested beneficial effects on respiratory infections[4 5 9–12] and malaria,[10] which this review does not confirm, and also reported variable effects on growth[5 6 8]; this review suggests that preventive zinc supplementation alone is unlikely to have large effects on linear growth and morbidity. Supplementation is associated with increased risk of vomiting, but there is no evidence of lasting adverse effects.

Critical outcomes included data for 2407–138 302 participants, so further placebo-controlled trials of
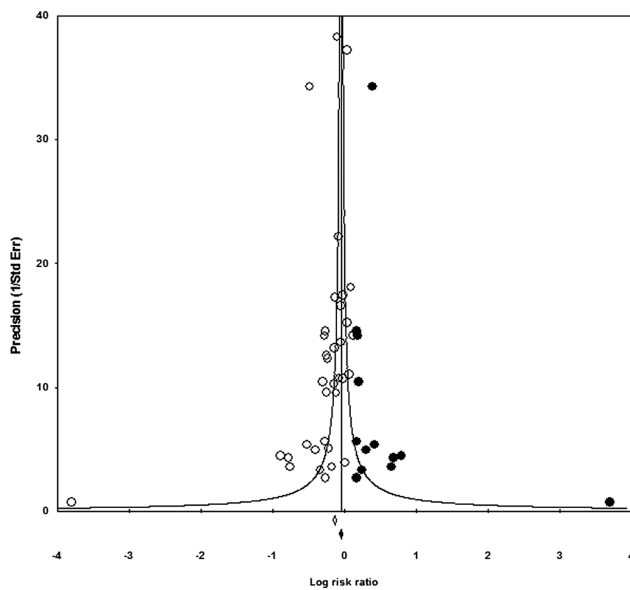
Figure 6 Incidence of diarrhoea funnel plot (trim-and-fill analysis).

preventive zinc supplementation for young children may not be necessary. However, subgroup analyses did not identify an optimal supplementation strategy (ie, dose, formulation and frequency), and large trials comparing active interventions could inform clinical guidelines. Subgroup analyses identify some sources of observed heterogeneity; however, subgroups that were statistically different included a large amount of residual heterogeneity, which is reflected in our judgements about the quality of the evidence (table 1). Analyses of group-level data are of limited value for identifying moderators, particularly in analyses dominated by a few large studies. Further analyses of individual patient data would be more conclusive.

Effects on biological indicators were inconsistent across studies, but large effects on these measures were not always reflected in clinical outcomes. Supplementation may increase serum zinc, but the magnitude of the effect appears to differ across populations and interventions. Effects on other micronutrients, including iron and copper, are uncertain. Researchers have suggested that iron supplementation may interfere with the absorption of zinc and, conversely, that zinc may interfere with iron and copper absorption[25 26]; however, the relationships between these biomarkers and clinical outcomes (ie, mortality and morbidities) have not been established.

Subgroup analyses comparing zinc with and without iron did not resolve uncertainty about the effect of cosupplementation. Only four studies with iron cosupplementation reported mortality outcomes, and evidence of outcome reporting bias for diarrhoea incidence leads to cautious interpretation of differences in this outcome. There was no evidence that larger doses or increased duration was associated with increased iron deficiency, but

these comparisons are observational and could be affected by uncontrolled covariates.

Direct comparisons within trials provide the only experimental evidence about the effects of cosupplementation with iron. For rare events like mortality, effects of zinc and iron can only be detected in large studies, so studies of interaction effects will need to be very large to detect real differences. Future studies are needed to identify the main effects and to explore how administration (ie, separate or combined) affects uptake and costs.

Dietary intake and supplementation have reduced micronutrient deficiencies in Asia, but micronutrient deficiencies remain common.[3 27] The prevalence of micronutrient deficiencies is declining in Africa, but the absolute number of deficient children is increasing.[3] This review suggests that the overall benefits of preventive zinc supplementation outweigh potential harms in areas with a high risk of zinc deficiency. Further research is needed to determine if these benefits extend to children over 5 years of age. Current estimates suggest that delivering 10 evidence-based nutrition-specific interventions, including preventive zinc supplements, could reduce global mortality in children under 5 years of age by 15%.[28] To that end, research is needed to identify the most effective strategies for delivering zinc supplements to populations in need.[29]

