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Interventions to improve water quality for preventing diarrhoea (Review)

Clasen TF, Alexander KT, Sinclair D, Boisson S, Peletz R, Chang HH, Majorin F, Cairncross S

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[Intervention Review]

Interventions to improve water quality for preventing diarrhoea

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ABSTRACT

Background

Diarrhoea is a major cause of death and disease, especially among young children in low-income countries. In these settings, many infectious agents associated with diarrhoea are spread through water contaminated with faeces.

In remote and low-income settings, source-based water quality improvement includes providing protected groundwater (springs, wells, and bore holes), or harvested rainwater as an alternative to surface sources (rivers and lakes). Point-of-use water quality improvement interventions include boiling, chlorination, flocculation, filtration, or solar disinfection, mainly conducted at home.

Objectives

To assess the effectiveness of interventions to improve water quality for preventing diarrhoea.

Search methods

We searched the Cochrane Infectious Diseases Group Specialized Register (11 November 2014), CENTRAL (the Cochrane Library, 7 November 2014), MEDLINE (1966 to 10 November 2014), EMBASE (1974 to 10 November 2014), and LILACS (1982 to 7 November 2014). We also handsearched relevant conference proceedings, contacted researchers and organizations working in the field, and checked references from identified studies through 11 November 2014.

Selection criteria

Randomized controlled trials (RCTs), quasi-RCTs, and controlled before-and-after studies (CBA) comparing interventions aimed at improving the microbiological quality of drinking water with no intervention in children and adults.

Data collection and analysis

Two review authors independently assessed trial quality and extracted data. We used meta-analyses to estimate pooled measures of effect, where appropriate, and investigated potential sources of heterogeneity using subgroup analyses. We assessed the quality of evidence using the GRADE approach.

Main results

Forty-five cluster-RCTs, two quasi-RCTs, and eight CBA studies, including over 84,000 participants, met the inclusion criteria. Most included studies were conducted in low- or middle-income countries (LMICs) (50 studies) with unimproved water sources (30 studies) and unimproved or unclear sanitation (34 studies). The primary outcome in most studies was self-reported diarrhoea, which is at high risk of bias due to the lack of blinding in over 80% of the included studies.

Source-based water quality improvements

There is currently insufficient evidence to know if source-based improvements such as protected wells, communal tap stands, or chlorination/filtration of community sources consistently reduce diarrhoea (one cluster-RCT, five CBA studies, *very low quality evidence*). We found no studies evaluating reliable piped-in water supplies delivered to households.

Point-of-use water quality interventions

On average, distributing water disinfection products for use at the household level may reduce diarrhoea by around one quarter (Home chlorination products: RR 0.77, 95% CI 0.65 to 0.91; 14 trials, 30,746 participants, *low quality evidence*; flocculation and disinfection sachets: RR 0.69, 95% CI 0.58 to 0.82, four trials, 11,788 participants, *moderate quality evidence*). However, there was substantial heterogeneity in the size of the effect estimates between individual studies.

Point-of-use filtration systems probably reduce diarrhoea by around a half (RR 0.48, 95% CI 0.38 to 0.59, 18 trials, 15,582 participants, *moderate quality evidence*). Important reductions in diarrhoea episodes were shown with ceramic filters, biosand systems and LifeStraw® filters; (Ceramic: RR 0.39, 95% CI 0.28 to 0.53; eight trials, 5763 participants, *moderate quality evidence*; Biosand: RR 0.47, 95% CI 0.39 to 0.57; four trials, 5504 participants, *moderate quality evidence*; LifeStraw®: RR 0.69, 95% CI 0.51 to 0.93; three trials, 3259 participants, *low quality evidence*). Plumbed in filters have only been evaluated in high-income settings (RR 0.81, 95% CI 0.71 to 0.94, three trials, 1056 participants, fixed effects model).

In low-income settings, solar water disinfection (SODIS) by distribution of plastic bottles with instructions to leave filled bottles in direct sunlight for at least six hours before drinking probably reduces diarrhoea by around a third (RR 0.62, 95% CI 0.42 to 0.94; four trials, 3460 participants, *moderate quality evidence*).

In subgroup analyses, larger effects were seen in trials with higher adherence, and trials that provided a safe storage container. In most cases, the reduction in diarrhoea shown in the studies was evident in settings with improved and unimproved water sources and sanitation.

Authors' conclusions

Interventions that address the microbial contamination of water at the point-of-use may be important interim measures to improve drinking water quality until homes can be reached with safe, reliable, piped-in water connections. The average estimates of effect for each individual point-of-use intervention generally show important effects. Comparisons between these estimates do not provide evidence of superiority of one intervention over another, as such comparisons are confounded by the study setting, design, and population.

Further studies assessing the effects of household connections and chlorination at the point of delivery will help improve our knowledge base. As evidence suggests effectiveness improves with adherence, studies assessing programmatic approaches to optimising coverage and long-term utilization of these interventions among vulnerable populations could also help strategies to improve health outcomes.

PLAIN LANGUAGE SUMMARY

Interventions to improve water quality and prevent diarrhoea

This Cochrane Review summarizes trials evaluating different interventions to improve water quality and prevent diarrhoea. After searching for relevant trials up to 11 November 2014, we included 55 studies enrolling over 84,000 participants. Most included

studies were conducted in low- or middle-income countries (LMICs) (50 studies), with unimproved water sources (30 studies), and unimproved or unclear sanitation (34 studies).

What causes diarrhoea and what water quality interventions might prevent diarrhoea?

Diarrhoea is a major cause of death and disease, especially among young children in low-income countries where the most common causes are faecally contaminated water and food, or poor hygiene practices.

In remote and low-income settings, source-based water quality improvement may include providing protected groundwater (springs, wells, and bore holes) or harvested rainwater as an alternative to surface sources (rivers and lakes). Alternatively water may be treated at the point-of-use in people's homes by boiling, chlorination, flocculation, filtration, or solar disinfection. These point-of-use interventions have the potential to overcome both contaminated sources and recontamination of safe water in the home.

What the research says

There is currently insufficient evidence to know if source-based improvements in water supplies, such as protected wells and communal tap stands or treatment of communal supplies, consistently reduce diarrhoea in low-income settings (*very low quality evidence*). We found no trials evaluating reliable piped-in water supplies to people's homes.

On average, distributing disinfection products for use in the home may reduce diarrhoea by around one quarter in the case of chlorine products (*low quality evidence*), and around a third in the case of flocculation and disinfection sachets (*moderate quality evidence*).

Water filtration at home probably reduces diarrhoea by around a half (*moderate quality evidence*), and effects were consistently seen with ceramic filters (*moderate quality evidence*), biosand systems (*moderate quality evidence*) and LifeStraw® filters (*low quality evidence*). Plumbed-in filtration has only been evaluated in high-income settings (*low quality evidence*).

In low-income settings, distributing plastic bottles with instructions to leave filled bottles in direct sunlight for at least six hours before drinking probably reduces diarrhoea by around a third (*moderate quality evidence*).

Research assessing the effects of household connections and chlorination at the point of delivery will help improve our knowledge base. Evidence indicates the more people use the various interventions for improving water quality, the larger the effects, so research into practical approaches to increase coverage and help assure long term use of them in poor groups will help improve impact.

SUMMARY OF FINDINGS FOR THE MAIN COMPARISON [Explanation]

Point-of-use water quality interventions for preventing diarrhoea in rural settings in low- and middle-income countries

Patient or population: adults and children

Settings: low- and middle-income countries in rural areas **Intervention:** point of use water quality interventions

Comparison: no intervention

| Outcomes | | | Relative effect (95% CI) | Number of participants (trials) | Quality of the evidence (GRADE) | |
|--------------------|--------------------------------|----------------------------------|-------------------------------|---------------------------------|-------------------------------------|-------------------------------|
| | Assumed risk | Corresponding risk | | | | |
| Diarrhoea episodes | No intervention | Chlorination | RR 0.77 | 30,746 | 0 000 | |
| | 3 episodes per person per year | 2.3 episodes (2.0 to 2.7) | (0.65 to 0.91) | (14 trials) | IOW ¹ 1-2-3-4 | low ^{1,2,3,4} |
| | No intervention | Flocculation/disinfection | RR 0.69 | 11,788 | 000 | |
| | 3 episodes per person per year | 2.1 episodes (1.7 to 2.5) | (0.58 to 0.82) | (4 trials) | (4 trials) mode | moderate ^{1,3,4,5,6} |
| | No intervention | Filtration | RR 0.48 | 15,582 | ⊕⊕⊕⊝ | |
| | 3 episodes per person per year | 1.4 episodes (1.1 to 1.8) | (0.38 to 0.59) | (18 trials) | moderate ^{1,3,4,5} | |
| | No intervention | Solar disinfection (SODIS) | RR 0.62 (0.42 to 0.94) | 3460 (4 trials) | ⊕⊕⊕⊖ moderate ^{1,3,4,5} | |
| | 3 episodes per person per year | 1.9 episodes (1.3 to 2.8) | | | | |

The **assumed risk** is taken from Fischer Walker 2012 and represents an estimated average for the incidence of diarrhoea in low- and middle-income countries. 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).

CI: confidence interval; RR: risk ratio.

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.

¹Downgraded by 1 for serious risk of bias: the outcome was measured as self-reported episodes of diarrhoea, and is susceptible to bias as most studies were unblinded.

²Downgraded by 1 for serious inconsistency: statistical heterogeneity was very high with six out of fourteen trials having point estimates close to no effect. A subgroup analysis by adherence with the intervention (assessed by measurements of residual chlorine in drinking water) found larger effects in the studies with better adherence but the results remained inconsistent.

³No serious indirectness: these studies are mainly from low- and middle-income countries, in settings with both improved and unimproved water sources and sanitation.

⁴No serious imprecision: The analysis is adequately powered to detect this effect.

⁵No serious inconsistency: The evidence of benefit is consistent across trials, but there is substantial statistical heterogeneity in the size of the effect.

⁶ This analysis excludes one additional study which found a much larger effect than seen in the other four trials and was considered an outlier (Doocy 2006 LBR).

BACKGROUND

Description of the condition

Diarrhoeal disease is the third leading cause of mortality in low-income countries, causing an estimated 1.4 million deaths in 2012 (WHO 2014;GBD 2015). Young children are especially vulnerable, with diarrhoea accounting for more than a quarter of all deaths in children aged under five years in Africa and Southeast Asia (Murray 2012; Lanata 2013; Walker 2013).

The bacterial, viral, and protozoan pathogens causing diarrhoeal disease are primarily transmitted via the faecal-oral route, through the consumption of faecally contaminated food and water (Byers 2001). Among the most important of these are rotavirus, Cryptosporidium sp., Escherichia coli, Salmonella sp., Shigella sp., Campylobacter jejuni, Vibrio cholerae, norovirus, Giardia lamblia, and Entamoeba histolytica (Leclerc 2002; Kotloff 2013), though the relative importance of these varies among settings, seasons, and population groups.

An estimated 1.1 billion people worldwide rely on water supplies that are at high risk of faecal contamination (Bain 2014). Moreover, nearly half the world's population lack household water connections (WHO/UNICEF 2015), and are at increased risk of unsafe water due to contamination during collection, storage, and use in the home (Wright 2004).

Description of the intervention

Interventions to improve the microbiological quality of water can be grouped into four main categories:

- Physical removal of pathogens (for example, filtration, adsorption, or sedimentation).
- Chemical treatment to kill or deactivate pathogens (most commonly with chlorine).
- Disinfection by heat (for example, boiling or pasturization) or ultraviolet (UV) radiation (for example, solar disinfection, or artificial UV lamps).
- Combination of these approaches (for example, filtration or flocculation combined with disinfection).

In higher-income countries, and in many urban settings world-wide, drinking water is treated centrally at the source of supply and distributed to consumers through a network of pipes and house-hold taps. Alternatively, water may be treated at any point in the distribution network, or at the 'point-of-use' (POU) in people's homes, schools, or workplaces.

In remote and low-income settings, source-based water quality improvement may include providing protected groundwater (springs, wells, and bore holes) or harvested rainwater as an alternative to surface sources (rivers and lakes). These improvements frequently also improve both the quantity and access to water by increasing the volume or frequency of water delivery or reducing

the time spent in collecting water. This may result in significant benefits not only in health but also in economic and social welfare (Hutton 2013; Stelmach 2015).

Potential and widely used POU interventions for remote or low-income settings include boiling, filtration, chlorination, flocculation, and solar disinfection. These interventions have the potential to overcome both contaminated sources and recontamination of safe water in the home (Wright 2004). A review commissioned by the World Health Organization (WHO) identified a wide variety of options for household-based water treatment and assessed the available evidence on their microbiological effectiveness, health impact, acceptability, affordability, sustainability, and scalability (Sobsey 2002).

How the intervention might work

Health authorities generally accept that microbiologically safe water plays an important role in preventing outbreaks of waterborne diseases (Reynolds 2008). Moreover, there is evidence that chlorination and filtration of municipal water supplies contributed to substantial health gains in the late 19th and early 20th century (Cutler 2005).

However, much of the epidemiological evidence for increased health benefits following improvements in the quality of drinking water has been equivocal, particularly in low-income settings (Clasen 2006; Waddington 2009; Cairncross 2010).

This may be due to the variety of alternative transmission pathways, such as ingestion of contaminated food, person-to-person contact, or direct contact with infected faeces. In addition, interventions which only target the home may fail if unsafe water is consumed at work or school. Consequently, effective programmes may require combined interventions to address not only water quality, but also water quantity and access, the proper disposal of human faeces (sanitation), and the promotion of hand washing and hygiene practices within communities.

The effectiveness of individual water quality interventions may also vary between settings due to the varied prevalence of the organisms causing diarrhoea. For instance, ceramic filters are only marginally protective against viral illness, while chlorination may provide little protection against *Cryptosporidium*.

Why it is important to do this review

This is an update of a Cochrane Review that was first completed in 2006 (Clasen 2006). The review concluded that, in general, interventions to improve microbiological quality of drinking water are effective in preventing diarrhoea, and that interventions at the household level were more effective than those at the source.

New studies have been recently published, and other unpublished studies have been made available to us. In this Cochrane Review update, we have reapplied the inclusion criteria, repeated data extraction, added new studies, and used the Grading of Recommendations Assessment, Development and Evaluation (GRADE) approach to assess the quality of the evidence. We were also able to apply statistical methods to unify the measures of effect and to apply additional criteria for subgrouping based on study design, setting, and length of follow-up.

OBJECTIVES

To assess the effectiveness of interventions to improve water quality for preventing diarrhoea.

METHODS

Criteria for considering studies for this review

Types of studies

Cluster-randomized controlled trials (cluster-RCTs), quasi-randomized controlled trials (quasi-RCTs) and controlled before-and-after studies (CBAs).

Types of participants

Children and adults.

Types of interventions

Intervention

Any intervention aimed at improving the microbiological quality of drinking water.

We included interventions that combined improvements in water quality with hygiene or health promotion, but excluded studies that combined water quality interventions with other water, sanitation, and hygiene (WASH) interventions, such as improvements in water quantity or sanitation. We also excluded studies where the water quality intervention was implemented away from the home, such as at schools, clinics, markets, or workplaces.

Control

No intervention, or a dummy intervention.

Types of outcome measures

Primary

• Diarrhoea episodes among individuals, whether or not confirmed by microbiological examination.

The WHO's definition of diarrhoea is three or more loose or fluid stools (that take the shape of the container) in a 24-hour period (WHO 1993). We defined diarrhoea and an episode in accordance with the case definitions used in each trial. In the 'Summary of findings' tables, we have converted the results to episodes per year from a baseline of three episodes/child year in 2010 (Fischer Walker 2012).

Secondary

- Death.
- Adverse events.

We excluded studies that had no clinical outcomes; for example, studies that only report on microbiological pathogens in the stool.

Search methods for identification of studies

We attempted to identify all relevant studies regardless of language or publication status (published, unpublished, in press, and in progress).

Electronic searches

We searched the following databases using the search terms and strategy described in Appendix 1: Cochrane Infectious Diseases Group Specialized Register (11 November 2014); Cochrane Central Register of Controlled Trials (CENTRAL), published in the Cochrane Library (7 November, 2014); MEDLINE (1966 to 10 November 2014); EMBASE (1974 to 10 November 2014); and LILACS (1982 to 7 November 2014).

Searching other resources

Conference proceedings

We searched the conference proceedings of the following organizations for relevant abstracts: International Water Association (IWA) (1990 to 11 November 2014); and Water, Engineering and Development Centre, Loughborough University, UK (WEDC) (1973 to 11 November 2014).

Researchers and organizations

We contacted individual researchers working in the field and the following organizations for unpublished and ongoing studies: Water, Sanitation and Health Programme of the WHO; World Bank Water and Sanitation Program; UNICEF Water, Sanitation and Hygiene; and IRC International Water and Sanitation Centre; Foodborne and Diarrhoeal Diseases Branch, Division of Bacterial and Mycotic Diseases, Centers for Disease Control and Prevention (CDC); US Agency for International Development (USAID), including its Environmental Health Project (EHP); and the UK Department for International Development (DFID).

Reference lists

We checked the reference lists of all studies identified by the above methods.

Data collection and analysis

Selection of studies

Two review authors (RP and SB) independently reviewed the titles and abstracts located in the searches and selected all potentially relevant studies. After obtaining the full-text articles, they independently determined whether they met the inclusion criteria. Where they were unable to agree, they consulted a third review author (TFC) and arrived at a consensus. We have listed the potentially relevant studies that were ultimately excluded together with the reasons for exclusion in the 'Characteristics of excluded studies' section.

Data extraction and management

Two review authors (RP and SB) used a pre-piloted form to extract and record the data described in Appendix 2. One review author entered the extracted data into Review Manager (RevMan) (KA).

Assessment of risk of bias in included studies

Two review authors (KA and FM) independently assessed the risk of bias of the included studies and resolved differences of opinion through discussion.

For cluster-RCTs we used the Cochrane 'Risk of bias' assessment tool (Higgins 2011). We followed the guidance to assess whether adequate steps were taken to reduce the risk of bias across five domains: sequence generation; allocation concealment; blinding of participants and personnel; blinding of outcome assessors; and incomplete outcome data.

For sequence generation and allocation concealment, we reported the methods used. For blinding, we described who was blinded and the blinding method. For incomplete outcome data, we reported the percentage and proportion of participants lost to follow-up. For selective outcome reporting, any discrepancies between the methods used and the results were stated in terms of the outcomes measured or the outcomes reported. For other biases, we described any other trial features that could have affected the trial result (for example, if the trial was stopped early).

We categorized our 'Risk of bias' judgements as 'low', 'high', or 'unclear'. Where risk of bias was unclear, we attempted to contact the study authors for clarification and we resolved any differences of opinion through discussion. We classified the inclusion of randomized participants in the analysis as 'low risk' if 90% or more of all participants randomized to the study were included in the analysis.

For quasi-RCTs and CBA studies, we used two additional criteria:

- 1. Comparability of baseline characteristics: we classified studies as 'low risk' if there were no substantial differences between groups with respect to water quality, diarrhoeal morbidity, age, socioeconomic status, access to water, hygiene practices, and sanitation facilities.
- 2. Contemporaneous data collection: we classified studies as 'low risk' if data were collected at similar points in time, 'unclear' if the relative timing was not reported or not clear from trial, or 'high risk' if data were not collected at similar points in time.

Measures of treatment effect

Two review authors independently extracted and, where necessary, calculated the measure of effect of the intervention on diarrhoea. We extracted the measure of effect as reported by the authors of each study, whether it be risk ratios (RRs), rate ratios, odds ratios (ORs), longitudinal prevalence ratios, or means ratios. In using these various measures of effect, we noted the design effect in treating all such measures of effect as equivalent for common outcomes such as diarrhoea and the debate about methodologies for converting such measures of effect into a single measure (Zhang 1998; McNutt 2003).

For purposes of analysis, we transformed ORs into RRs using the assumed control risk and the formula prescribed in Higgins 2011 (Section 12.5.4.4).

Unit of analysis issues

A number of the included studies had multiple intervention arms (for example, treating water with bleach or with a flocculant and disinfectant) and compared two or more intervention groups against a single control group. In some analyses, we included multiple comparisons from the same study, which double counts the control group participants and yields results in the meta-analysis that are artificially precise. Unfortunately, because of the way data was presented in included studies, it was not possible to correct for this error by dividing the control group participants between multiple groups.

Dealing with missing data

When data was missing or incomplete we attempted to contact the study authors.

Assessment of heterogeneity

We assessed the statistical heterogeneity between trials by visually examining the forest plots for overlapping confidence intervals (CIs), applying the Chi² test with a 10% level of statistical significance, and using the I² statistic with a value of 50% to denote moderate levels of heterogeneity.

Assessment of reporting biases

When there were sufficient studies, we assessed the possibility of publication bias by constructing funnel plots and looking for asymmetry.

Data synthesis

We entered the estimates of effect using the generic inverse variance method on the log scale (Higgins 2006), and analysed the data using Review Manager (RevMan).

We stratified our primary analysis by intervention type, and study design (cluster-RCT, quasi-RCT, or CBA). When appropriate we used meta-analyses to derive pooled estimates of effect using a random-effects model because of the substantial heterogeneity in study settings, interventions, and outcome measures.

We summarized the evidence using 'Summary of findings' tables that we created using the GRADE Guideline Development Tool (GRADEpro GDT). The quality of evidence was rated using the GRADE approach, which consists of five factors that are used to assess the quality of the evidence: study limitations (risk of bias), inconsistency, indirectness, imprecision, and publication bias (Guyatt 2008).

Subgroup analysis and investigation of heterogeneity

We investigated the potential causes of heterogeneity by conducting the following subgroup analyses: age (all ages versus children under five years old); adherence with intervention (< 50%, 50% to 85%, > 85%); water source; water access; water quantity; sanitation conditions; country income level; and length of follow-up. In the subgroup analyses based on water source, we followed terminology used by the WHO/UNICEF Joint Monitoring Programme (JMP) on Water and Sanitation (WHO/UNICEF 2015), using 'unimproved' to extend to unprotected wells or springs, vendoror tanker-provided water or bottled water, and 'improved' to extend to household connections, public standpipes, boreholes, protected dug wells or springs, or rainwater collection; we categorized studies as 'unclear' with respect to water supply if they contained insufficient information.

We used the same definitions from the WHO/UNICEF JMP criteria to classify sanitation conditions as 'improved' (connection

to a public sewer or septic system, pour-flush latrine, simple pit latrine, ventilated improved pit latrine) or 'unimproved' (service or bucket latrines, public latrines, open latrines); where the necessary information was unclear or unreported, we categorized the sanitation facilities as 'unclear'.

To subgroup studies based on access to water source, we used the classifications defined by the Sphere Project 2011, classifying access as 'sufficient' if a consistently available source was located within 500 m, with queuing no more than 15 minutes and filling time for a 20 L container no more than three minutes, 'insufficient' if any access failed any such criteria, and 'unclear' if such criteria was unreported or unclear.

The quantity of water available to study participants was considered 'sufficient' if consisting of a minimum of 15 L per person per day. For country income level, we used the World Bank classification of country income levels (high, upper middle, lower middle, low) (World Bank Country and Lending Groups).

Sensitivity analysis

We conducted a sensitivity analysis to investigate the robustness of the results to each of the 'Risk of bias' components by including only studies that were at low risk of bias. We used this information to guide our judgements on the quality of the evidence.

In addition, we explored the impact of non-blinding of POU interventions using a Bayesian meta-analysis with bias correction. For this purpose, we assumed the true log relative risks from non-blinding studies are subject to a multiplicative bias that results in the observed relative risks being inflated in magnitude. We assumed the bias is normally distributed with a mean 1.48 or 1.65 and a corresponding standard deviation (SD) of 0.17 or 0.13. These values were derived from the additive bias correction employed in Wood 2008 and Savovie 2012. While we believe an attempt to adjust for non-blinding is appropriate, we urge caution in relying on these adjusted estimates since the basis for the adjustment is from clinical (mainly drug) studies that may not be transferable to field studies of environmental interventions and because methodology for the adjustment has not been validated.

RESULTS

Description of studies

Results of the search

The search strategy identified 1088 titles and abstracts, 1076 from the databases and 12 from the other sources (Figure 1). We screened these titles and abstracts, and obtained the full-text articles of 161 studies for further assessment.

1076 records 12 additional records identified identified through through other sources database searching 1088 records screened 925 records excluded 163 full-text articles 108 full-text articles assessed for eligibility excluded, with reasons 55 studies included in 3 studies excluded from qualitative synthesis quantiative analysis 52 records, corresponding to 65 comparisons, included in quantitative synthesis (meta-analysis)

Figure I. Study flow diagram.

Included studies

Fifty-five studies, including 84,023 participants, met the inclusion criteria (see Characteristics of included studies). Of these, six studies had two relevant intervention arms (Austin 1993; URL 1995; Luby 2004; Crump 2005; Brown 2008; Lindquist 2014), two had three arms (Luby 2006; Opryszko 2010), and one had four arms (Reller 2003), making a total of 65 discrete comparisons. Three included studies had inadequate information on disease morbidity to include in the quantitative analysis (Torun 1982 GTM; Kremer 2011 KEN; Patel 2012 KEN). We contacted the study authors for

further information, but no data could be provided. Therefore we have only described these three studies and their results, but have not integrated these studies into the analysis.

Study design and length

Forty-five studies were cluster-RCTs, two were quasi-RCTs, and eight were CBA studies. Most included cluster-RCTs used house-holds as the unit of randomization, though some used neighbour-hoods, villages, or communities. Most CBA studies used villages or communities as the unit of allocation. The intervention period

ranged from eight weeks to four years. The duration of the cluster-RCTs (median seven months, range 9.5 weeks to 18 months) tended to be shorter than in the CBA studies (median 12 months, range two to 60 months). Studies of source-based interventions were also longer (median 24 months, range eight weeks to two years) than those of POU interventions (median six months, range 9.5 weeks to 17 months).

Participants and settings

Nine studies included data only for children under five years of age, and three studies included data only on adults. The other studies enrolled and presented results for all ages of participants. Most studies were undertaken in lower middle or low-income countries based on World Bank criteria, but three studies were conducted in the USA (Colford 2002 USA; Colford 2005 USA; Colford 2009 USA), one in Australia (Rodrigo 2011 AUS), and one in Saudi Arabia (Mahfouz 1995 KSA). Five studies were conducted in urban settings (Semenza 1998 UZB; Colford 2002 USA; Colford 2005 USA; Colford 2009 USA; Rodrigo 2011 AUS), five in peri-urban settings (Quick 1999 BOL; Quick 2002 ZMB; du Preez 2010 ZAF; Jain 2010 GHA; Peletz 2012 ZMB), two in urban informal or squatter settlements (Handzel 1998 BGD; Luby 2004), two in camps for refugees or displaced persons (Roberts 2001 MWI; Doocy 2006 LBR), five in multiple settings (URL 1995; Clasen 2005 COL; Stauber 2009 DOM; du Preez 2011 KEN; Boisson 2013 IND), and the others in villages or other rural settings.

Primary drinking water supply and sanitation facilities

The primary drinking water supply before the intervention was 'unimproved' in 30 studies, 'improved' in 15 studies, and 'unclear' or unreported in five studies. Sanitation facilities in trial settings were 'improved' in 12 studies, 'unimproved' in 15 studies, and 'unclear' or unreported in 19 studies. Access to a water source was deemed 'sufficient' in 14 studies, 'insufficient' in four studies, and 'unclear' or unreported in the remaining studies. The quantity of water available to study participants was considered 'sufficient' in eight studies, 'insufficient' in four studies, and 'unclear' in 43 studies.

Seventeen studies measured water quality before the introduction of the intervention as an indication of the ambient risk and the microbiological quality of the water consumed by the control group. Details on the indicators used varied among the studies (see Table 1). Thirty-five studies measured colony-forming units (CFUs) of thermotolerant coliforms, faecal coliforms, or *E. coli*, reporting geometric means, arithmetic means, number of CFUs/100 mL, mean faecal coliforms/100 mL, *E. coli* most probable number, median, or log₁₀CFUs/100 mL. Other studies measured the frequency of samples containing such bacteria, or the CFU of total coliforms or other indicators of microbial contamination. None

continually measured the microbiological performance of their interventions against the full range of bacterial, viral, and protozoan pathogens known to cause diarrhoea.

Eight studies did not report actually having measured microbiological water quality at all (Alam 1989 BGD; Xiao 1997 CHN; Luby 2006; Mäusezhal 2009 BOL; Opryszko 2010; Majuru 2011 ZAF; Rodrigo 2011 AUS; Lindquist 2014). Thus, it cannot be concluded definitively that the interventions investigated in these studies actually resulted in an improvement in drinking water quality.

Among the eight studies investigating interventions to improve water quality at the point of distribution, only four tested microbiological water quality (Torun 1982 GTM; Gasana 2002 RWA; Jensen 2003 PAK; Kremer 2011 KEN). As these tests were at the source or point of distribution and not the POU, their results do not reflect possible post-collection contamination.

Interventions

Eight studies evaluated source-based interventions: improved wells or boreholes (Alam 1989 BGD; Xiao 1997 CHN; Opryszko 2010b AFG; Opryszko 2010c AFG) or improved community sources and distribution to public tap stands (Torun 1982 GTM; Gasana 2002 RWA; Jensen 2003 PAK; Kremer 2011 KEN; Majuru 2011 ZAF); none evaluated reliable piped-in household connections.

Fourty-seven studies evaluated POU interventions: chlorination (17 studies), filtration (20 studies), combined flocculation and disinfection (five studies), SODIS solar disinfection (six studies), combination UV disinfection and filtration (one study), and improved storage (two studies). Significantly, there were no eligible studies that investigated the impact of boiling, even though that is by far the most common type of POU water treatment (Rosa 2010).

Many studies provided a supplementary hygiene education or instruction beyond the use of the intervention itself, and among POU interventions the primary intervention was often combined with some form of improved storage. In only three multiple-intervention arm studies did study authors establish different intervention groups with and without hygiene or other non-water improvement steps in order to isolate the impact of water quality (URL 1995; Opryszko 2010; Lindquist 2014).

Except in blinded trials involving placebos, control arms generally continued to use their pre-trial water supply and treatment practices. In one trial of POU chlorination plus a safe storage container, however, control households also received the container (Jain 2010 GHA). In two of the solar disinfection studies (Conroy 1996 KEN; Conroy 1999 KEN) both intervention and control households received plastic bottles for storing their drinking water. The intervention group was instructed to place the bottles on roofs to expose them to the sun, while the control group was told to keep the filled bottles indoors. It is important to note that since

improved storage even in the absence of treatment has been shown to improve microbial water quality (Wright 2004), the comparison between the intervention and control in these studies may understate the effectiveness of the intervention when compared to the controls following customary water handling practices.

Adherence with the intervention

Studies of source water interventions tended to assume adherence based on the fact that the primary water supply had been improved. Some studies of POU water treatment undertook indirect assessments of adherence by measuring residual chlorine levels in stored water, comparing microbiological water quality of intervention and control groups, conducting periodic or post-study surveys, or counting the amount of intervention product used. Most other studies measured adherence only by occasional observation, while eight cluster-RCTs did not report on adherence.

The studies of chlorine residuals reported adherence ranging from a high of 95% (Doocy 2006 LBR) to a low of 11% (Opryszko 2010a AFG). Even among these studies, however, investigators acknowledged that it was not possible to know to what extent intervention group participants actually consumed treated water or avoided consuming untreated water. For those studies that reported on adherence, three took the additional step of investigating and reporting on continued consumption of untreated water (Boisson 2010 DRC; Peletz 2012 ZMB; Boisson 2013 IND) a source of exposure that could be masked by less direct metrics of adherence.

Outcome measures

The studies' main outcome measure was diarrhoeal disease, but different methods were used to define, assess, and report this. Thirty-six studies used the WHO's definition of diarrhoea, while other studies used the following definitions: the mother's or respondent's definition (Austin 1993; Gasana 2002 RWA; Reller 2003; Crump 2005; Chiller 2006 GTM); 'watery diarrhoea as a component of gastroenteritis' (Colford 2002 USA; Colford 2005 USA; Colford 2009 USA; Rodrigo 2011 AUS); the local term (Conroy 1996 KEN; Conroy 1999 KEN; Boisson 2009 ETH); "significant change in bowel habits towards decreased consistency or increased frequency" (Kirchhoff 1985 BRA); or dysentery (du Preez 2010 ZAF; du Preez 2011 KEN). Four studies did not report the case definition used for diarrhoea (Torun 1982 GTM; Xiao 1997 CHN; Günther 2013 BEN; Lindquist 2014).

The method of diarrhoea surveillance and assessment also varied. In most cases, participants were visited on a periodic basis, either weekly (19 studies), fortnightly (16 studies), or more infrequently (14 studies). Participants were asked to recall and report on cases of diarrhoea during a previous period, usually seven days (30 studies) or 14 days (six studies), with four studies having recall periods of one to four days and one trial having a recall period of four weeks

(Günther 2013 BEN). Twelve studies asked each participant or a designated householder to keep a log or record to indicate days with or without diarrhoea, one procured data on diarrhoea from family records and disease registries (Mahfouz 1995 KSA), or used paediatricians to assess the participants during regular medical checkups (Gasana 2002 RWA). Only one trial did not report the method (Xiao 1997 CHN).

Using these data, study authors reported diarrhoeal disease using one or more of the following epidemiological measures of disease frequency: incidence (34 studies); period prevalence (12 studies); and longitudinal prevalence (nine studies). The studies also reported other measures of disease, including incidence of persistent diarrhoea, gastrointestinal illness, including specific symptoms thereof, incidence or prevalence of bloody diarrhoea, and days of work or school lost due to diarrhoea (Lule 2005 UGA). Seven studies also reported on mortality (Crump 2005; Colford 2009 USA; Boisson 2010 DRC; du Preez 2011 KEN; Kremer 2011 KEN; Peletz 2012 ZMB; Boisson 2013 IND). None reported adverse events.

None of these studies were primarily designed to investigate the impact of the intervention on death, and as such most were underpowered to evaluate this outcome.

Data presentation

Forty-three studies presented results both for children aged under five years (or a subgroup thereof) and for all ages or older age groups, three presented results only for adults, and nine presented results only for children under five years (or a subgroup thereof). Most of the studies adjusted raw data to account for possible covariates, including age, sex, sanitation or hygiene practices, area of residence, household income or proxies thereof, education or maternal literacy, age and occupation of the head of household, number of participants in the household or absent there from, baseline diarrhoea or conditions at baseline, or other variables associated with the household environment and participant behaviour.

Most studies of interventions at the POU also used statistical methods to adjust their results, either for repeated episodes of diarrhoea by the same participant or for clustering within the household, village or both. The studies that did not adjust for clustering may receive excess weight in meta-analysis due to artificial precision (Kirchhoff 1985 BRA; Austin 1993; Mahfouz 1995 KSA; URL 1995).

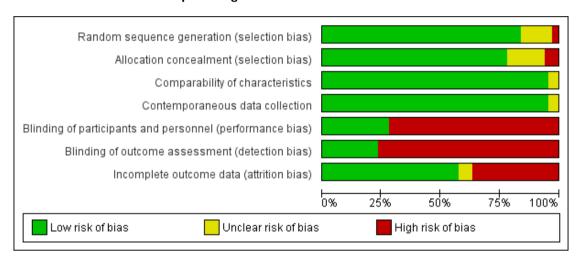
Excluded studies

We excluded 108 studies for the reasons given in the Characteristics of excluded studies table. Two studies that appear to meet this review's inclusion criteria are currently ongoing (see Characteristics of ongoing studies).

Risk of bias in included studies

We have summarized our judgements about the risk of bias of included studies in Figure 2.

Figure 2. Risk of bias graph: summary of authors' judgements about each 'Risk of bias' item presented as percentages across all included studies.



Allocation

The allocation sequence was generated using an adequate method and classified as 'low risk' in 36 of the 45 cluster-RCTs, 'high risk' in two, and 'unclear' in seven Figure 2. The method of allocation concealment was 'low risk' in 34 trials and 'high risk' in two and 'unclear' in nine.

Comparability of baseline characteristics (confounding bias)

All the quasi-RCTs and CBA studies were judged to be at low risk of bias for this criteria except Gasana 2002 RWA, which was at 'unclear' risk.

Contemporaneous data collection

We judged all the quasi-RCTs and CBA studies to be at low risk of bias for this criteria except Gasana 2002 RWA, which was at 'unclear' risk.

Blinding

Nine trials were blinded at the participant level (Kirchhoff 1985 BRA; Austin 1993; Colford 2002 USA; Colford 2005 USA;

Colford 2009 USA; Boisson 2010 DRC; Jain 2010 GHA; Rodrigo 2011 AUS; Boisson 2013 IND); all but two of these were blinded at the assessor level as well (Kirchhoff 1985 BRA; Austin 1993). The others followed an open design, classified as 'high risk' of bias. One of the principal objectives of Colford 2002 USA was to assess the effectiveness of its blinding methodology; it therefore provides the most comprehensive analysis of these issues. Colford 2002 USA, Colford 2005 USA, Boisson 2010 DRC and Rodrigo 2011 AUS all used household sham water filters. Austin 1993, Kirchhoff 1985 BRA, Jain 2010 GHA and Boisson 2013 IND, which were assessing the effectiveness of home-based chlorination, provided placebos to control households.

Incomplete outcome data

Twenty four studies were at 'low risk' of bias, 18 at 'high risk', and three studies were unclear.

Effects of interventions

See: Summary of findings for the main comparison Summary of findings table 1

Analysis I: Any water quality intervention versus no intervention

Diarrhoea episodes

An overall pooled analysis, across different trial designs, interventions and settings, finds the risk of diarrhoea to be lower with any water quality intervention compared to no intervention, both among all ages (RR 0.59, 95% CI 0.51 to 0.69, 81215 participants; 52 studies Analysis 1.1), and under fives (RR 0.61, 95% CI 0.49 to 0.75 Analysis 1.2). However, as would be expected given the diverse nature of the trials, statistical heterogeneity between trials is very high (I² statistic = 98% and 97%, respectively). Our primary analysis is therefore stratified by the specific intervention type (for example, interventions at water source, POU chlorination, POU filtration), and by study design (for example, cluster-RCT, quasi-RCT, CBAs).

Mortality

Only nine studies reported any deaths among study participants. Five reported the number of deaths in each study arm without differences evident (see Table 2). Two studies reported the total number of deaths without stating how many occurred in each group (du Preez 2010 ZAF; Boisson 2013 IND), and two reported recording deaths but the numbers were not presented in the papers (Boisson 2009 ETH; Kremer 2011 KEN).

None of these studies were primarily designed to investigate the impact of the intervention on mortality, and all were underpowered to investigate these effects.

Adverse events

No trial reported adverse events from the interventions.

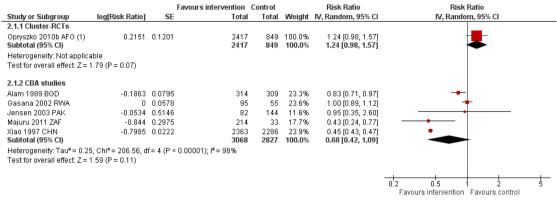
Analysis 2: Interventions at the water source

One cluster-RCT and five CBA studies evaluated interventions at the water source (Table 3). All but one study were from settings with 'unimproved' water sources (unprotected wells or surface water), and all had unclear levels of sanitation. Three studies evaluated improved wells or boreholes, two evaluated chlorination or filtration of community water sources, and one evaluated an improved community piped supply. No studies evaluated reliable household connections to a clean water source (see Table 4 and Table 5 for a description of study settings and interventions). The single cluster-RCT from Afghanistan reported no statistically similar and different at information of the study is a supposed wells connected to the study settings and interventions).

The single cluster-RCT from Afghanistan reported no statistically significant difference in diarrhoea with improved wells compared to no intervention (one trial, 3266 participants; Analysis 2.1; very low quality evidence).

The CBA studies evaluated different interventions, had variable findings, and were all at unclear risk of multiple sources of bias (see Figure 3). Three of the five studies reported statistically significant effects on diarrhoea (Analysis 2.1; Analysis 2.2): in Bangladesh, provision of one hand pump per four to six households (three times as many as control areas) was associated with a small reduction in diarrhoea over three-years follow-up (RR 0.83, 95% CI 0.71 to 0.97); in remote areas in South Africa a new community piped water supply was associated with around a 50% reduction in diarrhoea compared to untreated river water (RR 0.43, 95% CI 0.24 to 0.77); and in China structural well improvements were also associated with around a 50% reduction in diarrhoea (RR 0.45, 95% CI 0.43 to 0.47). In contrast, chlorination and filtration of community water supplies were not associated with positive benefits in Rwanda and Pakistan respectively. Overall, the body of evidence is judged to be of very low quality (Table 3). Given the variability in interventions, further subgroup analyses to try to understand the heterogeneity were not useful.

Figure 3. Forest plot of comparison: 2 Source: water supply improvement versus control, outcome: 2.1 Diarrhoea: CBA studies subgrouped by age.



Footnotes

(1) Opryszko 2010-ii AFG: Provided one well per 25 households providing 25 litres/person/day

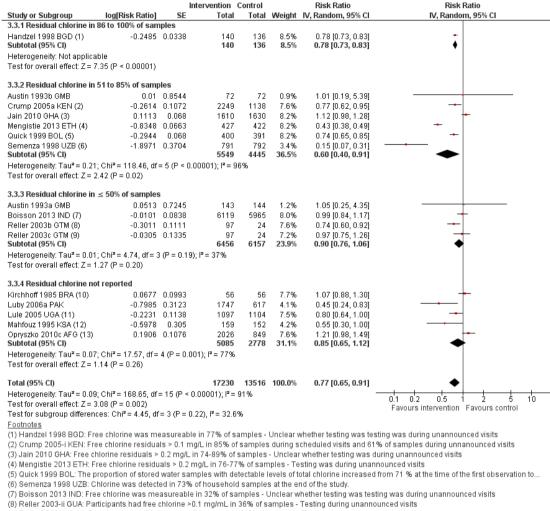
Analysis 3. POU chlorination

Fourteen cluster-RCTs, with 16 comparisons, evaluated POU chlorination versus control. Chlorine was delivered to households free of charge every one to four weeks, with instructions on how to use it, and in eight trials a water storage container was also provided (see Table 6 and Table 7 for a description of study settings and interventions).

On average, POU chlorination in cluster RCTs reduced the risk of diarrhoea episodes by around a quarter, both for all ages (RR 0.77, 95% CI 0.65 to 0.91; 14 trials, 30,746 participants; Analysis 3.2) and for children under five years of age (RR 0.77, 95% CI 0.64 to 0.92; Analysis 3.2). However, there was substantial heterogeneity in the size of the effect which was not well explained by a series of subgroup analyses (Analysis 3.2 to Analysis 3.9).

As might be expected from an effective intervention, the trials finding larger effects from chlorination tended to be those where adherence with the intervention was higher (as measured by residual chlorine) (Analysis 3.3; Figure 4), but in the four trials which had adequate blinding no effects of water chlorination were seen (Analysis 3.4). A subgroup analysis looking at interventions with and without the provision of water storage containers did not find statistical evidence of subgroup differences (Analysis 3.5). Effects were seen in trials with 3, 6, and 12 months of follow-up, but no effect was demonstrated in the two trials with follow-up longer than 12 months (Analysis 3.9). The funnel plot for this comparison has some asymmetry which may be the result of publication bias (see Figure 5). The overall quality of the evidence was therefore judged to be low (Table 8).

Figure 4. Forest plot of comparison: 3 POU: water chlorination versus control, outcome: 3.3 Diarrhoea: cluster-RCTs; subgrouped by adherence.



⁽⁹⁾ Relier 2003-iii GUA: Participants had free chlorine >0.1 mg/mL in 44% of samples - Testing during unannounced visit (10) Kirchhoff 1985 BRA: The chlorination was performed daily by blinded health staff.

⁽¹¹⁾ Lule 2005 UGA: Compliance not reported

⁽¹²⁾ Mahfouz 1995 KSA: The average free residual chlorine is reported as 0.13 ppm

⁽¹³⁾ Opryszko 2010-iii AFG: Self reported use of Chlorine in the previous two weeks was 82%

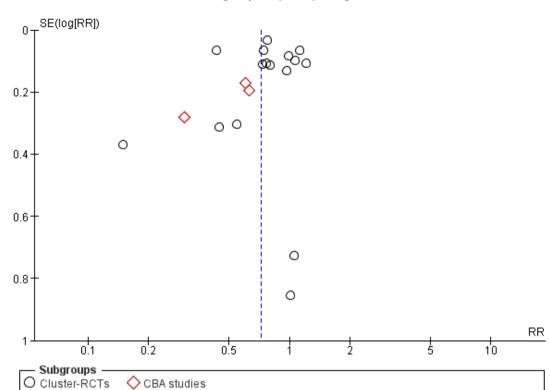


Figure 5. Funnel plot of comparison: 3 POU: water chlorination versus control, outcome: 3.1 Diarrhoea: subgrouped by study design.