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REFERENCES

1. Hotz C, Brown K; International Zinc Nutrition Consultative Group (IZiNCG). Assessment of the risk of zinc deficiency in populations and options for its control. *Food Nutr Bull* 2004;25(1 Suppl 2):S94–203.
2. Maggini S, Wenzlaff S, Hornig D. Essential role of vitamin C and zinc in child immunity and health. *J Int Med Res* 2010;38:386–414.
3. Black RE, Victora CG, Walker SP, *et al*. Maternal and child undernutrition and overweight in low-income and middle-income countries. *Lancet* 2013;382:427–51.
4. Aggarwal R, Sentz J, Miller MA. Role of zinc administration in prevention of childhood diarrhea and respiratory illnesses: a meta-analysis. *Pediatrics* 2007;119:1120–30.
5. Brown KH, Peerson JM, Baker SK, *et al*. Preventive zinc supplementation among infants, preschoolers, and older prepubertal children. *Food Nutr Bull* 2009;30:S12–40.
6. Imdad A, Bhutta ZA. Effect of preventive zinc supplementation on linear growth in children under 5 years of age in developing countries: a meta-analysis of studies for input to the lives saved tool. *BMC Public Health* 2011;11(Suppl 3):S22.
7. Patel AB, Mamtani M, Badhoniya N, *et al*. What zinc supplementation does and does not achieve in diarrhea prevention: a systematic review and meta-analysis. *BMC Infect Dis* 2011;11:122.
8. Ramakrishnan U, Nguyen P, Martorell R. Effects of micronutrients on growth of children under 5 y of age: meta-analyses of single and multiple nutrient interventions. *Am J Clin Nutr* 2009;89:191–203.
9. Roth DE, Richard SA, Black RE. Zinc supplementation for the prevention of acute lower respiratory infection in children in developing countries: meta-analysis and meta-regression of randomized trials. *Int J Epidemiol* 2010;39:795–808.
10. Yakoob MY, Theodoratou E, Jabeen A, *et al*. Preventive zinc supplementation in developing countries: impact on mortality and morbidity due to diarrhea, pneumonia and malaria. (Special issue: technical inputs, enhancements and applications of the Lives Saved Tool (LiST)). *BMC Public Health* 2011;11(Suppl 3):S23.
11. Bhutta ZA, Black RE, Brown KH, *et al*. Prevention of diarrhea and pneumonia by zinc supplementation in children in developing countries: pooled analysis of randomized controlled trials. Zinc Investigators' Collaborative Group. *J Pediatr* 1999;135:689–97.
12. Lassi ZS, Haider BA, Bhutta ZA. Zinc supplementation for the prevention of pneumonia in children aged 2 months to 59 months. *Cochrane Database Syst Rev* 2010;(12):CD005978.
13. Pasricha S-R, Hayes E, Kalumba K, *et al*. Effect of daily iron supplementation on health in children aged 4–23 months: a systematic review and meta-analysis of randomised controlled trials. *Lancet Global Health* 2013;1:e77–86.
14. Junior JA, Dean S, Mayo-Wilson E, *et al*. Zinc supplementation for preventing mortality and morbidity, and promoting growth, in children aged 6 months to 12 years of age (protocol). *Cochrane Database Syst Rev* 2011;10:CD009384.
15. Higgins JP, Green S. *Cochrane handbook for systematic reviews of interventions*. Version 5.1.0 [updated March 2011]. The Cochrane Collaboration, 2011.
16. Balshem H, Helfand M, Schunemann HJ, *et al*. GRADE guidelines: 3. Rating the quality of evidence. *J Clin Epidemiol* 2011;64:401–6.
17. Bhandari N, Taneja S, Mazumder S, *et al*. Adding zinc to supplemental iron and folic acid does not affect mortality and severe morbidity in young children. *J Nutr* 2007;137:112–17.
18. Sazawal S, Black RE, Ramsan M, *et al*. Effects of routine prophylactic supplementation with iron and folic acid on admission to hospital and mortality in preschool children in a high malaria transmission setting: community-based, randomised, placebo-controlled trial. [Erratum appears in *Lancet* 2006;367:302]. *Lancet* 2006;367:133–43.
19. Tielsch JM, Khatry SK, Stoltzfus RJ, *et al*. Effect of routine prophylactic supplementation with iron and folic acid on preschool child mortality in southern Nepal: community-based, cluster-randomised, placebo-controlled trial. *Lancet* 2006;367:144–52.
20. Sempertegui F, Estrella B, Correa E, *et al*. Effects of short-term zinc supplementation on cellular immunity, respiratory symptoms, and growth of malnourished Equadorian children. *Eur J Clin Nutr* 1996;50:42–6.
21. Alarcon K, Kolsteren PW, Prada AM, *et al*. Effects of separate delivery of zinc or zinc and vitamin A on hemoglobin response, growth, and diarrhea in young Peruvian children receiving iron therapy for anemia. *Am J Clin Nutr* 2004;80:1276–82.
22. Mazariegos M, Hambidge KM, Westcott JE, *et al*. Neither a zinc supplement nor phytate-reduced maize nor their combination enhance growth of 6- to 12-month-old Guatemalan infants. *J Nutr* 2010;140:1041–8.
23. Duval S, Tweedie R. Trim and fill: a simple funnel-plot-based method of testing and adjusting for publication bias in meta-analysis. *Biometrics* 2000;56:455–63.
24. Wessells KR, Brown KH. Estimating the global prevalence of zinc deficiency: results based on zinc availability in national food supplies and the prevalence of stunting. *PLoS ONE* 2012;7:e50568.
25. Maret W, Sandstead HH. Zinc requirements and the risks and benefits of zinc supplementation. *J Trace Elem Med Biol* 2006;20:3–18.
26. Sandstrom B. Micronutrient interactions: effects on absorption and bioavailability. *Br J Nutr* 2001;85(Suppl 2):S181–5.
27. Ram U, Jha P, Ram F, *et al*. Neonatal, 1–59 month, and under-5 mortality in 597 Indian districts, 2001 to 2012: estimates from national demographic and mortality surveys. *Lancet Global Health* 2013;1:e219–e226.
28. Bhutta ZA, Das JK, Rizvi A, *et al*. Evidence-based interventions for improvement of maternal and child nutrition: what can be done and at what cost? *Lancet* 2013;382:452–77.
29. Wazny K, Zipursky A, Black R, *et al*. Setting research priorities to reduce mortality and morbidity of childhood diarrhoeal disease in the next 15 years. *PLoS Med* 2013;10:e1001446.