An additional two CBA studies evaluated POU chlorination but only provide very low quality evidence of any effect (Analysis 3.1, Table 8).

Analysis 4. POU combined flocculation and disinfection

Five cluster-RCTs from low-income settings evaluated interventions where sachets of flocculant and disinfectant were distributed to households to treat water from unimproved sources (three trials), improved sources (one trial), and unclear sources (one trial). Four trials also provided water containers and mixing equipment (see Table 9 and Table 10 for a description of study settings and interventions). None of the trials blinded the outcome assessment. Four of the five trials found statistically significant reductions in diarrhoea with the intervention (Table 11), but statistical heterogeneity in the size of this effect made pooling the data difficult (I²

statistic = 99%; Analysis 4.1). This heterogeneity relates to one trial from Liberia IDP camps, Doocy 2006 LBR, where the flocculation and disinfection kits reduced diarrhoea by 88% (RR 0.12, 95% CI 0.11 to 0.13; one trial, 2191 participants). Exclusion of this potential outlier finds a more modest effect with the other four trials both for all ages (RR 0.69, 95% CI 0.58 to 0.82; four trials, 11788 participants; Analysis 4.2) and for children under five years of age (RR 0.71, 95% CI 0.61 to 0.84; Analysis 4.2). Adherence with the intervention, as measured by residual chlorine, was generally low (< 50%), but higher in the trial from Liberia showing large effects (Analysis 4.3). Larger effects tended to also be seen in the trials also providing water storage containers (Analysis 4.4). The effects were present in trials with both improved and unimproved water source and sanitation (Analysis 4.5; Analysis 4.6; Analysis 4.7). None of the trials had follow-up longer than

12 months (Analysis 4.8).

Analysis 5. POU filtration

Overall 20 cluster-RCTs evaluated POU filtration: ceramic filtration (nine trials), biosand filtration (five trials), LifeStraw® filters (three trials), and plumbed-in filtration (three trials) (see Table 12 and Table 13 for a description of study settings and interventions). On average, POU filtration technologies reduced diarrhoea by

around a half, both for all ages (RR 0.48, 95% CI 0.38 to 0.59; 18 trials, 15,582 participants; Analysis 5.1) and for children under five years of age (RR 0.49, 95% CI 0.38 to 0.62; Analysis 5.1). However, the number of trials and the quality of evidence was different for each specific intervention (Analysis 5.2; Figure 6). The lack of blinding in these studies is a major concern: of the five trials with adequate blinding only one found a statistically significant effect (Analysis 5.3). The quality of evidence was therefore downgraded for all types of filters due to risk of bias (Table 14).

Figure 6. Forest plot of comparison: 4 POU: filtration versus control, outcome: 4.2 Diarrhoea: cluster-RCTs: subgrouped by type of filtration.

| | | | Intervention | | | Risk Ratio | Risk Ratio |
|---|---------------------|-----------|----------------------------|----------|--------|--------------------|--------------------------------------|
| Study or Subgroup | log[Risk Ratio] | SE | Total | Total | Weight | IV, Random, 95% CI | IV, Random, 95% CI |
| 5.2.1 Ceramic filter | | | | | | | |
| Abebe 2014 ZAF | -1.5418 | | | | 9.9% | | - |
| Brown 2008a KHM | -0.6733 | | | | 9.7% | | |
| Brown 2008b KHM | -0.5447 | | | | 9.8% | | |
| Clasen 2004b BOL | -0.6733 | 0.3023 | 210 | 107 | 7.3% | | |
| Clasen 2004c BOL | -0.5852 | 0.1332 | 140 | 140 | 9.5% | 0.56 [0.43, 0.72] | |
| Clasen 2005 COL | -0.803 | 0.2132 | 415 | 265 | 8.5% | 0.45 [0.29, 0.68] | |
| du Preez 2008 ZAF/ZWE | -1.5606 | 0.2855 | 60 | 55 | 7.5% | 0.21 [0.12, 0.37] | |
| Lindquist 2014a BOL | -1.5606 | 0.1717 | 330 | 140 | 9.1% | 0.21 [0.15, 0.29] | |
| Lindquist 2014b BOL | -1.3093 | 0.1045 | 285 | 139 | 9.8% | 0.27 [0.22, 0.33] | |
| Rodrigo 2011 AUS | -0.1625 | 0.2039 | 698 | 654 | 8.6% | 0.85 [0.57, 1.27] | |
| URL 1995a GTM | | 0.4476 | | 134 | 5.4% | 0.47 [0.20, 1.13] | |
| URL 1995b GTM | -1.0498 | 0.4931 | 297 | 135 | 4.9% | | |
| Subtotal (95% CI) | | | 3556 | 2207 | 100.0% | | • |
| Heterogeneity: Tau ² = 0.2 | 2: Chi² = 116.38, d | f = 11 (P | < 0.00001); I ² | = 91% | | | |
| Test for overall effect: Z= | | | ~ | | | | |
| 5.2.2 Sand filtration | | | | | | | |
| Fabiszewski 2012 HND | -0.4748 | 0.2905 | 532 | 488 | 11.3% | 0.62 [0.35, 1.10] | |
| Stauber 2009 DOM | -0.755 | 0.1221 | 447 | 460 | 63.8% | 0.47 [0.37, 0.60] | - |
| Stauber 2012a KHM | -0.8916 | 0.2732 | 546 | 601 | 12.8% | 0.41 [0.24, 0.70] | |
| Stauber 2012b GHA | -0.8916 | 0.42 | 1012 | 1031 | 5.4% | 0.41 [0.18, 0.93] | |
| Tiwari 2009 KEN | -0.7765 | 0.3763 | 206 | 181 | 6.7% | 0.46 [0.22, 0.96] | |
| Subtotal (95% CI) | | | 2743 | 2761 | 100.0% | 0.47 [0.39, 0.57] | ◆ |
| Heterogeneity: Tau² = 0.0 | | | 86); I² = 0% | | | | |
| Test for overall effect: Z= | 7.68 (P < 0.00001) | | | | | | |
| 5.2.3 LifeStraw® | | | | | | | |
| Boisson 2009 ETH | -0.2877 | 0.1139 | 731 | | | | |
| Boisson 2010 DRC | -0.1625 | 0.1777 | 546 | 598 | 31.7% | 0.85 [0.60, 1.20] | |
| Peletz 2012 ZMB | -0.7765 | 0.2181 | 300 | | 26.2% | | |
| Subtotal (95% CI) | | | 1577 | 1682 | 100.0% | 0.69 [0.51, 0.93] | • |
| Heterogeneity: Tau ² = 0.0 | | 2 (P = 0 | 07); I²= 62% | | | | |
| Test for overall effect: Z = | 2.43 (P = 0.02) | | | | | | |
| 5.2.4 Plumbed | | | | | | | |
| Colford 2002 USA | -0.6061 | 0.1939 | 118 | 118 | 33.6% | 0.55 [0.37, 0.80] | |
| Colford 2005 USA | -0.2399 | | | | 15.0% | | |
| Colford 2009 USA | -0.1393 | | | | 51.3% | | - |
| Subtotal (95% CI) | | | 527 | | 100.0% | | • |
| Heterogeneity: Tau² = 0.0 Test for overall effect: Z = | | 2 (P = 0 | 09); I² = 59% | | | _ | |
| | (/ | | | | | | |
| | | | | | | - | 0.1 0.2 0.5 1 2 5 10 |
| Test for subgroup differer | nces: Chi² = 11 82 | df= 3 /5 | = 0.000\ 12= | 74.2% | | | Favours intervention Favours control |
| reation adaptions differen | 1003. OIII - 11.02, | ui – 5 (F | - 0.000), 1 - | 1 7.2 /0 | | | |

POU ceramic filters reduced diarrhoea by around 60% in nine trials mainly from low- or middle-income countries, regardless of whether the water source or sanitation was classified as improved or unimproved (RR 0.39, 95% CI 0.29 to 0.53, eight trials, 5763

participants; Analysis 5.3; Analysis 5.4; *moderate quality evidence*). Similarly, biosand filtration reduced diarrhoea by around a half consistently across five trials from low- or middle-income settings,

again regardless of whether the water source or sanitation was improved or unimproved (RR 0.47, 95% CI 0.39 to 0.57, four trials, 5504 participants; Analysis 5.6; Analysis 5.7; moderate quality evidence).

On average, the use of LifeStraw® filters reduced diarrhoea by around a third in three trials from low-income settings with unimproved water sources (RR 0.69, 95% CI 0.51 to 0.93; three trials, 3259 participants; Analysis 5.2; low quality evidence).

Plumbed-in filtration has only been evaluated in high-income settings (USA). There is a modest effect in all three trials, although only one reaches standard levels of statistical significance. The overall meta-analysis has similar effect sizes with both fixed effects and random effects models, but wider confidence intervals with random effects (Fixed-effects: RR 0.81, 95% CI 0.70 to 0.94; Random-effects: RR 0.73, 95% CI 0.52 to 1.03; three trials, 1056 participants; Analysis 5.2; moderate quality evidence).

Adherence with the filtration systems was reported by 14 trials, of which eight assessed this by self-reported use which is at high risk of bias due to the lack of blinding. Adherence was generally reported as high, and larger effects were apparent in trials with higher adherence (Analysis 5.8). A subgroup analysis looking at filtration interventions with and without the provision of water storage containers (excluding the trials evaluating plumbed in filtration), found larger effects in the nine trials providing containers (Analysis 5.9). Effects were seen in trials with 3, 6, and 12 months of follow-up, but no effect was demonstrated in the one trial with follow-up longer than 12 months (Analysis 5.10).

Analysis 6. POU solar disinfection (SODIS)

Four cluster-RCTs and two quasi-RCTs evaluated solar disinfection of water from improved sources (one study) and unimproved sources (five studies) in low-income settings. Plastic bottles were distributed to households with instructions to leave filled bottles in direct sunlight for at least six hours before drinking (see Table 15 and Table 16 for a description of study settings and interventions).

Overall in the cluster-RCTs, solar disinfection reduced diarrhoea by around a third for all ages (RR 0.62, 95% CI 0.42 to 0.94; four trials, 3460 participants; Analysis 6.1), and almost a half in children under five years of age (RR 0.55, 95% CI 0.34 to 0.91; Analysis 6.2). The largest effect was seen in the trial with the highest adherence (Analysis 6.3). The quality of evidence was downgraded to moderate due to the lack of blinding and the inherent risk of bias (Table 17).

In the quasi-RCTs the observed effect was lower (RR 0.82, 95% CI 0.69 to 0.97; two trials, 555 participants; Analysis 6.1).

Analysis 7. POU UV disinfection

One cluster-RCT from Mexico evaluated an UV tube disinfection technology (Gruber 2013 MEX; see Table 18 and Table 19 for a description of the study setting and intervention).

The effect on diarrhoea among all age populations did not reach standard levels of statistical significance (RR 0.79, 95% CI 0.49 to 1.27; one trial, 1913 participants; Analysis 7.1), and did not report separately for children under five years of age.

Analysis 8. POU improved storage

Two trials from Malawi and Benin evaluated the distribution of improved water storage containers in settings with improved water sources (see Table 20 and Table 21 for a description of the study setting and intervention).

Overall, there was no statistically significant effect on diarrhoea for all ages (RR 0.91, 95% CI 0.74 to 1.11; two trials, 1871 participants; Analysis 8.1), or children under five years of age (RR 0.69, 95% CI 0.47 to 1.01; Analysis 8.1). Both studies were at high risk of bias due to being non-blinded, and the overall quality of the evidence was judged to be low (Table 22).

Analyses adjusted for non-blinding

In Table 23 we have presented meta-analysis results adjusted for non-blinding using an approach described in the Methods section and based in part on those employed by other researchers (Hunter 2009; Wolf 2014). In these analyses, the effects of POU chlorination and filtration are smaller but remain statistically significant; the effect of POU solar disinfection becomes borderline non-significant.

DISCUSSION

Summary of main results

There is currently insufficient evidence to know if source-based improvements such as protected wells, communal tap stands, or chlorination/filtration of community sources consistently reduce diarrhoea (*very low quality evidence*).

The distribution and promotion of point-of-use water chlorination products may reduce diarrhoea by around one quarter (*low quality evidence*). Similarly, distribution and promotion of flocculation and disinfection sachets probably reduces diarrhoea but had highly variable effects (*moderate quality evidence*).

Point-of-use filtration systems probably reduce diarrhoea by around a half (*moderate quality evidence*). This reduction was apparent for ceramic filters, biosand systems and LifeStraw® filters, but plumbed in filtration has only been evaluated in high-income settings and a statistically significant effect has not been demonstrated.

In low-income settings, distribution of plastic bottles with instructions to leave filled bottles in direct sunlight for at least six hours before drinking (SODIS) probably reduces diarrhoea by around a third (*moderate quality evidence*).

In subgroup analyses, larger effects were seen in trials with higher adherence, and trials that provided a safe storage container.

Overall completeness and applicability of evidence

Fifty-five studies met the inclusion criteria, of which most studies were conducted in low- or middle-income countries (50 studies), with unimproved water sources (30 studies), and unimproved or unclear sanitation (34 studies).

For water source interventions, there are simply too few studies to make conclusions about what may or may not be effective in different settings. While protective effects were seen in some individual trials, it is unclear whether these effects could be expected to be reproducible in other settings, and all of the trials had multiple potential sources of bias. Significantly, we found no studies evaluating reliable, piped-in water supplies.

In contrast, some POU interventions do appear to be broadly protective against diarrhoea across many settings regardless of whether water sources and sanitation are 'improved' or 'unimproved'. This finding affirms the current strategy of the WHO and UNICEF to promote POU water treatment and safe storage, even though this will not increase the number of households with access to improved water supplies and therefore will not contribute towards achieving current international water targets (WHO 2011). The effectiveness of POU interventions in settings without improved sanitation contradicts earlier findings that interventions to improve water quality are effective only where sanitation has already been addressed (Esrey 1986; VanDerslice 1995), or that environmental interventions to prevent diarrhoea are effective only by employing an integrated approach (Eisenberg 2007).

Although we provide average estimates of effect for each individual POU intervention, we recommend caution in using these estimates to conclude the superiority of one intervention over another. Such an observational analysis would be highly susceptible to confounding by study setting and population, and may not represent true differences in the size of the effects. Head-to-head trials would be necessary to reliably conclude superiority and these were not the focus of this review.

As few studies continued follow-up beyond 12 months, we are unable to comment reliably on the long-term sustainability of these effects. While pooled estimates of studies with follow-up periods under 12 months were generally protective, those with follow-up periods in excess of 12 months were not.

Quality of the evidence

The quality of evidence for the effects of the individual interventions on diarrhoea ranged from moderate (for ceramic filters and biosand filtration), to low (for distribution of chlorination kits,

flocculation and disinfection sachets, and LifeStraw® filters), to very low (for water source improvements).

The primary reason for downgrading the quality of evidence was the risk of bias inherent in unblinded studies evaluating the efficacy of an intervention on a self-reported outcome. Notably, only one of the nine blinded trials reported a statistically significant protective effect, but this observation may be explained by other confounding factors present in these nine trials (see Table 24):

- 1. Four studies were conducted in high-income countries where the water was of good microbiological quality even in the control groups (Colford 2002 USA; Colford 2005 USA; Colford 2009 USA; Rodrigo 2011 AUS).
- 2. One further trial from Ghana found very low levels of faecal contamination of water supplies in the control group which were likely to present only minimal risk (Jain 2010 GHA).
- 3. Three studies had either low adherence with the intervention (Austin 1993; Boisson 2013 IND), or very high reported use of drinking untreated water from other sources (Boisson 2010 DRC).
- 4. Two studies employed control interventions which may have improved water quality: Boisson 2010 DRC employed a "placebo" that actually removed one log (90%) of faecal indicator bacteria and Jain 2010 GHA provided control households with safe storage.

The second common reason for downgrading the quality of evidence was unexplained heterogeneity. For some of the POU interventions, the protective effect varied considerably across studies. Some of this variability could be explained by adherence with the intervention, with larger effects in studies with higher adherence, but some variability remained which we were unable to explain despite multiple subgroup analyses. This is likely to reflect important underlying clinical heterogeneity: the aetiology and epidemiology of diarrhoea is complex and variable, transmission pathways are multiple, and even the portion of diarrhoea that is waterborne is not well understood (Eisenberg 2012).

There was also some evidence of possible publication bias in the trials evaluating home chlorination but this was not strong enough to further downgrade the quality of evidence.

Potential biases in the review process

A number of the included studies had multiple intervention arms comparing two or more intervention groups against a single control group. In some analyses, we included multiple comparisons from the same trial which double counts the control group participants and yields results in the meta-analysis that are artificially precise. However, this bias is unlikely to have significantly impacted the overall quality of evidence or conclusions.

Agreements and disagreements with other

studies or reviews

Our results are generally consistent with the prior version of this Cochrane Review (Clasen 2006) and with other reviews of water quality interventions (Fewtrell 2005; Arnold 2007; Waddington 2009; Cairncross 2010; Wolf 2014).

One additional review of water quality interventions reports no effect with POU interventions once blinding is taken into account (Engell 2013). While we share the concerns about the lack of blinding in many of these trials (and have downgraded the quality of evidence accordingly), and also found no effect in any of the trials with adequate blinding, we have identified several possible confounders in this observation (discussed above), and retain low to moderate confidence that these interventions are effective.

Although we found no controlled trials evaluating piped-in water supplies, a recent review that also included some observational studies reported some evidence of a protective effect with this intervention (Wolf 2014).

The finding of larger effects with increased adherence is consistent with modelling data based on quantitative microbial risk assessment which suggest a dose-response association between water quality and diarrhoea (Brown 2012; Enger 2013).

AUTHORS' CONCLUSIONS

Implications for practice

Interventions that address the microbial contamination of water at the POU are important interim measures to improve drinking water quality until homes can be reached with safe, reliable, household piped-water connections.

Implications for research

Rigorously conducted RCTs that compare various approaches to improving drinking water quality will help clarify the potential for water quality interventions to prevent endemic diarrhoea. It is particularly important that such trials be designed to minimize reporting bias, such as through the use of objective outcomes.

Among source-based interventions, there is a need for studies to assess household connections and other approaches (such as chlorination at the point of delivery) that are more likely to ensure safe drinking water from source through to the POU.

There is also a need for longer-term studies in programmatic settings on approaches to optimise the coverage and long-term utilization of these interventions among vulnerable populations.

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CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Abebe 2014 ZAF

| Methods | RCT | |
|---------------|---|--|
| Participants | Number: 74 individuals Inclusion criteria: 18 years or older, receiving anti-retroviral therapy for at least 6 month | |
| Interventions | 1. Ceramic water filter impregnated with silver nanoparticles | |
| Outcomes | Incidence of diarrhoea Water quality Presence of <i>Cryptosporidium</i> in stool | |
| Notes | Location: rural South Africa Length: 12 months Publication status: journal | |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--------------------------------------|
| Random sequence generation (selection bias) | Low risk | Permuted block randomization system. |
| Allocation concealment (selection bias) | Low risk | Permuted block randomization system. |
| Comparability of characteristics | Unclear risk | Not described. |
| Contemporaneous data collection | Unclear risk | Not described. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | > 20% loss to follow-up. |

Alam 1989 BGD

| 1111111 1707 202 | | | |
|---|--|-----------------------------|--|
| Methods | Quasi-RCT | | |
| Participants | Number: 623 children Inclusion criteria: households with children aged 6 to 23 months | | |
| Interventions | Improved water supply and hygiene education (3 subunits) Primary drinking supply (2 subunits) | | |
| Outcomes | 1. Incidence of diarrhoea among children aged 6 to 23 months by water source, hygiene practices, and household socioeconomic characteristics | | |
| Notes | Location: 5 political subunits in a village in rural Bangladesh Length: 3 years Publication status: journal | | |
| Risk of bias | | | |
| Bias | Authors' judgement | Support for judgement | |
| Random sequence generation (selection bias) | Low risk | Irrevelant to study design. | |
| Allocation concealment (selection bias) | Low risk | Irrevelant to study design. | |
| | | | |

Comparability of characteristics Low risk No substantial differences at baseline. Contemporaneous data collection Low risk Data collected at similar points in time. Blinding of participants and personnel Low risk Irrevelant to study design. (performance bias) All outcomes Blinding of outcome assessment (detection Low risk Irrevelant to study design. bias) All outcomes Incomplete outcome data (attrition bias) Low risk Irrevelant to study design. All outcomes

Austin 1993a GMB

| Methods | RCT |
|--------------|---|
| Participants | Number: 287 children Inclusion criteria: households with children aged 25 to 60 months (group B) from villages primarily using open, shallow wells for drinking water |

Austin 1993a GMB (Continued)

| Interventions | Sodium hypochlorite solution used at household level (11 villages) Primary drinking supply (11 villages) | |
|---------------|---|--|
| Outcomes | Longitudinal prevalence of diarrhoea Change in nutritional status using weight-for-height Z-score | |
| Notes | Location: 22 rural villages in The Gambia Length: 20 weeks Publication status: PhD dissertation | |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Random number table. |
| Allocation concealment (selection bias) | Low risk | Numbers assigned to villages. |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Placebo. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | 89.4% of participants included in analysis. |

Austin 1993b GMB

| Methods | See Austin 1993a GMB |
|---------------|--|
| Participants | Number: 144 children between 6 and 24 months Inclusion criteria: as above |
| Interventions | As above |
| Outcomes | As above |
| Notes | As above |

Austin 1993b GMB (Continued)

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Random number table. |
| Allocation concealment (selection bias) | Low risk | Numbers assigned to villages. |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Placebo. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | 89.4% of participants included in analysis. |

Boisson 2009 ETH

| 0005001 2007 1.111 | | | |
|--------------------|---|-----------------------|--|
| Methods | RCT | | |
| Participants | Number: 196 children under 5, 1516 people, 313 households Inclusion criteria: householders were eligible to participate in the study if (i) at least one member of the household worked away from home during the day in a setting without adequate water supply, and (ii) the household was not already practicing an effective POU water treatment method | | |
| Interventions | 1. LifeStraw® personal distributed to each household member over the age of six months. A special attachment was given for children under 3 | | |
| Outcomes | Incidence of diarrhoea among young children in the preceding seven days (recorded fortnightly); other health conditions also recorded Water quality, flow rate and iodine residual Acceptability and use | | |
| Notes | Location: rural Oromia, Ethiopia Length: 5 months Publication status: journal | | |
| Risk of bias | Risk of bias | | |
| Bias | Authors' judgement | Support for judgement | |

Boisson 2009 ETH (Continued)

| Random sequence generation (selection bias) | Low risk | Lottery used to randomly allocate eligible households into intervention and control groups |
|---|-----------|--|
| Allocation concealment (selection bias) | Low risk | Lottery used to randomly allocate eligible households into intervention and control groups |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 4% of person-weeks data lost to follow-up. |

Boisson 2010 DRC

| Methods | RCT |
|---------------|--|
| Participants | Number: 190 children under 5, 1144 people, 240 households Inclusion criteria: unimproved water sources that tested over 1000 thermotolerant coliforms (TTC)/100 ml, reported low use of household water treatment, were easily accessible all year round and were motivated to take part in the project |
| Interventions | 1. LifeStraw® Family filter |
| Outcomes | Incidence of diarrhoea among young children in the preceding seven days (recorded monthly); cough and fever also recorded Filter and water quality monitoring Compliance |
| Notes | Location: rural eastern province of Kasai, Democratic Republic of Congo Length: 12 months Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--------------------------|
| Random sequence generation (selection bias) | Low risk | Random number generator. |

Boisson 2010 DRC (Continued)

| Allocation concealment (selection bias) | Low risk | "Randomisation was stratified by village and was conducted by the trial manager who played no part in the collection of the data" |
|---|-----------|---|
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Double-blinded; however filters removed turbidity, so controls were not always successfully blinded |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Double-blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | 18.2% person-weeks missing due to families moving out of study area, or not being home at time of visit |

Boisson 2013 IND

| Methods | RCT |
|---------------|---|
| Participants | Number: 2986 children under 5, 12,454 people, 2163 households Inclusion criteria: households were eligible if there was at least one child under 5, and they lived permanently in the study area |
| Interventions | 1. Sodium dichloroisocyanurate (NaDCC) disinfection tablets |
| Outcomes | Longitudinal prevalence of diarrhoea among children under 5 Diarrhoea among participants of all ages Weight-for-age z-score, school absenteeism, health care expenditures; adherence; water quality |
| Notes | Location: informal settlements of Orissa, India Length: 12 months Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | "The randomisation list was generated using Stata 10 and was conducted by the trial manager who played no part in the collection of the data" |

Boisson 2013 IND (Continued)

| Allocation concealment (selection bias) | Low risk | "The randomisation list was generated using Stata 10 and was conducted by the trial manager who played no part in the collection of the data" |
|---|-----------|---|
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | "The active and placebo tablets were packaged in identical boxes of three strips containing ten tablets each" |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | "The labeling of the boxes was conducted by members of staff who were neither involved in the implementation nor data col- lection or analysis" |
| Incomplete outcome data (attrition bias) All outcomes | High risk | 12% days of observation lost to follow-up. |

Brown 2008a KHM

| Methods | RCT |
|---------------|---|
| Participants | Number: 239 children under 5, 1196 people, 180 households (across both interventions) Inclusion criteria: households were eligible if they stored drinking water at the household level, if they have at least one child under 5, and if the household was located in the study village |
| Interventions | Iron-rich Cambodian Ceramic Water Purifier Water quality |
| Outcomes | 1. Longitudinal prevalence of diarrhoea for all household members |
| Notes | Location: rural Kandal Province, Cambodia Length: 18 weeks Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Random numbers table. |
| Allocation concealment (selection bias) | Low risk | Households were approached in group-randomized order. |
| Comparability of characteristics | Low risk | Irrelevant to study design. |

Brown 2008a KHM (Continued)

| Contemporaneous data collection | Low risk | Irrelevant to study design. |
|---|-----------|----------------------------------|
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 2% households lost to follow-up. |

Brown 2008b KHM

| Methods | See Brown 2008a KHM |
|---------------|-------------------------------------|
| Participants | As above |
| Interventions | 1. Cambodian Ceramic Water Purifier |
| Outcomes | As above |
| Notes | As above |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Random numbers table. |
| Allocation concealment (selection bias) | Low risk | Households were approached in group-randomized order. |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 2% households lost to follow-up. |

Chiller 2006 GTM

| Methods | RCT |
|---------------|--|
| Participants | Number: 3401 persons from 514 households Inclusion criteria: households with at least one child under 1 year |
| Interventions | Flocculant-disinfectant sachets used at household level Primary drinking supply |
| Outcomes | Longitudinal prevalence of diarrhoea (portion of total days of diarrhoea out of total days of observation) among all ages Incidence of persistent diarrhoea |
| Notes | Location: 42 neighbourhood clusters in 12 rural villages in Guatemala Length: 13 weeks Publication status: journal |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Random number generator used to assigned neighbourhoods to intervention or control group |
| Allocation concealment (selection bias) | Low risk | Random number generator used to assigned neighbourhoods to intervention or control group |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Less than 8% of households lost to follow-up. |

Clasen 2004b BOL

| Methods | RCT |
|--------------|---|
| Participants | Number: 324 persons of all ages from 60 households Inclusion criteria: all households in the community |

Clasen 2004b BOL (Continued)

| Interventions | Household gravity water filter system using imported ceramic filter elements Primary drinking supply |
|---------------|---|
| Outcomes | Period prevalence of diarrhoea (7-day recall) among all ages Microbial water quality |
| Notes | Location: rural Bolivian community Length: 9 months Publication status: unpublished |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Households were randomly allocated by names drawn from a hat in a public assembly |
| Allocation concealment (selection bias) | Low risk | Households were randomly allocated by names drawn from a hat in a public assembly |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | No participants lost to follow-up. |

Clasen 2004c BOL

| Methods | RCT | |
|---------------|---|--|
| Participants | Number: 50 households with 280 persons, of which 32 (11%) were under age 5 Inclusion criteria: all households in the community | |
| Interventions | Household gravity water filter system using imported ceramic filter elements Primary drinking supply | |
| Outcomes | 1. Period prevalence of diarrhoea (7-day recall) among householders assessed at approximately 6-week intervals | |

Clasen 2004c BOL (Continued)

| Notes | Location: rural Bolivia |
|-------|-----------------------------|
| | Length: 6 months |
| | Publication status: journal |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|--|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Households were randomly allocated by lottery, half to an intervention group and half to a control group |
| Allocation concealment (selection bias) | Low risk | Households were randomly allocated by lottery, half to an intervention group and half to a control group |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Less than 1% participants lost to follow-up. |

Clasen 2005 COL

| Methods | RCT |
|---------------|---|
| Participants | Number: 140 children under 5, 680 people, 140 households Inclusion criteria: all households in the community |
| Interventions | 1. Ceramic water filter |
| Outcomes | Diarrhoea prevalence during previous seven days Water quality |
| Notes | Location: three rural villages in Colombia Length: six months Publication status: journal |
| Risk of bias | |

Clasen 2005 COL (Continued)

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Public lottery. |
| Allocation concealment (selection bias) | Low risk | Lottery conducted at each study site to randomly allocate households |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 5% of households lost to follow-up. |

Colford 2002 USA

| Methods | RCT | | |
|---------------|--|-----------------------|--|
| Participants | Number: 236 people from 77 households Inclusion criteria: families were required to own their own homes, use municipal tap water as their main drinking water and have no seriously immunocompromised household members | | |
| Interventions | Household reverse osmosis filters Primary drinking supply | | |
| Outcomes | Incidence of watery diarrhoea Gastrointestinal illness and various other symptoms Water consumption Effectiveness of blinding | | |
| Notes | Location: urban community in California, USA Length: 4 months Publication status: journal | | |
| Risk of bias | | | |
| Bias | Authors' judgement | Support for judgement | |

Colford 2002 USA (Continued)

| Random sequence generation (selection bias) | Low risk | Two random sequences generated to allocated households to intervention or control groups |
|---|--------------|---|
| Allocation concealment (selection bias) | Low risk | Two random sequences generated to allocated households to intervention or control groups |
| Comparability of characteristics | Unclear risk | Irrelevant to study design. |
| Contemporaneous data collection | Unclear risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | One investigator, not involved in analyses prepared coded labels for the placebo and active devices |
| | | |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Triple-blinded. |

Colford 2005 USA

| Methods | RCT |
|---------------|--|
| Participants | Number: 50 HIV+ people, all over 30 years Inclusion criteria: confirmed HIV+ status, uses tap water 75% of the time, no children residing in the home |
| Interventions | 1. Countertop water filtration device |
| Outcomes | Episodes of "highly credible gastrointestinal illness" Diarrhoea episodes calculated |
| Notes | Location: San Francisco, USA Length: 12 months Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Computer generated random numbers. |
| Allocation concealment (selection bias) | Low risk | The manufacturer provided a list of device serial numbers and their corresponding active/sham status to facilitate device assign- |

Colford 2005 USA (Continued)

| | | ment. All study participants, the study investigators (including clinic personnel and those performing data analysis) and the device installer were blinded throughout the trial as to device assignment |
|---|----------|--|
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | All study participants, the study investigators (including clinic personnel and those performing data analysis) and the device installer were blinded throughout the trial as to device assignment |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | All study participants, the study investigators (including clinic personnel and those performing data analysis) and the device installer were blinded throughout the trial as to device assignment |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 10% participants withdrew from study (mixed from active and sham devices) |

Colford 2009 USA

| Methods | Randomized controlled (crossover) trial | |
|---------------|--|--|
| Participants | Number: 988 people, 714 households Inclusion criteria: households were eligible if they had one or more persons 55 or older | |
| Interventions | 1. Countertop water filtration and UV device | |
| Outcomes | Episodes of "highly credible gastrointestinal illness" Diarrhoea episodes calculated | |
| Notes | Location: Sonoma County, USA Length: 13.5 months Publication status: journal | |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Households were block-randomized in blocks of 10, with an equal probability of receiving either a sham or an active device |
| Allocation concealment (selection bias) | Low risk | Households were block-randomized in blocks of 10, with an equal probability of receiving either a sham or an active device |

Colford 2009 USA (Continued)

| Comparability of characteristics | Low risk | Irrelevant to study design. |
|---|-----------|---|
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | All study staff involved in installation and contact with participants were blinded to device assignments throughout the trial |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Assessors blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | "Among households initially assigned to receive an active device, 89% completed cycle 1 and 83% also completed cycle 2; among households initially assigned to receive a sham device, 90% completed cycle 1 and 82% also completed cycle 2" |

Conroy 1996 KEN

| Methods | RCT | |
|---------------|--|--|
| Participants | Number: 206 Maasai children aged 5 to 16 years in 3 adjoining areas of single province Inclusion criteria: all households in the village | |
| Interventions | Solar disinfection in plastic bottles at household level Primary drinking supply | |
| Outcomes | 1. Period prevalence of diarrhoea | |
| Notes | Location: single province of rural Kenya Length: 12 weeks Publication status: journal | |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Interventions assigned by alternate household. |
| Allocation concealment (selection bias) | High risk | Interventions assigned by alternate household. |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |

Conroy 1996 KEN (Continued)

| Blinding of participants and personnel (performance bias) All outcomes | High risk | Not blinded. |
|--|-----------|-----------------------|
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | No loss to follow-up. |

Conroy 1999 KEN

| Methods | RCT | |
|---------------|---|--|
| Participants | Number: 349 Maasai children < 6 years in 140 households Inclusion criteria: all households in the village | |
| Interventions | Solar disinfection in plastic bottles at household level Primary drinking supply | |
| Outcomes | 1. Period prevalence of diarrhoea | |
| Notes | Location: rural Kenya Length: 1 year Publication status: journal | |

Risk of bias

Cochrane Collaboration.

| Bias | Authors' judgement | Support for judgement |
|--|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Interventions assigned by alternate household. |
| Allocation concealment (selection bias) | High risk | Interventions assigned by alternative household. |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Not blinded. |

Conroy 1999 KEN (Continued)

| Incomplete outcome data (attrition bias) All outcomes | High risk | > 20% children lost to follow-up. |
|---|-----------|-----------------------------------|
|---|-----------|-----------------------------------|

Crump 2005a KEN

| Methods | RCT |
|---------------|---|
| Participants | Number: 6650 persons of all ages in 604 family compounds Inclusion criteria: family compounds with at least 1 child < 2 years and likely to be using highly turbid source water |
| Interventions | Sodium hypochlorite used at household level Primary drinking water supply |
| Outcomes | Longitudinal prevalence (weeks with diarrhoea/weeks of observation) among all ages Breastfeeding and consumption of food and water for children < 2 years Deaths Use of intervention Mothers' knowledge of and acceptance of intervention (weeks 5 and 15) Microbial water quality and turbidity Mothers' knowledge of and attitudes to intervention |
| Notes | Location: 49 rural villages in western Kenya Length: 20 weeks Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|-----------------------------|
| Random sequence generation (selection bias) | Unclear risk | Insufficient detail. |
| Allocation concealment (selection bias) | Unclear risk | Insufficient detail. |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |

Crump 2005a KEN (Continued)

Crump 2005b KEN

| Methods | See Crump 2005a KEN |
|---------------|---|
| Participants | As above |
| Interventions | Flocculant-disinfectant sachets used at household level Primary drinking water supply |
| Outcomes | As above |
| Notes | As above |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|-------------------------------------|
| Random sequence generation (selection bias) | Unclear risk | Insufficient detail. |
| Allocation concealment (selection bias) | Unclear risk | Insufficient detail. |
| Comparability of characteristics | Low risk | Irrelevant to study design, |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | 82% participants lost to follow-up. |

Doocy 2006 LBR

| RCT |
|---|
| Number: 2191 persons of all ages (1138 intervention, 1053 controls), of which 735 are children < 5 (395 intervention, 340 controls) Inclusion criteria: households in settlement area not using treated water for drinking |
|] |

Doocy 2006 LBR (Continued)

| Interventions | Flocculant-disinfectant sachets used at household level, plus water storage vessel Primary drinking supply; also received vessel |
|---------------|--|
| Outcomes | Longitudinal prevalence (days with diarrhoea/total days of observation) Prevalence of bloody diarrhoea Utilization and acceptability data from exit survey |
| Notes | Location: Liberian camp for displaced persons Length: 12 weeks Publication status: journal |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Random division of households by blocks and subsections. |
| Allocation concealment (selection bias) | Low risk | Households were systematically selected based on their assigned plot number |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 1% of households lost to follow-up. |

du Preez 2008 ZAF/ZWE

| Methods | RCT |
|---------------|--|
| Participants | Number: 115 children < 5 years Inclusion criteria: households were randomly selected from a list of eligible households from an earlier study: if they had no in-house piped water, and if they had at least one child 12 to 24 months of age |
| Interventions | Household commercial ceramic filter using imported components (60 children) Primary drinking supply (55 children) |

du Preez 2008 ZAF/ZWE (Continued)

| Outcomes | Incidence of diarrhoea Incidence of bloody diarrhoea and non-bloody diarrhoea Microbiological water quality |
|----------|---|
| Notes | Location: rural South Africa and Zimbabwe Length: 6 months Publication status: journal |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Unclear risk | Reported to be randomized, but no description of method of randomization process |
| Allocation concealment (selection bias) | Unclear risk | Insufficient detail. |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Unclear risk | Insufficient detail. |

du Preez 2010 ZAF

| Methods | RCT |
|---------------|---|
| Participants | Number: 824 children, 649 households Inclusion criteria: households were eligible if they had no in-house piped water, and if they had at least one child over 6 months and under 5 years |
| Interventions | SODIS (438 children) Primary drinking supply (386 children) |
| Outcomes | Incidence of dysentery Incidence of non-dysentery diarrhoea |

du Preez 2010 ZAF (Continued)

| Notes | Location: four peri-urban districts of Gauteng Province, South Africa Length: 12 months |
|-------|--|
| | Publication status: journal |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|--|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Random number table. |
| Allocation concealment (selection bias) | Low risk | This table was not available to field workers until after the sample frame was drawn up |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | 13% of children lost to follow-up. |

du Preez 2011 KEN

| Methods | RCT |
|---------------|---|
| Participants | Number: 1089 children, 765 households Inclusion criteria: eligible households stored water in containers in-house, did not have a drinking water tap in the house or yard, and had at least one child (but not more than 5) between 6 months and 5 years old residing in the house |
| Interventions | SODIS (404 households) Primary drinking supply (361) |
| Outcomes | Episodes of dysentery and non-dysentery diarrhoea Height-for-age and weight-for-age Microbial water quality |
| Notes | Location: three urban slums, three rural areas near Nakuru, Kenya\ Length: 17 months Publication status: journal |

| Risk of bias | | |
|--|--------------------|---|
| Bias | Authors' judgement | Support for judgement |
| Random sequence generation (selection bias) | Low risk | Random numbers between zero and one were generated and allocated to the households. If the random number allocated to a household was less than 0.5 the household was randomized to the test group. If the allocated number was above 0.5 the house was randomized to the control group |
| Allocation concealment (selection bias) | Low risk | Field workers were unaware of how the numbers were allocated |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 4% children lost to follow-up. |

Fabiszewski 2012 HND

| RCT |
|--|
| Number: 230 children < 5, 1020 people, 178 households Inclusion criteria: households were eligible if they had a least one child under 5, did not have year-round access to piped water, and did not use bottled water |
| Biosand filter (90 households) Primary drinking supply (86 households) |
| Incidence of diarrhoea Microbial water quality |
| Location: 11 rural communities in Copan, Honduras Length: six month follow-up Publication status: journal |
| |

Fabiszewski 2012 HND (Continued)

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Random number generation. |
| Allocation concealment (selection bias) | Low risk | No one knew which group they were assigned to until the day before |
| Comparability of characteristics | Low risk | Irrelevant to this study design. |
| Contemporaneous data collection | Low risk | Irrelevant to this study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Less than 1% lost to follow-up. |

Gasana 2002 RWA

| Methods | Quasi-RCT |
|---------------|---|
| Participants | Number: 150 children < 5 years Inclusion criteria: all households with at least one child < 5 |
| Interventions | Improved source: pipes to stand post; sedimentation tank; ceramic filter; storage tank; and communal tap (95 children) Primary drinking supply (55 children) |
| Outcomes | 1. Incidence of diarrhoea |
| Notes | Location: rural Rwanda Length: 24 months Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|-----------------------------|
| Random sequence generation (selection bias) | Low risk | Irrelevant to study design. |
| Allocation concealment (selection bias) | Low risk | Irrelevant to study design. |

Gasana 2002 RWA (Continued)

| Comparability of characteristics | Unclear risk | Not described. |
|---|--------------|-----------------------------|
| Contemporaneous data collection | Unclear risk | Not described. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Irrelevant to study design. |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Irrelevant to study design. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Irrelevant to study design. |

Gruber 2013 MEX

| Methods | RCT |
|---------------|--|
| Participants | Number: 1916 people, 444 households Inclusion criteria: households were eligible if they did not have access to centrally treated drinking water and collected water from local sources year-round |
| Interventions | 1. UV water treatment and storage system (Mesita Azul) |
| Outcomes | Diarrhoea prevalence Microbial water quality |
| Notes | Location: rural Baja California Sur, Mexico Length: 15 months Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Eligible communities assigned a random number between zero and one by an investigator using STATA |
| Allocation concealment (selection bias) | Low risk | Every 2 months another community was randomly allocated to intervention group; no one knew in advance |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |

Gruber 2013 MEX (Continued)

| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
|---|-----------|-------------------------------------|
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | 15% participants lost to follow-up. |

Günther 2013 BEN

| Methods | RCT |
|---------------|--|
| Participants | Number: 364 intervention households; 347 control households Inclusion criteria: all households in intervention villages |
| Interventions | Improved water vessel for fetching Improved water vessel for storing |
| Outcomes | Water quality of stored water Diarrhoea prevalence |
| Notes | Location: rural Benin Length: 3 months Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|--|--------------------|-----------------------------|
| Random sequence generation (selection bias) | Unclear risk | Insufficient detail. |
| Allocation concealment (selection bias) | Unclear risk | Insufficient detail. |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |

Günther 2013 BEN (Continued)

| Incomplete outcome data (attrition bias) All outcomes | High risk | 64% of sample with follow-up data (due to budgetary constraints) |
|---|-----------|--|
|---|-----------|--|

Handzel 1998 BGD

| Methods | RCT |
|---------------|--|
| Participants | Number: 447 children aged 3 to 60 months from 276 households Inclusion criteria: households with children 3 to 60 months of age using municipal water (household taps) as primary source of drinking water which had tested positive at baseline for <i>E. coli</i> |
| Interventions | Household chlorination using sodium hypochlorite solution, special storage vessel and hygiene instruction about why and how to treat water (140 households) Primary drinking supply (136 households) |
| Outcomes | Incidence of diarrhoea Microbial water quality |
| Notes | Location: informal settlement in urban Bangladesh Length: 8 months Publication status: PhD dissertation |

| Bias | Authors' judgement | Support for judgement |
|--|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Lottery. |
| Allocation concealment (selection bias) | Low risk | Consent was obtained from participating households; none knew whether they would be placed into the intervention or comparison group |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |

Handzel 1998 BGD (Continued)

Jain 2010 GHA

| Methods | RCT |
|---------------|---|
| Participants | Number: 549 children under five, 3240 individuals, 240 households Inclusion criteria: households were eligible if there was at least one child < 5 |
| Interventions | Chlorine (NaDCC) tablets (120 households) Placebo-tablets without chlorine (120 households) |
| Outcomes | Diarrhoeal episodes Chlorine residuals Microbiological water quality |
| Notes | Location: peri-urban communities of Tamale, Ghana Length: 12 weeks Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Random number table. |
| Allocation concealment (selection bias) | Low risk | Only technical staff at Medentech, Ltd knew which tablets were placebo and which were NaDCC |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Triple blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Triple blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Less than 1% of households lost to follow-up. |

Jensen 2003 PAK

| Methods | Quasi-RCT |
|---------------|---|
| Participants | Number: 226 children < 5 years of age Inclusion criteria: all households that had children aged less than five years and that primarily obtained drinking-water from the water supply systems |
| Interventions | Village level chlorination of water supply using calcium hypochlorite (82 children) Primary drinking supply (144 children) |
| Outcomes | Incidence of diarrhoea Microbial water quality |
| Notes | Location: 2 villages in Pakistan Length: 6 months Publication status: journal Controlled for sanitation and water storage status of households, and for seasonality |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Irrelevant to study design. |
| Allocation concealment (selection bias) | Low risk | Irrelevant to study design. |
| Comparability of characteristics | Low risk | Water quality at baseline significantly different between intervention and control villages |
| Contemporaneous data collection | Low risk | Data collected at similar points in time. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Irrelevant to study design. |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Irrelevant to study design. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Irrelevant to study design. |

Kirchhoff 1985 BRA

| Methods | RCT |
|---------------|--|
| Participants | Number: 112 persons (all ages) from 20 families Inclusion criteria: households with at least 2 children living at home and using water from pond exclusively |
| Interventions | Household level chlorination with sodium hypochlorite Primary drinking supply |
| Outcomes | Longitudinal prevalence of diarrhoea Microbial water quality Acceptability of intervention to study population |
| Notes | Location: rural Brazil Length: 18 weeks Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | High risk | Sequences could be related to outcomes (eligible households which agreed to participate were enrolled) |
| Allocation concealment (selection bias) | High risk | Sequences could be related to outcomes (eligible households which agreed to participate were enrolled) |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Study staff and participants blinded (placebo). |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | Approximately 20% participants lost to follow-up. |

Kremer 2011 KEN

| Methods | RCT |
|---------------|---|
| Participants | Number: 184 springs; 1354 households Inclusion criteria: springs that were not seasonally dry, landownder gave approval to be protected |
| Interventions | 1. Protected springs |
| Outcomes | Diarrhoeal episodes Microbiological water quality |
| Notes | Location: rural western Kenya Length: 2 years Publication status: economics quarterly journal |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Random number generator assigned springs into year of treatment |
| Allocation concealment (selection bias) | Low risk | Random selection of households at each intervention spring. |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 95% of all households were surveyed for baseline and at least two follow-up rounds |

Lindquist 2014a BOL

| Methods | RCT |
|--------------|--|
| Participants | Number: 330 intervention households; 279 control households Inclusion criteria: households: with children less than 60 months of age, in squatter or low-income rental housing, receive their primary drinking/household water from a non- municipal source, and no access to a direct municipal sewer line. Enrollment was limited to one child per household |

Lindquist 2014a BOL (Continued)

| Interventions | 1. Filter |
|---------------|--|
| Outcomes | 1. Diarrhoea period prevalence |
| Notes | Location: rural Bolivia Length: 3 months Publication status: journal |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Random number generator. |
| Allocation concealment (selection bias) | Low risk | Randomization done at neighbourhood level. |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | > 20% lost to follow-up. |

Lindquist 2014b BOL

| Methods | RCT |
|---------------|---|
| Participants | Number: 285 intervention households; 279 control households Inclusion criteria: as above |
| Interventions | Filter WASH behaviour change education |
| Outcomes | As above |
| Notes | As above |
| Risk of bias | |

Lindquist 2014b BOL (Continued)

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Random number generator. |
| Allocation concealment (selection bias) | Low risk | Randomization done at neighbourhood level. |
| Comparability of characteristics | Low risk | Irrelevant to study design. |
| Contemporaneous data collection | Low risk | Irrelevant to study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | > 20% lost to follow-up. |

Luby 2004a PAK

| Methods | Quasi-RCT | | |
|---------------|--|-----------------------|--|
| Participants | Number: 2365 persons < 15 years from 285 households Inclusion criteria: eligible households included at least one child less than five years of age and two children less than 15 years of age, had sufficient water supply for the children to bathe daily, and planned to continue to reside in their homes for at least the ensuing four months | | |
| Interventions | Bleach + regular vessel (640 people) Primary drinking supply (1027 people) | | |
| Outcomes | Longitudinal prevalence of diarrhoea Use of intervention by certain household characteristics | | |
| Notes | Location: 3 neighbourhoods in squatter settlements in Karachi, Pakistan Length: 6 months Publication status: journal | | |
| Risk of bias | | | |
| Bias | Authors' judgement | Support for judgement | |

Luby 2004a PAK (Continued)

| Random sequence generation (selection bias) | Low risk | Irrelevant for study design. |
|---|----------|--|
| Allocation concealment (selection bias) | Low risk | Irrelevant for study design. |
| Comparability of characteristics | Low risk | Baseline characteristics did not differ significantly between groups |
| Contemporaneous data collection | Low risk | Data collected at similar points in time. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Irrelevant for study design. |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Irrelevant for study design. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Irrelevant for study design. |

Luby 2004b PAK

| Methods | See Luby 2004a PAK |
|---------------|---|
| Participants | As above |
| Interventions | Bleach + insulated vessel (697 people) Primary drinking supply (1027 people) |
| Outcomes | As above |
| Notes | As above |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Irrelevant for study design. |
| Allocation concealment (selection bias) | Low risk | Irrelevant for study design. |
| Comparability of characteristics | Low risk | No substantial differences at baseline. |
| Contemporaneous data collection | Low risk | Data collected at similar points in time. |

Luby 2004b PAK (Continued)

| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Irrelevant for study design. |
|--|----------|------------------------------|
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Irrelevant for study design. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Irrelevant for study design. |

Luby 2006a PAK

| Methods | RCT |
|---------------|--|
| Participants | Number: 5520 persons of all ages Inclusion criteria: running water at least one hour twice a week and at least one child under 5 |
| Interventions | Dilute bleach + vessel (1747 people) Primary drinking supply (1852 people) |
| Outcomes | 1. Incidence and longitudinal prevalence of diarrhoea |
| Notes | Location: 47 squatter settlements of Karachi, Pakistan Length: 8 months Publication status: unpublished |

| Bias | Authors' judgement | Support for judgement |
|--|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Computer-generated random number assigned households to groups |
| Allocation concealment (selection bias) | Low risk | Households consented to study before computer randomly assigned them to specific groups |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |

Luby 2006a PAK (Continued)

| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
|--|-----------|---|
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Overall less than 8% of participants lost to follow-up (averaged across all groups) |

Luby 2006b PAK

| Methods | See Luby 2006a PAK | |
|---------------|---|--|
| Participants | As above | |
| Interventions | Flocculant-disinfectant + soap (1806 in flocculant-disinfection group) Primary drinking supply (1852 people) | |
| Outcomes | As above | |
| Notes | As above | |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Computer-generated random number assigned households to groups |
| Allocation concealment (selection bias) | Low risk | Households consented to study before computer randomly assigned them to specific groups |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Overall less than 8% of participants lost to follow-up (averaged across all groups) |

Luby 2006c PAK

| Methods | See Luby 2006a PAK | |
|---------------|--|--|
| Participants | As above | |
| Interventions | Flocculant-disinfectant + vessel (1833 in flocculant-disinfection group) Primary drinking supply (1852 people, 40.0%) | |
| Outcomes | As above | |
| Notes | As above | |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Computer-generated random number assigned households to groups |
| Allocation concealment (selection bias) | Low risk | Households consented to study before computer randomly assigned them to specific groups |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Overall less than 8% of participants lost to follow-up (averaged across all groups) |

Lule 2005 UGA

| Methods | RCT |
|---------------|---|
| Participants | Number: 2201 persons of all ages among 458 households Inclusion criteria: households without access to chlorinated municipal water; at least 1 resident of each household was HIV+ |
| Interventions | Household level chlorination using sodium hypochlorite + special vessel (1097 people) Primary drinking supply (1104 people) Note: hygiene education was provided to both groups |

Lule 2005 UGA (Continued)

| Outcomes | Incidence of diarrhoea Days with diarrhoea (longitudinal prevalence) Days lost from work or school Aetiology of diarrhoea Frequency of clinic visits and hospitalization Mortality |
|----------|---|
| Notes | Location: households in rural Uganda Length: 5 months Publication status: unpublished Succeeded by 18-month RCT that included cotrimoxazole prophylaxis |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Unclear risk | Insufficient detail. |
| Allocation concealment (selection bias) | Unclear risk | Insufficient detail. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Less than 8% of participants lost to follow-up. |

Mahfouz 1995 KSA

| Methods | RCT |
|---------------|---|
| Participants | Number: 311 children < 5 years (among intervention households, among controls) among 171 families Inclusion criteria: households with at least one child less than 5 years of age |
| Interventions | Household level chlorination using calcium hypochlorite (159 children) Primary drinking supply (152 children) |
| Outcomes | 1. Reported cases of diarrhoea in intervention year compared with previous year |

Mahfouz 1995 KSA (Continued)

| Notes | Location: 9 villages in rural Saudi Arabia Length: 6 months |
|-------|---|
| | Publication status: journal |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Unclear risk | No description of randomization process (for villages). No description of how households were chosen |
| Allocation concealment (selection bias) | Unclear risk | No description of how chosen families were selected or contacted |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded |
| Incomplete outcome data (attrition bias) All outcomes | High risk | Large loss to follow-up in intervention and control groups |

Majuru 2011 ZAF

| Methods | Quasi-RCT | |
|---------------|---|--|
| Participants | Number: community 1, 234 individuals; community 2, 173 individuals; reference community, 146 individuals Inclusion criteria: new community level piped water supply | |
| Interventions | Community-level piped water supply (2 communities, 407 individuals) Primary drinking water supply, unimproved sources (1 community, 146 individuals) | |
| Outcomes | 1. Diarrhoeal episodes | |
| Notes | Location: rural, remote communities, Limpopo Province, South Africa Length: approximately 10 months of follow-up Publication status: journal | |

Majuru 2011 ZAF (Continued)

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Irrelevant for study design. |
| Allocation concealment (selection bias) | Low risk | Irrelevant for study design. |
| Comparability of characteristics | Low risk | No substantial differences at baseline. |
| Contemporaneous data collection | Low risk | Data collected at similar points in time. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Irrelevant for study design. |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Irrelevant for study design. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Irrelevant for study design. |

McGuigan 2011 KHM

| Methods | RCT |
|---------------|---|
| Participants | Number: 964 children in 782 households Inclusion criteria: households were eligible if they were permanent residents in the area, had at least one child 6 months to 5 years old, and did not use other methods of household water treatment |
| Interventions | SODIS (407 households, 502 children < 5) Primary drinking water supply (375 households, 426 children < 5) |
| Outcomes | Days of dysentery diarrhoea for < 5s Days of non-dysentery diarrhoea for < 5s |
| Notes | Location: rural communities in Prey Veng and Svey Rieng provinces, Cambodia Length: 12 months Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Randomized raffle system of interested households during initial meeting |

McGuigan 2011 KHM (Continued)

| Allocation concealment (selection bias) | Low risk | Households were randomly allocated to intervention or control groups at community meeting |
|---|-----------|---|
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessor not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 5% of participants had less than 10 months of follow-up. |

Mengistie 2013 ETH

| Methods | RCT |
|---------------|--|
| Participants | Number: 36 clusters, 569 households, 845 children < 5 Inclusion criteria: households were eligible if they had at least one child < 5 |
| Interventions | Chlorine disinfection (WaterGuard) (427 children < 5) Primary drinking supply (422 children < 5) |
| Outcomes | Diarrhoeal episodes for children < 5 Intervention compliance |
| Notes | Location: rural communities, Kersa district, Ethiopia Length: 16 weeks Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Computer generated random sample. |
| Allocation concealment (selection bias) | Low risk | Randomization of clusters done in community meeting. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |

Mengistie 2013 ETH (Continued)

| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
|---|-----------|---|
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 2% to 3% of person-weeks of observation lost. |

Mäusezhal 2009 BOL

| Methods | RCT |
|---------------|--|
| Participants | Number: 484 households, 819 children < 5 Inclusion criteria: communities had to have at least 30 children < 5 and rely on contaminated drinking water sources |
| Interventions | SODIS (11 communities, 262 households, 441 children) Primary drinking water supply, unimproved sources (11 communities, 222 households, 378 children) |
| Outcomes | Diarrhoeal episodes for children < 5 Dysentery episodes for children < 5 |
| Notes | Location: rural Totora District, Cochabamba Department, Bolivia Length: 12 months Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Random assignment during public event. |
| Allocation concealment (selection bias) | Low risk | Balls with community codes inscribed on them were drawn from a box; the first ball drawn would be the intervention community |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |

Mäusezhal 2009 BOL (Continued)

| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
|--|-----------|--|
| Incomplete outcome data (attrition bias) All outcomes | High risk | 21% of person-days of observation missing. |

Opryszko 2010a AFG

| Methods | RCT |
|---------------|--|
| Participants | Number: 553 households, 4507 individuals Inclusion criteria: inadequate access to improved water sources; high areas of diarrhoeal disease according to 2004 census |
| Interventions | Chlorine disinfection (with improved storage vessel); Improved water supply (tube wells); hygiene promotion (261 households, 1958 individuals) Primary drinking supply (292 households, 2549 individuals) |
| Outcomes | Diarrhoea prevalence Dysentery-diarrhoea prevalence |
| Notes | Location: rural communities, Wardak province, Afghanistan Length: 16 months Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|--|--------------------|---------------------------------------|
| Random sequence generation (selection bias) | Low risk | Randomly allocated. |
| Allocation concealment (selection bias) | Low risk | Randomly allocated by numbered lists. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |

Opryszko 2010a AFG (Continued)

| All outcomes | 1 | Low risk | 10% of households data missing at follow-up. |
|--------------|---|----------|--|
|--------------|---|----------|--|

Opryszko 2010b AFG

| Methods | See Opryszko 2010a AFG |
|---------------|--|
| Participants | Number: 600 households, 4,966 individuals Inclusion criteria: inadequate access to improved water sources; high areas of diarrhoeal disease according to 2004 census |
| Interventions | Improved water supply (tube wells) Primary drinking supply (292 households, 2549 individuals) |
| Outcomes | As above |
| Notes | As above |

| Bias | Authors' judgement | Support for judgement |
|--|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Randomly allocated. |
| Allocation concealment (selection bias) | Low risk | Randomly allocated by numbered lists. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 10% of households data missing at follow-up. |

Opryszko 2010c AFG

| Methods | See Opryszko 2010a AFG |
|---------------|--|
| Participants | Number: 591 households, 4575 individuals Inclusion criteria: inadequate access to improved water sources; high areas of diarrhoeal disease according to 2004 census |
| Interventions | Chlorine disinfection (Clorin); Improved storage vessel (299 households, 2026 individuals) Primary drinking supply (292 households, 2549 individuals) |
| Outcomes | As above |
| Notes | As above |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Randomly allocated. |
| Allocation concealment (selection bias) | Low risk | Randomly allocated by numbered lists. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 10% of households data missing at follow-up. |

Patel 2012 KEN

| Methods | RCT |
|---------------|---|
| Participants | Number: 42 schools Inclusion criteria: schools were eligible if they were not near urban centres and did not have pre-existing water-treatment promotion activities |
| Interventions | Chlorine disinfection (WaterGuard); improved vessel (22 schools) Primary drinking supply (20 schools) |

Patel 2012 KEN (Continued)

| Outcomes | Student's knowledge and practice of using WaterGuard Any illness Diarrhoeal illness Acute respiratory illness |
|----------|--|
| Notes | Location: rural Nyanza province, Kenya Length: 2 years Publication status: journal |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|-------------------------------------|
| Random sequence generation (selection bias) | Low risk | Random allocation from census list. |
| Allocation concealment (selection bias) | Low risk | Random allocation from census list. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | 32% students lost to follow-up. |

Peletz 2012 ZMB

| Methods | RCT | |
|---------------|--|--|
| Participants | Number: 120 households, 599 individuals, 121 children < 2 Inclusion criteria: mothers who disclosed their HIV status, had a child 6-12 months old, and permanently resided in the catchment area | |
| Interventions | Filter (LifeStraw® Family); two 5 L storage vessels (61 households, 299 individuals, 61 children < 2) Primary drinking supply (59 households, 300 individuals, 60 children < 2) | |
| Outcomes | Use of filter Microbiological water quality Longitudinal diarrhoeal prevalence | |

Peletz 2012 ZMB (Continued)

| | 4. Weight-for-age Z-scores | |
|--|--|--|
| Notes | Location: two peri-urban neighbourhoods, Chongwe district, Zambia Length: 12 month Publication status: journal | |
| Risk of bias | | |
| Bias | Authors' judgement | Support for judgement |
| Random sequence generation (selection bias) | Low risk | Random number generator. |
| Allocation concealment (selection bias) | Low risk | Randomization conducted by person not involved in study. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |

Quick 1999 BOL

All outcomes

Incomplete outcome data (attrition bias)

| Methods | RCT |
|---------------|--|
| Participants | Number: 791 persons of all ages from 127 households Inclusion criteria: all households in the community |
| Interventions | Household level chlorination + vessel + hygiene education (400 people, 64 households) Primary drinking supply (391 people, 63 households) |
| Outcomes | Mean episodes of diarrhoea per person Microbiological water quality |
| Notes | Location: 2 peri-urban communities in Bolivia Length: 5 months Publication status: journal |

High risk

More than 80% of person-weeks of observation completed.

| Risk of bias | | |
|--|--------------------|--|
| Bias | Authors' judgement | Support for judgement |
| Random sequence generation (selection bias) | Low risk | Randomized by public lottery into two groups. |
| Allocation concealment (selection bias) | Low risk | Randomized by public lottery into two groups. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Less than 10% of participants lost to follow-up. |

Quick 2002 ZMB

| Methods | Quasi-RCT |
|---------------|---|
| Participants | Number: 1584 persons of all ages from 260 households Inclusion criteria: lack of piped water and presence of health clinic in community |
| Interventions | Household level chlorination + vessel + hygiene education (166 households) Primary drinking supply (94 households) |
| Outcomes | Incidence of diarrhoea Microbiological water quality |
| Notes | Location: 2 peri-urban communities in Zambia Length: 3 months Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|------------------------------|
| Random sequence generation (selection bias) | Low risk | Irrelevant for study design. |

Quick 2002 ZMB (Continued)

| Allocation concealment (selection bias) | Low risk | Irrelevant for study design. |
|---|----------|---|
| Comparability of characteristics | Low risk | No substantial differences at baseline. |
| Contemporaneous data collection | Low risk | Data collected at similar points in time. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Irrelevant for study design. |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Irrelevant for study design. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Irrelevant for study design. |

Reller 2003a GTM

| Methods | RCT | |
|---------------|---|--|
| Participants | Number: 492 households Inclusion criteria: household with a child < 12 months or mother in last trimester o pregnancy | |
| Interventions | Flocculant-disinfectant (102 households) Primary drinking supply (96 households) | |
| Outcomes | Incidence of diarrhoea Intervention knowledge and acceptability Microbiological water quality Intervention utilization | |
| Notes | Location: 12 villages in rural Guatemala Length: 12 months Publication status: journal | |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Random number generator assigned eligible households to groups |
| Allocation concealment (selection bias) | Low risk | Random number generator assigned eligible households to groups |

Reller 2003a GTM (Continued)

| Comparability of characteristics | Low risk | Irrelevant for study design. |
|---|-----------|--|
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | Approximately 13% of participants lost to follow-up. |

Reller 2003b GTM

| Methods | See Reller 2003a GTM |
|---------------|---|
| Participants | As above |
| Interventions | Bleach only (97 households) Primary drinking supply (as above) |
| Outcomes | As above |
| Notes | As above |

| Bias | Authors' judgement | Support for judgement |
|--|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Random number generator assigned eligible households to groups |
| Allocation concealment (selection bias) | Low risk | Random number generator assigned eligible households to groups |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |

Reller 2003b GTM (Continued)

| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
|--|-----------|--|
| Incomplete outcome data (attrition bias) All outcomes | High risk | Approximately 13% of participants lost to follow-up. |

Reller 2003c GTM

| Methods | See Reller 2003a GTM |
|---------------|---|
| Participants | As above |
| Interventions | Bleach + vessel (97 households) Primary drinking supply (as above) |
| Outcomes | As above |
| Notes | As above |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Random number generator assigned eligible households to groups |
| Allocation concealment (selection bias) | Low risk | Random number generator assigned eligible households to groups |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | Approximately 13% of participants lost to follow-up. |

Reller 2003d GTM

| Methods | See Reller 2003a GTM |
|---------------|---|
| Participants | As above |
| Interventions | Flocculant-disinfectant + vessel (100 households) Primary drinking supply (as above) |
| Outcomes | As above |
| Notes | As above |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Random number generator assigned eligible households to groups |
| Allocation concealment (selection bias) | Low risk | Random number generator assigned eligible households to groups |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | Approximately 13% of participants lost to follow-up. |

Roberts 2001 MWI

| Methods | RCT |
|---------------|--|
| Participants | Number: 1160 persons of all ages; of these, 208 were children < 5 years Inclusion criteria: all households in refugee camp |
| Interventions | Improved storage: bucket with spout and narrow opening to limit hand entry (310 people including 51 children, 100 households) Primary drinking supply (850 people including 157 children, 300 households) |

Roberts 2001 MWI (Continued)

| Outcomes | Incidence of diarrhoea Microbiological water quality Incidence of diarrhoea by selected environmental factors |
|----------|---|
| Notes | Location: Malawi refugee camp Length: 4 months Publication status: journal |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|--|--------------------|---|
| Random sequence generation (selection bias) | High risk | "One fourth of the interviewed households were selected at ran- dom to receive the improved buckets" |
| Allocation concealment (selection bias) | High risk | "One fourth of the interviewed households were selected at ran- dom to receive the improved buckets" |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | 88.8% of participants lost to follow-up. |

Rodrigo 2011 AUS

| Methods | RCT |
|---------------|---|
| Participants | Number: 300 households, 1352 individuals, 185 children < 5 Inclusion criteria: households were eligible if they use untreated rainwater as their primary drinking source |
| Interventions | Water filters (Freshwater systems) (152 households, 698 individuals) Sham-water filters (148 households, 654 individuals) |
| Outcomes | Episodes of Highly Credible Gastrenteritis Episodes of diarrhoea |

Rodrigo 2011 AUS (Continued)

| Notes | Location: Adelaide, Australia |
|-------|-------------------------------|
| | Length: 12 months |
| | Publication status: journal |
| | <i>'</i> |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Random number sequence by independent researcher. |
| Allocation concealment (selection bias) | Low risk | Random number sequence by independent researcher. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Sham device (placebo) utilised. |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | 31% households lost to follow-up. |

Semenza 1998 UZB

| Methods | RCT |
|---------------|---|
| Participants | Number and inclusion criteria: 1583 persons of all ages from 240 households, half with access to piped water (first control group) and half without (of which 62 received intervention, and 58 served as a second control group); these included 344 children < 5 |
| Interventions | Household level chlorination + vessel + hygiene education Primary drinking supply |
| Outcomes | Incidence of diarrhoea Incidence of diarrhoea by selected household and water management practices |
| Notes | Location: urban Uzbekistan Length: 9.5 weeks Publication status: journal |
| Risk of bias | |

Semenza 1998 UZB (Continued)

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Households randomly selected from map of neighbourhoods. |
| Allocation concealment (selection bias) | Low risk | Households randomly selected from map of neighbourhoods. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Unclear risk | Lost to follow-up not discussed. |

Stauber 2009 DOM

| Methods | RCT |
|---------------|---|
| Participants | Number: 167 households, 907 individuals, 243 children < 5 Inclusion criteria: households were eligible if there was no biosand filter in the house, and there was at least one child < 5 |
| Interventions | Biosand filter (81 households, 447 individuals) Primary drinking supply (86 households, 460 individuals) |
| Outcomes | Diarrhoeal incidence Microbiological water quality |
| Notes | Location: one semi-rural and one urban community, Bonao, Dominican Republic Length: six months follow-up Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Random number generation assigned 50% of households to intervention group |

Stauber 2009 DOM (Continued)

| Allocation concealment (selection bias) | Unclear risk | Households were unaware of whether they would be assigned to the intervention or control group until one week before BSF installation, but it is not clear whether this was foreknowledge of group assignment |
|---|--------------|---|
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 7% participants lost to follow-up. |

Stauber 2012a KHM

| Methods | RCT |
|---------------|---|
| Participants | Number: 189 households, 1147 individuals, 242 children < 5 Inclusion criteria: households were eligible if there was at least one child < 5 |
| Interventions | Plastic Biosand filter (7 villages, 90 households, 546 individuals) Primary drinking supply (6 villages, 99 households, 601 individuals) |
| Outcomes | Diarrhoeal incidence Microbiological water quality |
| Notes | Location: 13 rural communities, Angk Snoul district, Cambodia Length: four months follow-up Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Random number generation assigned 7 of 13 villages to intervention group |
| Allocation concealment (selection bias) | Low risk | All villages were told they would not know to which group they were assigned until halfway through the study (due to surveillance period, pre-intervention) |

Stauber 2012a KHM (Continued)

| Comparability of characteristics | Low risk | Irrelevant for study design. |
|---|-----------|---|
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | 4% of person-observation weeks missing. |

Stauber 2012b GHA

| Methods | RCT |
|---------------|---|
| Participants | Number: 2043 individuals, of which 440 were children < 5, from 260 households Inclusion criteria: households were eligible if there was at least one child < 5 |
| Interventions | Plastic Biosand filter (117 households, 1012 individuals) Primary drinking supply (143 households, 1031 individuals) |
| Outcomes | Diarrhoeal incidence Microbiological water quality |
| Notes | Location: six rural communities, Tamale, Ghana Length: three months follow-up Publication status: journal |

| Bias | Authors' judgement | Support for judgement |
|--|--------------------|--|
| Random sequence generation (selection bias) | Low risk | Random number generator assigned 3 of the 6 villages to the intervention group |
| Allocation concealment (selection bias) | Unclear risk | Not discussed. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |

Stauber 2012b GHA (Continued)

| Blinding of outcome assessment (debias) All outcomes | etection High risk | Assessors not blinded. |
|--|--------------------|---|
| Incomplete outcome data (attrition All outcomes | n bias) Low risk | Less than 3% of households lost to follow-up. |

Tiwari 2009 KEN

| Methods | RCT | |
|---------------|---|--|
| Participants | Number: 387 individuals, of which 114 were children < 5, from 60 households Inclusion criteria: households were eligible if they had at least one child < 3, used rive water as their primary or secondary drinking water source, stable residence for next 12 months, and indicators of lower socio-economic status | |
| Interventions | Biosand filter (30 households, 118 children) Primary drinking water supply (30 households, 104 children) | |
| Outcomes | Microbiological water quality Diarrhoea prevalence in children | |
| Notes | Location: rural households in River Njoro watershed, Nakuru and Molo districts, Kenya Length: six months Publication status: journal | |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Unclear risk | No description of randomization process. |
| Allocation concealment (selection bias) | Unclear risk | No description of steps to conceal allocation. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |

Tiwari 2009 KEN (Continued)

| Incomplete outcome data (attrition bias) All outcomes | Low risk | After randomization, 75 (93%) and 79 (92%) of BSF and control households, respectively, completed the study |
|---|----------|---|
| All outcomes | | tiof flousefloids, respectively, completed the study |

Torun 1982 GTM

| Methods | Quasi-RCT | |
|---------------|---|--|
| Participants | Number: 2103 persons of all ages from 2 villages Inclusion critera: all households within 2 villages | |
| Interventions | Source protection (spring), chlorination facilities, "adequate storage", and water mains with faucets to yards of intervention village (1006 people) Primary drinking supply (1097 people) | |
| Outcomes | 1. Incidence of diarrhoea | |
| Notes | Location: 2 small villages in Guatemala Length: 12 months Publication status: book | |

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Irrelevant for study design. |
| Allocation concealment (selection bias) | Low risk | Irrelevant for study design. |
| Comparability of characteristics | Low risk | No substantial differences at baseline. |
| Contemporaneous data collection | Low risk | Data collected at similar points in time. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Irrelevant for study design. |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Irrelevant for study design. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Irrelevant for study design. |

URL 1995a GTM

| Methods | RCT | |
|---------------|--|--|
| Participants | Number: 1120 children < 5 years (265 and 289 allocated to the water quality intervention arms, 297 to an education only arm, and 269 to the control arm) from 680 families from three demographic regions Inclusion criteria: households must have children < 5 and have indicators of low sociole economic status and microbiological contamination of water source | |
| Interventions | Locally fabricated ceramic filters (265 children or 23.6%) Primary drinking supply (269 children) | |
| Outcomes | Incidence of diarrhoea Nutritional status (weight/age) | |
| Notes | Location: 3 demographic regions of Guatemala Length: 12 months Publication status: unpublished | |

| Bias | Authors' judgement | Support for judgement |
|--|--------------------|--|
| Random sequence generation (selection bias) | Unclear risk | Reported to be randomized, but no description of method of randomization process |
| Allocation concealment (selection bias) | Unclear risk | No description of allocation concealment. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | Participants not blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Unclear risk | Not discussed. |

URL 1995b GTM

| Methods | See URL 1995a GTM | |
|---------------|--|--|
| Participants | As above | |
| Interventions | Locally fabricated ceramic filters + hygiene education Primary drinking supply (as above) | |
| Outcomes | As above | |
| Notes | As above | |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence generation (selection bias) | Unclear risk | Reported to be randomized, but no description of method of randomization process |
| Allocation concealment (selection bias) | Unclear risk | No description of allocation concealment. |
| Comparability of characteristics | Low risk | Irrelevant for study design. |
| Contemporaneous data collection | Low risk | Irrelevant for study design. |
| Blinding of participants and personnel (performance bias) All outcomes | High risk | No placebo used. |
| Blinding of outcome assessment (detection bias) All outcomes | High risk | Assessors not blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Unclear risk | Not discussed. |

Xiao 1997 CHN

| Methods | Quasi-RCT | |
|---------------|---|--|
| Participants | Number: 4649 people of all ages Inclusion criteria: all households within villages | |
| Interventions | Improved water supply + sanitation + hygiene education (2363 people) Primary drinking supply (2286 people) | |
| Outcomes | Incidence of diarrhoea | |

Xiao 1997 CHN (Continued)

| Notes | Location: 2 villages in rural China Length: 3 years Publication status: journal |
|-------|---|
| | |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence generation (selection bias) | Low risk | Irrelevant for study design. |
| Allocation concealment (selection bias) | Low risk | Irrelevant for study design. |
| Comparability of characteristics | Low risk | No substantial differences at baseline. |
| Contemporaneous data collection | Low risk | Data collected at similar points in time. |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Irrelevant for study design. |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Irrelevant for study design. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Irrelevant for study design. |

Characteristics of excluded studies [ordered by study ID]

| Study | Reason for exclusion | | | | |
|----------------|--|--|--|--|--|
| Ahoyo 2011 | Allocation was neither randomized nor quasi-randomized. | | | | |
| Aiken 2011 | Allocation was neither randomized nor quasi-randomized. | | | | |
| Alexander 2013 | Outcome measures did not include diarrhoea. | | | | |
| Arnold 2009 | Allocation was neither randomized nor quasi-randomized. | | | | |
| Arnold 2012a | Comment paper. | | | | |
| Arnold 2013 | Design paper. | | | | |
| Asaolu 2002 | Allocation was neither randomized nor quasi-randomized; outcome measures did not include diarrhoea | | | | |

| Aziz 1990 BGD | The intervention included the provision of sanitation facilities | | | |
|-----------------------|--|--|--|--|
| Azurin 1974 | Outcome measures did not include diarrhoea. | | | |
| Bahl 1976 | Allocation was neither randomized nor quasi-randomized. | | | |
| Bajer 2012 | Allocation was neither randomized nor quasi-randomized, outcome not diarrhoea | | | |
| Barreto 2007 | Allocation was neither randomized nor quasi-randomized. | | | |
| Barzilay 2011 | Allocation was neither randomized nor quasi-randomized. | | | |
| Bersh 1985 | Allocation was neither randomized nor quasi-randomized. | | | |
| Boubacar 2014 | Allocation was neither randomized nor quasi-randomized. | | | |
| Brown 2012a | Modelling paper. | | | |
| Capuno 2011 | Allocation was neither randomized nor quasi-randomized. | | | |
| Cavallaro 2011 | Allocation was neither randomized nor quasi-randomized, outcomes did not include diarrhoea | | | |
| Chang 2012 | Outcomes did not include diarrhoea. | | | |
| Chongsuvivatwong 1994 | Allocation was neither randomized nor quasi-randomized. | | | |
| Christen 2011 | Allocation was neither randomized nor quasi-randomized, outcomes did not include diarrhoea | | | |
| Clasen 2012 | No water quality intervention. | | | |
| Colford 2005 | Outcomes did not include diarrhoea. | | | |
| Colwell 2003 | Outcomes did not include diarrhoea. | | | |
| Conroy 2001 | Outcomes did not include diarrhoea. | | | |
| Coulliette 2013 | Allocation was neither randomized nor quasi-randomized, outcomes did not include diarrhoea | | | |
| Crump 2007 | Allocation was neither randomized nor quasi-randomized, outcomes did not include diarrhoea | | | |
| Davis 2011 | Outcomes did not include diarrhoea. | | | |
| Deb 1986 | Outcomes did not include diarrhoea. | | | |
| Denslow 2010 | Allocation was neither randomized nor quasi-randomized. | | | |
| Devoto 2011 | Intervention did not affect water quality. | | | |

| Dorevitch 2011 | Outcomes did not include diarrhoea. | | | | |
|----------------------|--|--|--|--|--|
| Dreibelbis 2014a KEN | School-based study. | | | | |
| Dreibelbis 2014b KEN | School-based study. | | | | |
| Dreibelbis 2014c KEN | School-based study. | | | | |
| du Preez 2012 | Response to comments. | | | | |
| Eisenberg 2006 | Study on risk assessment. | | | | |
| Enger 2012 | Modelling paper. | | | | |
| Esrey 1988 | Allocation was neither randomized nor quasi-randomized. | | | | |
| Fewtrell 1994 | Allocation was neither randomized nor quasi-randomized, outcomes did not include diarrhoea | | | | |
| Fewtrell 1997 | Allocation was neither randomized nor quasi-randomized, outcomes did not include diarrhoea | | | | |
| Firth 2010 | Outcomes did not include diarrhoea. | | | | |
| Fisher 2011 | Allocation was neither randomized nor quasi-randomized. | | | | |
| Freeman 2012 | Outcomes did not include diarrhoea. | | | | |
| Freeman 2014a KEN | School-based study. | | | | |
| Freeman 2014b KEN | School-based study. | | | | |
| Freeman 2014c KEN | School-based study. | | | | |
| Fry 2010 | Modelling paper. | | | | |
| Galiani 2009 | Allocation was neither randomized nor quasi-randomized | | | | |
| Garrett 2008 KEN | The intervention included the provision of sanitation facilities | | | | |
| Ghannoum 1981 | Allocation was neither randomized nor quasi-randomized, outcomes did not include diarrhoea | | | | |
| Gorelick 2011 | Allocation was neither randomized nor quasi-randomized. | | | | |
| Greene 2012 | Outcome not diarrhoea, see Freeman 2012. | | | | |
| Gómez-Couso 2012 | Allocation was neither randomized nor quasi-randomized, outcome not diarrhoea | | | | |
| Habib 2013 | Water quality intervention applied once children had experienced diarrhoea | | | | |

| Harris 2009 | Allocation was neither randomized nor quasi-randomized. | | | | | |
|-----------------|---|--|--|--|--|--|
| Harshfield 2012 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Hartinger 2011 | Design paper. | | | | | |
| Hartinger 2012 | Outcome measures did not include diarrhoea. | | | | | |
| Hellard 2001 | Outcome measures did not include diarrhoea. | | | | | |
| Hoque 1996 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Huda 2012 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Hunter 2010 | Allocation was neither randomized nor quasi-randomized | | | | | |
| Iijima 2001 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Islam 2011 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Jensen 2002 | Outcome not diarrhoea. | | | | | |
| Kariuki 2012 | Intervention not water. | | | | | |
| Karon 2011 | Outcome not diarrhoea. | | | | | |
| Keraita 2007 | Outcome not diarrhoea. | | | | | |
| Khan 1984 | Outcome not diarrhoea. | | | | | |
| Luby 2008 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Luoto 2011 | Outcome not diarrhoea. | | | | | |
| Luoto 2012 | Outcome not diarrhoea. | | | | | |
| Macy 1998 | Allocation was neither randomized nor quasi-randomized; intervention not an improvement in water quality; outcome not diarrhoea | | | | | |
| McCabe 1957 | Intervention not an improvement in water quality. | | | | | |
| Mertens 1990 | Allocation was neither randomized nor quasi-randomized, intervention not an improvement in water quality; outcome not diarrhoea | | | | | |
| Messou 1997 | The intervention included the provision of sanitation facilities | | | | | |
| Mäusezahl 2003 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| | | | | | | |

| Nanan 2003 | Allocation was neither randomized nor quasi-randomized. | | | | | |
|-----------------|---|--|--|--|--|--|
| Nerkar 2014 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Nnane 2011 | Allocation was neither randomized nor quasi-randomized, no intervention | | | | | |
| Oluyege 2011 | Allocation was neither randomized nor quasi-randomized, no intervention | | | | | |
| Palit 2012 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Pavlinac 2014 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Payment 1991a | Allocation was neither randomized nor quasi-randomized, outcomes did not include diarrhoea | | | | | |
| Payment 1991b | Outcomes did not include diarrhoea. | | | | | |
| Peletz 2013 | Outcomes did not include diarrhoea. | | | | | |
| Pinfold 1990 | Intervention not an improvement in water quality; outcome not diarrhoea | | | | | |
| Psutka 2012 | Allocation was neither randomized nor quasi-randomized, outcomes did not include diarrhoea | | | | | |
| Rosa 2014 | Outcomes did not include diarrhoea. | | | | | |
| Rose 2006 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Rubenstein 1969 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Russo 2012 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Sathe 1996 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Shah 2012 | Review paper. | | | | | |
| Sharan 2011 | Allocation was neither randomized nor quasi-randomized, outcome not diarrhoea | | | | | |
| Sheth 2010 | Allocation was neither randomized nor quasi-randomized, outcome not diarrhoea | | | | | |
| Shiffman 1978 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Shrestha 2006 | Cost-effectiveness paper. | | | | | |
| Shum 1971 | Allocation was neither randomized nor quasi-randomized, intervention not an improvement in water quality; outcome not diarrhoea | | | | | |
| Sima 2012 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| | | | | | | |

| Sorvillo 1994 | Outcomes did not include diarrhoea. | | | | | |
|------------------|--|--|--|--|--|--|
| Stauber 2013 | Outcomes did not include diarrhoea. | | | | | |
| Sutha 2011 | Review paper. | | | | | |
| Tonglet 1992 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Trivedi 1971 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| VanDerslice 1995 | Allocation was neither randomized nor quasi-randomized, intervention not an improvement in water quality | | | | | |
| Varghese 2002 | Allocation was neither randomized nor quasi-randomized. | | | | | |
| Wiedenmann 2006 | Intervention not an improvement in water quality. | | | | | |
| Wolf 2014 | Review. | | | | | |
| Wood 2012 | Qualitative study. | | | | | |
| Wu 2011 | Allocation was neither randomized nor quasi-randomized. | | | | | |

Characteristics of ongoing studies [ordered by study ID]

Chlorination, Dhaka

| Trial name or title | Impact of Low-Cost In-Line Chlorination Systems in Urban Dhaka on Water Quality and Child Health | | | | | | |
|---------------------|---|--|--|--|--|--|--|
| Methods | RCT | | | | | | |
| Participants | All poor households, with at least one child under five, that access one of 160 studied shared water point Dhaka | | | | | | |
| Interventions | In-line chlorination | | | | | | |
| Outcomes | Water quality, diarrhoea in children under five, weight of children, cost of instilling and maintaining system, hospital visits, health care expenditures, other household expenditures | | | | | | |
| Starting date | Early 2015 | | | | | | |
| Contact information | | | | | | | |
| Notes | Funded by SIEF, World Bank | | | | | | |

WASH-B, Bangladesh

| Trial name or title | WASH Benefits Bangladesh: A Cluster Randomized Controlled Trial of the Benefits of Water, Sanitation, Hygiene Plus Nutrition Interventions on Child Growth | | | | | |
|---------------------|--|--|--|--|--|--|
| Methods | Parallel, cluster-RCT | | | | | |
| Participants | Estimated enrolment: 5040 | | | | | |
| Interventions | Water quality: Storage vessel and chlorine tablets. Sanitation: a) a sani-scoop hoe dedicated to the removal of human and animal faeces from the compound, b) plastic child potties for children ages 6 months and older until they are using the latrine, and c) a new or upgraded dual pit latrine for each household in the compound. The behavior change components of the intervention will emphasize the use of the latrine for defecation and the safe disposal of faeces in the compound courtyard to prevent contact with young children. Handwashing: The hardware components of the Bangladesh handwashing intervention include two handwashing stations. The first station will be located in the kitchen (location of food preparation), and will include a 16 L bucket with a tap fitting, a stool, bowl and soapy water bottle. The second station will be located near the toilet, and will include a 40 L bucket with tap fitting, stool, bowl and soapy water bottle. The study will provide detergent soap to families free of charge to replenish the soapy water bottles. The behavior change component of the intervention will focus messaging for handwashing at two critical times: after defecation and before food preparation. Nutrition: Mothers will be encouraged to exclusively breastfeed their children through age 6 months. When newborns reach 6 months of age, mothers will be encouraged to continue breastfeeding their children until 24 months, and will receive education about supplementing breastfeeding with healthy complementary foods following infant and young child feeding best practice guidelines from Unicef and the WHO. From ages 6 to 24 months, study children will receive a daily lipid-based nutritional supplement (LNS) that has been developed and tested through the iLiNS project. | | | | | |
| Outcomes | Length-for-Age Z-scores (time frame: measured 24 months after intervention) (Designated as safety issue: no). Child's recumbent length, standardized to Z-scores using the WHO 2006 growth standards. Diarrhoea Prevalence (time frame: measured 12- and 24-months after intervention). | | | | | |
| Starting date | May 2012 | | | | | |
| Contact information | International Centre for Diarrhoeal Disease Research, Bangladesh | | | | | |
| Notes | | | | | | |

WASH-B, Kenya

| Trial name or tit | tle WASH-Benefits study, Kenya |
|-------------------|--|
| Methods | Parallel, cluster-RCT |
| Participants | Estimated: 8000 |
| Interventions | 1. Water quality: intervention villages will receive chlorine dispensers at spring water sources. After filling their plastic jerry can of water from the source, users can place the jerry can under the dispenser, and turn a knob to release 3 mL of chlorine. Behavior change messages will focus on the consistent provision of treated |

WASH-B, Kenya (Continued)

| | water to all children living in the household. 2. Sanitation: a) a sani-scoop hoe dedicated to the removal of human and animal faeces from the compound; b) plastic child potties for children ages 6 months and older until they are using the latrine; and c) a new or upgraded pit latrine for each household in the compound. If participants have a latrine, its structure will be improved if necessary. Plastic slabs will be installed to improve mud or wood floors, and the intervention delivery team will make sure that all latrine structures have walls, doors, roofs that ensure safety and privacy. The behaviour change components of the intervention will emphasize the use of the latrine for defecation and the safe disposal of faeces in the compound courtyard to prevent contact with young children. 3. Handwashing: two handwashing stations in the compound of each respondent, one near the latrine, and one by the cooking area. The handwashing stations are constructed from locally available materials and are of a dual tippy-tap design with independent pedals attached to 5 L jerry cans of clean water and jugs of soapy water. The behavior change component of the intervention will focus messaging for handwashing at two critical times: after defecation and before food preparation. 4. Nutrition: mothers will be encouraged to exclusively breastfeed their children through to 6 months of age. When newborns reach 6 months of age, mothers will be encouraged to continue breastfeeding their children until 24 months, and will receive education about supplementing breastfeeding with healthy complementary foods following infant and young child feeding best practice guidelines from Unicef and WHO. From ages six to 24 months, study children will receive a daily lipid-based nutritional supplement (LNS) that has been developed and tested through the iLiNS project. |
|---------------------|--|
| Outcomes | Length-for-age Z-scores (time frame: measured 24 months after intervention) (designated as safety issue: no). Child's recumbent length, standardized to Z-scores using the WHO 2006 growth standards. Diarrhoea prevalence (time frame: measured 12 and 24 months after intervention) |
| Starting date | September 2012 |
| Contact information | Innovations for Poverty Action, Kenya |
| Notes | |

DATA AND ANALYSES

Comparison 1. Water quality intervention versus control

| Outcome or subgroup title | No. of studies | No. of participants | Statistical method | Effect size |
|---------------------------------|----------------|---------------------|-----------------------------|-------------------|
| 1 Diarrhoea: all ages | 64 | 81215 | Risk Ratio (Random, 95% CI) | 0.59 [0.51, 0.69] |
| 1.1 Source water improvement | 6 | 9161 | Risk Ratio (Random, 95% CI) | 0.76 [0.48, 1.19] |
| 1.2 POU treatment | 58 | 72054 | Risk Ratio (Random, 95% CI) | 0.58 [0.48, 0.69] |
| 2 Diarrhoea: children < 5 years | 49 | | Risk Ratio (Random, 95% CI) | 0.61 [0.49, 0.75] |
| 2.1 Source water improvement | 4 | | Risk Ratio (Random, 95% CI) | 0.96 [0.82, 1.12] |
| 2.2 POU treatment | 45 | | Risk Ratio (Random, 95% CI) | 0.58 [0.46, 0.73] |

Comparison 2. Source: water supply improvement versus control

| Outcome or subgroup title | No. of studies | No. of participants | Statistical method | Effect size |
|---|----------------|---------------------|-----------------------------|-------------------|
| 1 Diarrhoea: CBA studies subgrouped by age | 6 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 1.1 Cluster-RCTs | 1 | 3266 | Risk Ratio (Random, 95% CI) | 1.24 [0.98, 1.57] |
| 1.2 CBA studies | 5 | 5895 | Risk Ratio (Random, 95% CI) | 0.68 [0.42, 1.09] |
| 2 Diarrhoea: CBA studies subgrouped by age | 5 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 2.1 All ages | 5 | 5895 | Risk Ratio (Random, 95% CI) | 0.68 [0.42, 1.09] |
| 2.2 < 5 years | 3 | 999 | Risk Ratio (Random, 95% CI) | 0.92 [0.79, 1.07] |

Comparison 3. POU: water chlorination versus control

| Outcome or subgroup title | No. of studies | No. of participants | Statistical method | Effect size |
|---|----------------|---------------------|-----------------------------|-------------------|
| 1 Diarrhoea: subgrouped by study design | 19 | 34694 | Risk Ratio (Random, 95% CI) | 0.72 [0.61, 0.84] |
| 1.1 Cluster-RCTs | 16 | 30746 | Risk Ratio (Random, 95% CI) | 0.77 [0.65, 0.91] |
| 1.2 CBA studies | 3 | 3948 | Risk Ratio (Random, 95% CI) | 0.51 [0.34, 0.75] |
| 2 Diarrhoea: cluster-RCTs: subgrouped by age | 16 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 2.1 All ages | 16 | | Risk Ratio (Random, 95% CI) | 0.77 [0.65, 0.91] |
| 2.2 < 5 years | 15 | | Risk Ratio (Random, 95% CI) | 0.77 [0.64, 0.92] |
| 3 Diarrhoea: cluster-RCTs; subgrouped by adherence | 16 | 30746 | Risk Ratio (Random, 95% CI) | 0.77 [0.65, 0.91] |
| 3.1 Residual chlorine in 86 to 100% of samples | 1 | 276 | Risk Ratio (Random, 95% CI) | 0.78 [0.73, 0.83] |

| 3.2 Residual chlorine in 51 to 85% of samples | 6 | 9994 | Risk Ratio (Random, 95% CI) | 0.60 [0.40, 0.91] |
|--|----|-------|---------------------------------|-------------------|
| 3.3 Residual chlorine in ≤ 50% of samples | 4 | 12613 | Risk Ratio (Random, 95% CI) | 0.90 [0.76, 1.06] |
| 3.4 Residual chlorine not reported | 5 | 7863 | Risk Ratio (Random, 95% CI) | 0.85 [0.65, 1.12] |
| 4 Diarrhoea: cluster-RCTs by risk of bias by blinding of participants | 16 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 4.1 Low risk | 5 | 15867 | Risk Ratio (Random, 95% CI) | 1.07 [0.97, 1.17] |
| 4.2 High risk | 11 | 14879 | Risk Ratio (Random, 95% CI) | 0.68 [0.56, 0.83] |
| 5 Diarrhoea: cluster-RCTs; subgrouped by additional water storage intervention | 16 | | Risk Ratio (Random, 95% CI) | 0.77 [0.65, 0.91] |
| 5.1 Chlorination kit alone | 8 | | Risk Ratio (Random, 95% CI) | 0.75 [0.54, 1.05] |
| 5.2 Chlorination kit plus | 8 | | Risk Ratio (Random, 95% CI) | 0.80 [0.66, 0.97] |
| water storage | O | | rusik rukto (rumqom, 7576 Ci) | 0.00 [0.00, 0.57] |
| 6 Diarrhoea: cluster-RCTs: subgrouped by sufficiency of | 16 | 30746 | Risk Ratio (Random, 95% CI) | 0.77 [0.65, 0.91] |
| water quantity | 2 | 5252 | D' 1 D ' (D 1 050(CD) | 0.00 [0.60, 1.17] |
| 6.1 Sufficient | 3 | 5352 | Risk Ratio (Random, 95% CI) | 0.90 [0.69, 1.17] |
| 6.2 Insufficient | 2 | 3499 | Risk Ratio (Random, 95% CI) | 0.91 [0.66, 1.26] |
| 6.3 Unclear | 11 | 21895 | Risk Ratio (Random, 95% CI) | 0.67 [0.50, 0.88] |
| 7 Diarrhoea: cluster-RCTs: subgrouped by water source | 16 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 7.1 Improved water source | 3 | 5880 | Risk Ratio (Random, 95% CI) | 0.82 [0.59, 1.14] |
| 7.2 Unimproved water source | 13 | 24866 | Risk Ratio (Random, 95% CI) | 0.75 [0.59, 0.93] |
| 8 Diarrhoea: cluster-RCTs: subgrouped by sanitation level | 16 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 8.1 Improved sanitation | 3 | 4876 | Risk Ratio (Random, 95% CI) | 0.64 [0.44, 0.92] |
| 8.2 Unimproved sanitation | 6 | 17352 | Risk Ratio (Random, 95% CI) | 0.81 [0.63, 1.05] |
| 8.3 Unclear | 7 | 8518 | Risk Ratio (Random, 95% CI) | 0.75 [0.54, 1.05] |
| 9 Diarrhoea: cluster-RCTs; | 16 | | Risk Ratio (Random, 95% CI) | 0.77 [0.65, 0.91] |
| subgrouped by length of follow-up | 10 | | 14011 1410 (14114011, 75770 G2) | 01/7 [0105, 0151] |
| $9.1 \leq 3$ months | 2 | | Risk Ratio (Random, 95% CI) | 0.42 [0.06, 3.03] |
| 9.2 > 3 to 6 months | 7 | | Risk Ratio (Random, 95% CI) | 0.71 [0.51, 0.99] |
| 9.3 > 6 to 12 months | 5 | | Risk Ratio (Random, 95% CI) | 0.82 [0.71, 0.96] |
| 9.4 > 12 months | 2 | | Risk Ratio (Random, 95% CI) | 0.99 [0.66, 1.48] |

Comparison 4. POU: flocculation and disinfection versus control

| Outcome or subgroup title | No. of studies | No. of participants | Statistical method | Effect size |
|--|----------------|---------------------|-----------------------------|-------------------|
| 1 Diarrhoea: cluster-RCTs | 7 | | Risk Ratio (Random, 95% CI) | 0.48 [0.20, 1.16] |
| 2 Diarrhoea: cluster-RCTs: subgrouped by age; excluding Doocy 2006 LBR | 6 | | Risk Ratio (Random, 95% CI) | Subtotals only |

| 2.1 All ages | 6 | 11788 | Risk Ratio (Random, 95% CI) | 0.69 [0.58, 0.82] |
|---|----------|--------------|--|--|
| 2.2 < 5 | 6 | 0 | Risk Ratio (Random, 95% CI) | 0.71 [0.61, 0.84] |
| 3 Diarrhoea: cluster-RCTs: | 7 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| subgrouped by adherence | | | , | , |
| 3.2 Residual chlorine 51 to | 1 | 2191 | Risk Ratio (Random, 95% CI) | 0.12 [0.11, 0.13] |
| 85% | | | | |
| 3.3 Residual chlorine < 50% | 4 | 6914 | Risk Ratio (Random, 95% CI) | 0.76 [0.67, 0.85] |
| 3.4 Residual chlorine not | 2 | 4874 | Risk Ratio (Random, 95% CI) | 0.41 [0.26, 0.64] |
| measured | | | | |
| 4 Diarrhoea: cluster-RCTs: | 7 | | Risk Ratio (Random, 95% CI) | 0.48 [0.20, 1.16] |
| subgrouped by additional | | | | |
| storage container | | | | |
| 4.1 No storage container | 2 | | Risk Ratio (Random, 95% CI) | 0.81 [0.69, 0.95] |
| 4.2 Storage container | 5 | | Risk Ratio (Random, 95% CI) | 0.39 [0.14, 1.08] |
| 5 Diarrhoea: cluster-RCTs: | 7 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| subgrouped by sufficiency of | | | | |
| water quantity | | | | |
| 5.1 Sufficient | 1 | 3401 | Risk Ratio (Random, 95% CI) | 0.62 [0.47, 0.82] |
| 5.2 Insufficient | 2 | 5454 | Risk Ratio (Random, 95% CI) | 0.31 [0.05, 2.09] |
| 5.3 Unclear | 4 | 5124 | Risk Ratio (Random, 95% CI) | 0.64 [0.49, 0.85] |
| 6 Diarrhoea: cluster-RCTs: | 7 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| subgrouped by water source | | // | | |
| 6.1 Improved water source | 2 | 4874 | Risk Ratio (Random, 95% CI) | 0.41 [0.26, 0.64] |
| 6.2 Unimproved water source | 4 | 5704 | Risk Ratio (Random, 95% CI) | 0.49 [0.14, 1.68] |
| 6.3 Unclear | 1 | 3401 | Risk Ratio (Random, 95% CI) | 0.62 [0.47, 0.82] |
| 7 Diarrhoea: cluster-RCTs: | 7 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| subgrouped by sanitation level | 2 | 4874 | Did Deci (Decident 050/ CI) | 0 /1 [0 2/ 0 //] |
| 7.1 Improved sanitation7.2 Unimproved sanitation | 2 2 | 48/4 5592 | Risk Ratio (Random, 95% CI) Risk Ratio (Random, 95% CI) | 0.41 [0.26, 0.64] 0.27 [0.05, 1.36] |
| 7.3 Unclear | 3 | 3513 | Risk Ratio (Random, 95% CI) | 0.27 [0.05, 1.36] |
| 8 Diarrhoea: cluster-RCTs: | <i>7</i> | 13979 | Risk Ratio (Random, 95% CI) | 0.48 [0.20, 1.16] |
| subgrouped by length of | / | 139/9 | Risk Ratio (Random, 93% CI) | 0.48 [0.20, 1.16] |
| follow-up | | | | |
| $8.1 \le 3$ months | 2 | 5592 | Risk Ratio (Random, 95% CI) | 0.27 [0.05, 1.36] |
| 8.2 > 3 to 6 months | 1 | 3263 | Risk Ratio (Random, 95% CI) | 0.83 [0.67, 1.03] |
| 8.3 > 6 to 12 months | 4 | 5124 | Risk Ratio (Random, 95% CI) | 0.64 [0.49, 0.85] |
| 5.5 × 6 to 12 months | | 7121 | 14011 14010 (140100111, 7770 O1) | 0.01 [0.15, 0.05] |

Comparison 5. POU: filtration versus control

| Outcome or subgroup title | No. of studies | No. of participants | Statistical method | Effect size |
|---|----------------|---------------------|-----------------------------|-------------------|
| 1 Diarrhoea: cluster-RCTs: subgrouped by age | 23 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 1.1 All ages | 23 | | Risk Ratio (Random, 95% CI) | 0.48 [0.38, 0.59] |
| 1.2 < 5 years | 19 | | Risk Ratio (Random, 95% CI) | 0.49 [0.38, 0.62] |
| 2 Diarrhoea: cluster-RCTs: subgrouped by type of filtration | 23 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 2.1 Ceramic filter | 12 | 5763 | Risk Ratio (Random, 95% CI) | 0.39 [0.29, 0.53] |

| 2.2 Sand filtration | 5 | 5504 | Risk Ratio (Random, 95% CI) | 0.47 [0.39, 0.57] |
|--|---------|------|---|---|
| 2.3 LifeStraw® | 3 | 3259 | Risk Ratio (Random, 95% CI) | 0.69 [0.51, 0.93] |
| 2.4 Plumbed | 3 | 1056 | Risk Ratio (Random, 95% CI) | 0.73 [0.52, 1.03] |
| 3 Diarrhoea: cluster-RCTs: | 23 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| subgrouped by blinding of | | | ((, / / / / / | 0.0000000000000000000000000000000000000 |
| participants | | | | |
| 3.1 Low risk | 5 | | Risk Ratio (Random, 95% CI) | 0.80 [0.68, 0.94] |
| 3.2 High risk | 18 | | Risk Ratio (Random, 95% CI) | 0.41 [0.33, 0.52] |
| 4 Diarrhoea: ceramic filter studies | 12 | 5763 | Risk Ratio (Random, 95% CI) | 0.39 [0.29, 0.53] |
| subgrouped by water source | | | | |
| 4.1 Improved water source | 8 | 3607 | Risk Ratio (Random, 95% CI) | 0.33 [0.23, 0.46] |
| 4.2 Unimproved water source | 4 | 2156 | Risk Ratio (Random, 95% CI) | 0.54 [0.48, 0.61] |
| 5 Diarrhoea: ceramic filter studies | 12 | 5763 | Risk Ratio (Random, 95% CI) | 0.39 [0.29, 0.53] |
| subgrouped by sanitation level | | | | |
| 5.1 Improved sanitation | 7 | 4198 | Risk Ratio (Random, 95% CI) | 0.49 [0.38, 0.64] |
| 5.2 Unimproved sanitation | 4 | 1491 | Risk Ratio (Random, 95% CI) | 0.35 [0.22, 0.56] |
| 5.3 Unclear | 1 | 74 | Risk Ratio (Random, 95% CI) | 0.21 [0.18, 0.25] |
| 6 Diarrhoea: sand filter studies: | 5 | | Risk Ratio (Random, 95% CI) | 0.47 [0.39, 0.57] |
| subgrouped by water source | | | | |
| 6.1 Improved water source | 2 | | Risk Ratio (Random, 95% CI) | 0.50 [0.33, 0.75] |
| 6.2 Unimproved water source | 2 | | Risk Ratio (Random, 95% CI) | 0.44 [0.25, 0.76] |
| 6.3 Unclear | 1 | | Risk Ratio (Random, 95% CI) | 0.47 [0.37, 0.60] |
| 7 Diarrhoea: sand filter studies: | 5 | | Risk Ratio (Random, 95% CI) | 0.47 [0.39, 0.57] |
| subgrouped by sanitation level | | | | |
| 7.1 Improved sanitation | 1 | | Risk Ratio (Random, 95% CI) | 0.47 [0.37, 0.60] |
| 7.2 Unimproved sanitation | 3 | | Risk Ratio (Random, 95% CI) | 0.48 [0.34, 0.68] |
| 7.3 Unclear | 1 | | Risk Ratio (Random, 95% CI) | 0.46 [0.22, 0.96] |
| 8 Diarrhoea: cluster-RCTs: | 23 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| subgrouped by adherence | | | | |
| 8.1 86 to 100% | 12 | 7300 | Risk Ratio (Random, 95% CI) | 0.43 [0.34, 0.55] |
| 8.2 51 to 85% | 4 | 2346 | Risk Ratio (Random, 95% CI) | 0.56 [0.33, 0.95] |
| $8.3 \le 50\%$ | 1 | 1516 | Risk Ratio (Random, 95% CI) | 0.75 [0.60, 0.94] |
| 8.4 Not reported | 6 | 4420 | Risk Ratio (Random, 95% CI) | 0.46 [0.28, 0.75] |
| 9 Diarrhoea: cluster-RCTs: | 19 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| subgrouped by additional water | | | | |
| storage intervention | 0 | | D' 1 D .' (D 1 050/ CI) | 0 (0 [0 (0 0 7() |
| 9.1 Filtration alone | 8 | | Risk Ratio (Random, 95% CI) | 0.60 [0.48, 0.76] |
| 9.2 Filtration plus storage | 11 | | Risk Ratio (Random, 95% CI) | 0.38 [0.29, 0.49] |
| 10 Diarrhoea: cluster-RCTs; | 23 | | Risk Ratio (Random, 95% CI) | 0.48 [0.38, 0.59] |
| subgrouped by length of follow-up | | | | |
| | 2 | | Risk Ratio (Random, 95% CI) | 0.26 [0.20, 0.33] |
| $10.1 \le 3$ months $10.2 > 3$ to 6 months | 3 11 | | Risk Ratio (Random, 95% CI) Risk Ratio (Random, 95% CI) | 0.26 [0.20, 0.33] 0.52 [0.44, 0.60] |
| 10.2 > 5 to 6 months | 8 | | Risk Ratio (Random, 95% CI) | 0.52 [0.44, 0.00] |
| 10.4 > 12 months | 1 | | Risk Ratio (Random, 95% CI) | 0.87 [0.74, 1.02] |
| 10.1 × 12 monus | 1 | | Mon Natio (Nationalli, 7) /0 Ci) | 0.0/ [0./4, 1.02] |

| Outcome or subgroup title | No. of studies | No. of participants | Statistical method | Effect size |
|--|----------------|---------------------|-----------------------------|-------------------|
| 1 Diarrhoea: subgrouped by study | 6 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| design | | | | |
| 1.1 Cluster-RCTs | 4 | 3460 | Risk Ratio (Random, 95% CI) | 0.62 [0.42, 0.94] |
| 1.2 Quasi-RCTs | 2 | 555 | Risk Ratio (Random, 95% CI) | 0.82 [0.69, 0.97] |
| 2 Diarrhoea: cluster-RCTs; subgrouped by age | 4 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 2.1 All ages | 4 | | Risk Ratio (Random, 95% CI) | 0.62 [0.42, 0.94] |
| 2.2 < 5 | 3 | | Risk Ratio (Random, 95% CI) | 0.55 [0.34, 0.91] |
| 3 Diarrhoea: cluster-RCTs; subgrouped by adherence | 4 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 3.1 86 to 100% | 1 | 928 | Risk Ratio (Random, 95% CI) | 0.37 [0.29, 0.47] |
| 3.2 51 to 85% | 0 | 0 | Risk Ratio (Random, 95% CI) | 0.0 [0.0, 0.0] |
| $3.3 \le 50\%$ | 2 | 1443 | Risk Ratio (Random, 95% CI) | 0.80 [0.57, 1.11] |
| 3.4 Not reported | 1 | 1089 | Risk Ratio (Random, 95% CI) | 0.73 [0.63, 0.85] |
| 4 Diarrhoea: cluster-RCTs; subgrouped by sufficiency of water supply level | 4 | 3460 | Risk Ratio (Random, 95% CI) | 0.62 [0.42, 0.94] |
| 4.1 Sufficient | 2 | 1443 | Risk Ratio (Random, 95% CI) | 0.80 [0.57, 1.11] |
| 4.3 Unclear | 2 | 2017 | Risk Ratio (Random, 95% CI) | 0.52 [0.27, 1.02] |
| 5 Diarrhoea: cluster-RCTs; subgrouped by water source | 4 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 5.1 Improved water source | 1 | 718 | Risk Ratio (Random, 95% CI) | 0.64 [0.39, 1.05] |
| 5.2 Unimproved water source | 3 | 2742 | Risk Ratio (Random, 95% CI) | 0.62 [0.38, 1.02] |
| 6 Diarrhoea: cluster-RCTs; subgrouped by sanitation level | 4 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 6.1 Improved sanitation | 0 | 0 | Risk Ratio (Random, 95% CI) | 0.0 [0.0, 0.0] |
| 6.2 Unimproved sanitation | 2 | 1653 | Risk Ratio (Random, 95% CI) | 0.57 [0.24, 1.39] |
| 6.3 Unclear | 2 | 1807 | Risk Ratio (Random, 95% CI) | 0.72 [0.63, 0.83] |
| 7 Diarrhoea: cluster-RCTs; | 4 | 3460 | Risk Ratio (Random, 95% CI) | 0.62 [0.42, 0.94] |
| subgrouped by length of follow-up | | | | |
| 7.2 > 6 to 12 months | 3 | 2371 | Risk Ratio (Random, 95% CI) | 0.59 [0.32, 1.09] |
| 7.3 > 12 months | 1 | 1089 | Risk Ratio (Random, 95% CI) | 0.73 [0.63, 0.85] |

Comparison 7. POU: UV disinfection versus control

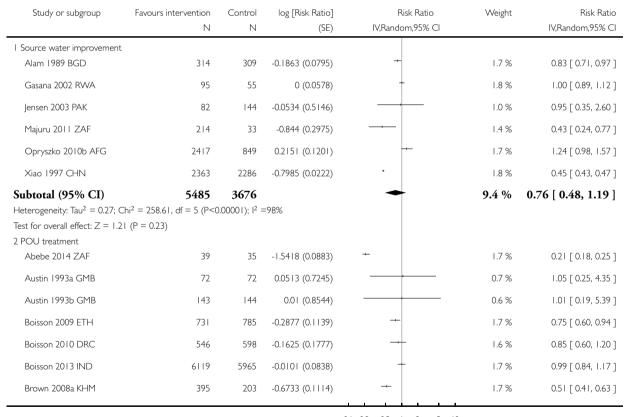
| Outcome or subgroup title | No. of studies | No. of participants | Statistical method | Effect size |
|---------------------------|----------------|-----------------------------|--------------------|----------------|
| 1 Diarrhoea: cluster-RCT | 1 | Risk Ratio (Random, 95% CI) | | Subtotals only |

| Outcome or subgroup title | No. of studies | No. of participants | Statistical method | Effect size |
|--|----------------|---------------------|-----------------------------|-------------------|
| 1 Diarrhoea: cluster-RCTs: subgrouped by age | 2 | | Risk Ratio (Random, 95% CI) | Subtotals only |
| 1.1 All ages | 2 | | Risk Ratio (Random, 95% CI) | 0.91 [0.74, 1.11] |
| 1.2 < 5 | 1 | | Risk Ratio (Random, 95% CI) | 0.69 [0.47, 1.01] |

Analysis I.I. Comparison I Water quality intervention versus control, Outcome I Diarrhoea: all ages.

Comparison: I Water quality intervention versus control

Outcome: I Diarrhoea: all ages



0.1 0.2 0.5 1 2 5 10

Favours intervention Favours control

(Continued ...)

| Study or subgroup | Favours intervention | Control N | log [Risk Ratio] (SE) | Risk Ratio IV.Random,95% Cl | Weight | (Continued) Risk Ratio IV,Random,95% CI |
|-----------------------|----------------------|--------------|--------------------------|--------------------------------|--------|--|
| Brown 2008b KHM | 398 | 200 | -0.5447 (0.1073) | - | 1.7 % | 0.58 [0.47, 0.72] |
| Chiller 2006 GTM | 1702 | 1699 | -0.478 (0.1426) | - | 1.7 % | 0.62 [0.47, 0.82] |
| Clasen 2004b BOL | 210 | 107 | -0.6733 (0.3023) | | 1.4 % | 0.51 [0.28, 0.92] |
| Clasen 2004c BOL | 140 | 140 | -0.5852 (0.1332) | | 1.7 % | 0.56 [0.43, 0.72] |
| Clasen 2005 COL | 415 | 265 | -0.803 (0.2132) | | 1.6 % | 0.45 [0.29, 0.68] |
| Colford 2002 USA | 118 | 118 | -0.6061 (0.1939) | | 1.6 % | 0.55 [0.37, 0.80] |
| Colford 2005 USA | 24 | 26 | -0.2399 (0.3853) | | 1.3 % | 0.79 [0.37, 1.67] |
| Colford 2009 USA | 385 | 385 | -0.1393 (0.0826) | + | 1.7 % | 0.87 [0.74, 1.02] |
| Conroy 1996 KEN | 108 | 98 | -0.2194 (0.147) | | 1.7 % | 0.80 [0.60, 1.07] |
| Conroy 1999 KEN | 175 | 174 | -0.1924 (0.1092) | - | 1.7 % | 0.82 [0.67, 1.02] |
| Crump 2005a KEN | 2249 | 1138 | -0.2614 (0.1072) | - | 1.7 % | 0.77 [0.62, 0.95] |
| Crump 2005b KEN | 2124 | 1139 | -0.1863 (0.1101) | + | 1.7 % | 0.83 [0.67, 1.03] |
| Doocy 2006 LBR | 1138 | 1053 | -2.1203 (0.0408) | + | 1.8 % | 0.12 [0.11, 0.13] |
| du Preez 2008 ZAF/ZWE | 60 | 55 | -1.5606 (0.2855) | | 1.4 % | 0.21 [0.12, 0.37] |
| du Preez 2010 ZAF | 383 | 335 | -0.4463 (0.2527) | | 1.5 % | 0.64 [0.39, 1.05] |
| du Preez 2011 KEN | 555 | 534 | -0.3147 (0.0752) | + | 1.7 % | 0.73 [0.63, 0.85] |
| Fabiszewski 2012 HND | 532 | 488 | -0.4748 (0.2905) | | 1.4 % | 0.62 [0.35, 1.10] |
| Gruber 2013 MEX | 957 | 956 | -0.2357 (0.2437) | -+ | 1.5 % | 0.79 [0.49, 1.27] |
| Günther 2013 BEN | 364 | 347 | -0.0192 (0.0761) | + | 1.7 % | 0.98 [0.85, 1.14] |
| Handzel 1998 BGD | 140 | 136 | -0.2485 (0.0338) | + | 1.8 % | 0.78 [0.73, 0.83] |
| Jain 2010 GHA | 1610 | 1630 | 0.1113 (0.068) | + | 1.8 % | 1.12 [0.98, 1.28] |
| Kirchhoff 1985 BRA | 56 | 56 | 0.0677 (0.0993) | + | 1.7 % | 1.07 [0.88, 1.30] |
| Lindquist 2014a BOL | 330 | 140 | -1.5606 (0.1717) | - | 1.6 % | 0.21 [0.15, 0.29] |
| Lindquist 2014b BOL | 285 | 139 | -1.3093 (0.1045) | - | 1.7 % | 0.27 [0.22, 0.33] |
| Luby 2004a PAK | 697 | 513 | -1.204 (0.2806) | | 1.5 % | 0.30 [0.17, 0.52] |
| Luby 2004b PAK | 640 | 514 | -0.5108 (0.1716) | | 1.6 % | 0.60 [0.43, 0.84] |
| Luby 2006a PAK | 1747 | 617 | -0.7985 (0.3123) | | 1.4 % | 0.45 [0.24, 0.83] |
| Luby 2006b PAK | 1806 | 617 | -0.7985 (0.3062) | | 1.4 % | 0.45 [0.25, 0.82] |
| Luby 2006c PAK | 1833 | 618 | -1.0217 (0.3465) | | 1.3 % | 0.36 [0.18, 0.71] |
| Lule 2005 UGA | 1097 | 1104 | -0.2231 (0.0489) | + | 1.8 % | 0.80 [0.73, 0.88] |
| Mahfouz 1995 KSA | 159 | 152 | -0.5978 (0.305) | | 1.4 % | 0.55 [0.30, 1.00] |

Favours intervention $\begin{tabular}{ll} Favours control \\ \hline & (Continued \dots) \end{tabular}$

0.1 0.2 0.5 1 2 5 10

| Study or subgroup | Favours intervention N | Control N | log [Risk Ratio] (SE) | Risk Ratio IV,Random,95% CI | Weight | (Continued Risk Ratio IV,Random,95% CI |
|---|--|-------------------------------|--------------------------|--------------------------------|---------|---|
| McGuigan 2011 KHM | 426 | 502 | -0.9943 (0.1243) | - | 1.7 % | 0.37 [0.29, 0.47] |
| Mengistie 2013 ETH | 427 | 422 | -0.8348 (0.0663) | + | 1.8 % | 0.43 [0.38, 0.49] |
| Mäusezhal 2009 BOL | 376 | 349 | -0.0943 (0.1796) | - | 1.6 % | 0.91 [0.64, 1.29] |
| Opryszko 2010c AFG | 2026 | 849 | 0.1906 (0.1076) | + | 1.7 % | 1.21 [0.98, 1.49] |
| Peletz 2012 ZMB | 300 | 299 | -0.7765 (0.2181) | _ _ | 1.6 % | 0.46 [0.30, 0.71] |
| Quick 1999 BOL | 400 | 391 | -0.2944 (0.068) | + | 1.8 % | 0.74 [0.65, 0.85] |
| Quick 2002 ZMB | 1000 | 584 | -0.4604 (0.1933) | | 1.6 % | 0.63 [0.43, 0.92] |
| Reller 2003a GTM | 102 | 24 | -0.2357 (0.1151) | - | 1.7 % | 0.79 [0.63, 0.99] |
| Reller 2003b GTM | 97 | 24 | -0.3011 (0.1111) | - | 1.7 % | 0.74 [0.60, 0.92] |
| Reller 2003c GTM | 97 | 24 | -0.0305 (0.1335) | + | 1.7 % | 0.97 [0.75, 1.26 |
| Reller 2003d GTM | 100 | 24 | -0.3011 (0.1221) | | 1.7 % | 0.74 [0.58, 0.94 |
| Roberts 2001 MWI | 310 | 850 | -0.2357 (0.1353) | - | 1.7 % | 0.79 [0.61, 1.03 |
| Rodrigo 2011 AUS | 698 | 654 | -0.1625 (0.2039) | | 1.6 % | 0.85 [0.57, 1.27 |
| Semenza 1998 UZB | 791 | 792 | -1.8971 (0.3704) | | 1.3 % | 0.15 [0.07, 0.31 |
| Stauber 2009 DOM | 447 | 460 | -0.755 (0.1221) | - | 1.7 % | 0.47 [0.37, 0.60 |
| Stauber 2012a KHM | 546 | 601 | -0.8916 (0.2732) | | 1.5 % | 0.41 [0.24, 0.70 |
| Stauber 2012b GHA | 1012 | 1031 | -0.8916 (0.42) | | 1.2 % | 0.41 [0.18, 0.93] |
| Tiwari 2009 KEN | 206 | 181 | -0.7765 (0.3763) | | 1.3 % | 0.46 [0.22, 0.96 |
| URL 1995a GTM | 289 | 134 | -0.755 (0.4476) | | 1.1 % | 0.47 [0.20, 1.13 |
| URL 1995b GTM | 297 | 135 | -1.0498 (0.4931) | | 1.1 % | 0.35 [0.13, 0.92 |
| ubtotal (95% CI) | 40429 | 31625 | | • | 90.6 % | 0.58 [0.48, 0.69] |
| Heterogeneity: Tau ² = 0.47; Ch | ` | 2<0.00001); 1 | 2 =98% | | | |
| est for overall effect: Z = 5.84 Total (95% CI) leterogeneity: Tau ² = 0.36; Chest for overall effect: Z = 6.55 | 45914 mi ² = 2577.10, df = 63 (F | 35301 P<0.00001); I | ² =98% | • | 100.0 % | 0.59 [0.51, 0.69 |

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 0.2
 0.5
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 2
 5
 I0

 Favours intervention
 Favours control

Analysis 1.2. Comparison I Water quality intervention versus control, Outcome 2 Diarrhoea: children < 5 years.

Comparison: I Water quality intervention versus control

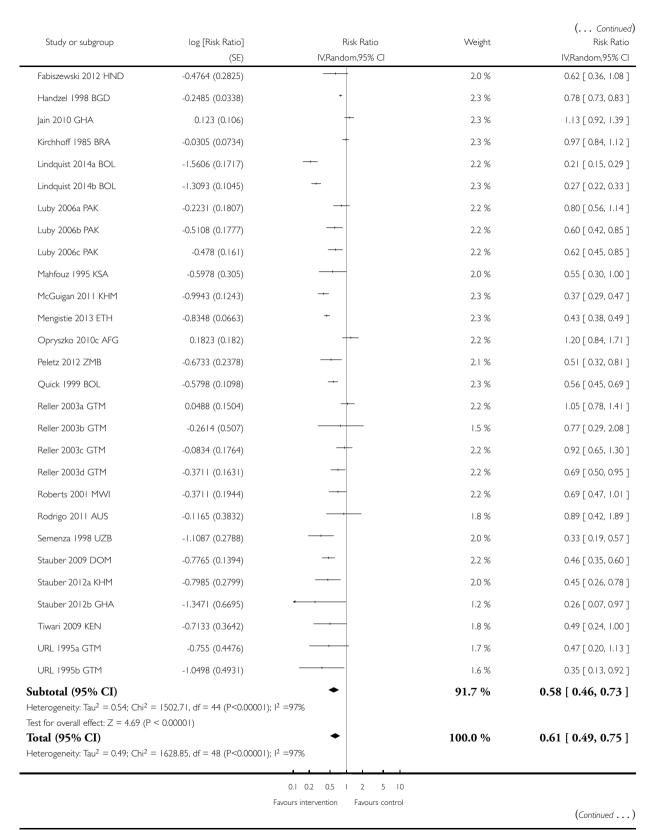
Outcome: 2 Diarrhoea: children < 5 years

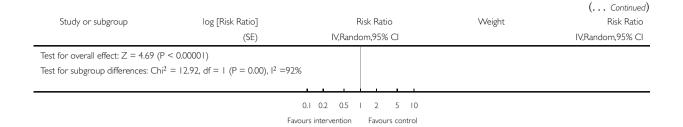
| | (SE) | IV,Random,95% CI | Weight | Risk Ratio IV,Random,95% CI |
|--|---|------------------|---|---|
| Source water improvement | | | | |
| Alam 1989 BGD | -0.1863 (0.0795) | + | 2.3 % | 0.83 [0.71, 0.97] |
| Gasana 2002 RWA | 0 (0.0578) | + | 2.3 % | 1.00 [0.89, 1.12] |
| Jensen 2003 PAK | -0.0534 (0.5146) | | 1.5 % | 0.95 [0.35, 2.60] |
| Opryszko 2010b AFG | 0.1989 (0.1784) | +- | 2.2 % | 1.22 [0.86, 1.73] |
| subtotal (95% CI) eterogeneity: $Tau^2 = 0.01$: $Chi^2 = 0.01$ | 5.63. df = 3 (P = 0.13): l ² =47% | • | 8.3 % | 0.96 [0.82, 1.12] |
| st for overall effect: $Z = 0.53$ (P = | , | | | |
| POU treatment | | | | |
| Austin 1993a GMB | 0.0513 (0.7245) | | 1.1 % | 1.05 [0.25, 4.35] |
| Austin 1993b GMB | 0.01 (0.8544) | | 0.9 % | 1.01 [0.19, 5.39] |
| Boisson 2009 ETH | -0.0305 (0.1888) | + | 2.2 % | 0.97 [0.67, 1.40] |
| Boisson 2010 DRC | -0.1625 (0.2129) | + | 2.1 % | 0.85 [0.56, 1.29] |
| Boisson 2013 IND | -0.0513 (0.0941) | + | 2.3 % | 0.95 [0.79, 1.14] |
| Brown 2008a KHM | -0.5447 (0.177) | | 2.2 % | 0.58 [0.41, 0.82] |
| Brown 2008b KHM | -0.4308 (0.1764) | | 2.2 % | 0.65 [0.46, 0.92] |
| Chiller 2006 GTM | -0.462 (0.1345) | | 2.2 % | 0.63 [0.48, 0.82] |
| Clasen 2004b BOL | -0.3827 (0.2878) | -+ | 2.0 % | 0.68 [0.39, 1.20] |
| Clasen 2004c BOL | -0.8867 (0.4638) | | 1.6 % | 0.41 [0.17, 1.02] |
| Clasen 2005 COL | -0.4589 (0.1722) | | 2.2 % | 0.63 [0.45, 0.89] |
| Crump 2005a KEN | -0.1863 (0.1151) | + | 2.3 % | 0.83 [0.66, 1.04] |
| Crump 2005b KEN | -0.2877 (0.1206) | - | 2.3 % | 0.75 [0.59, 0.95] |
| Doocy 2006 LBR | -2.5257 (0.0601) | • | 2.3 % | 0.08 [0.07, 0.09] |
| du Preez 2008 ZAF/ZWE | -1.5606 (0.2855) | | 2.0 % | 0.21 [0.12, 0.37] |
| du Preez 2010 ZAF | -0.4463 (0.2527) | | 2.1 % | 0.64 [0.39, 1.05] |
| du Preez 2011 KEN | -0.3147 (0.0752) | + | 2.3 % | 0.73 [0.63, 0.85] |
| ty: Tau ² = 0.01; Chi ² = 9 all effect: Z = 0.53 (P = ment 93a GMB 93b GMB 9 | 0.0513 (0.7245) 0.01 (0.8544) -0.0305 (0.1888) -0.1625 (0.2129) -0.0513 (0.0941) -0.5447 (0.177) -0.4308 (0.1764) -0.462 (0.1345) -0.3827 (0.2878) -0.8867 (0.4638) -0.4589 (0.1722) -0.1863 (0.1151) -0.2877 (0.1206) -2.5257 (0.0601) -1.5606 (0.2855) -0.4463 (0.2527) | | 1.1 % 0.9 % 2.2 % 2.1 % 2.3 % 2.2 % 2.2 % 2.2 % 2.2 % 2.3 % 2.3 % 2.3 % 2.3 % 2.3 % 2.3 % 2.1 % | 1.05 [0.25, 4.35] 1.01 [0.19, 5.39] 0.97 [0.67, 1.40] 0.85 [0.56, 1.29] 0.95 [0.79, 1.14] 0.58 [0.41, 0.82] 0.65 [0.46, 0.92] 0.63 [0.48, 0.82] 0.68 [0.39, 1.20] 0.41 [0.17, 1.02] 0.63 [0.45, 0.89] 0.83 [0.66, 1.04] 0.75 [0.59, 0.95] 0.08 [0.07, 0.09] 0.21 [0.12, 0.37] 0.64 [0.39, 1.05] |

0.1 0.2 0.5 I 2 5 I0

Favours intervention Favours control

(Continued ...)

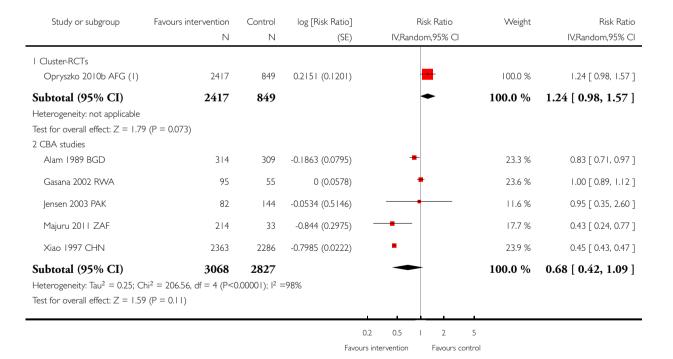




Analysis 2.1. Comparison 2 Source: water supply improvement versus control, Outcome I Diarrhoea: CBA studies subgrouped by age.

Comparison: 2 Source: water supply improvement versus control

Outcome: I Diarrhoea: CBA studies subgrouped by age

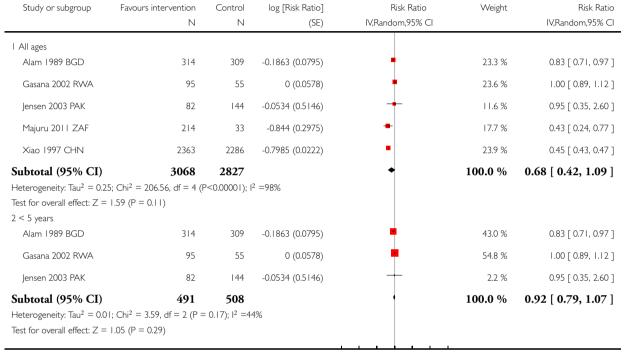


 $[\]hbox{(1) Opryszko 2010-ii AFG: Provided one well per 25 households providing 25 litres/person/day}$

Analysis 2.2. Comparison 2 Source: water supply improvement versus control, Outcome 2 Diarrhoea: CBA studies subgrouped by age.

Comparison: 2 Source: water supply improvement versus control

Outcome: 2 Diarrhoea: CBA studies subgrouped by age

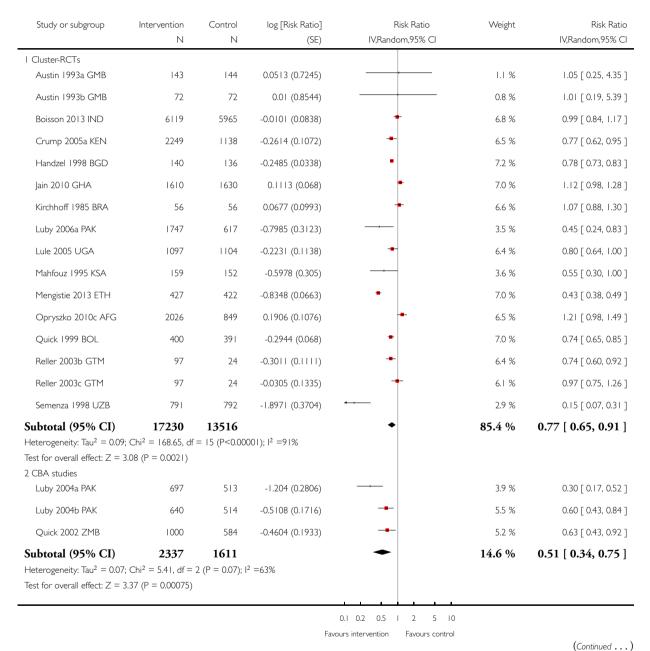


0.001 0.01 0.1 1 10 100 1000 Favours intervention Favours control

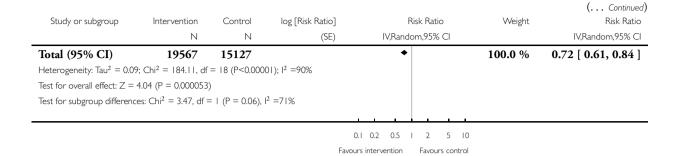
Analysis 3.1. Comparison 3 POU: water chlorination versus control, Outcome 1 Diarrhoea: subgrouped by study design.

Comparison: 3 POU: water chlorination versus control

Outcome: I Diarrhoea: subgrouped by study design



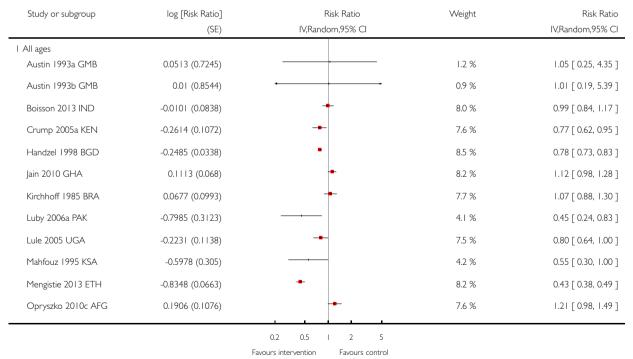
(Continued . .



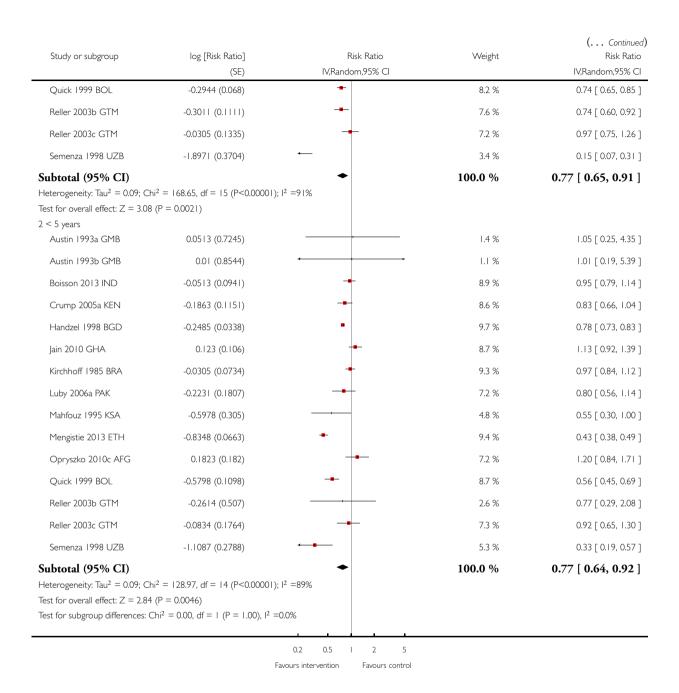
Analysis 3.2. Comparison 3 POU: water chlorination versus control, Outcome 2 Diarrhoea: cluster-RCTs: subgrouped by age.

Comparison: 3 POU: water chlorination versus control

Outcome: 2 Diarrhoea: cluster-RCTs: subgrouped by age



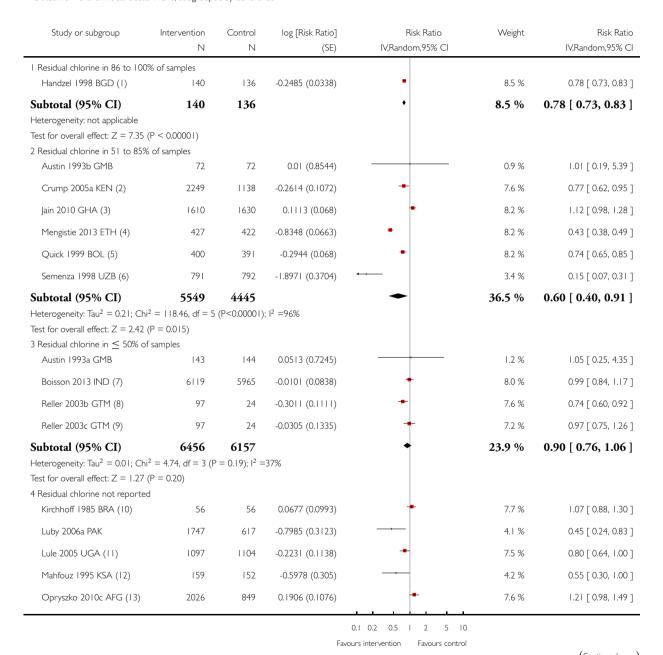
(Continued . . .)



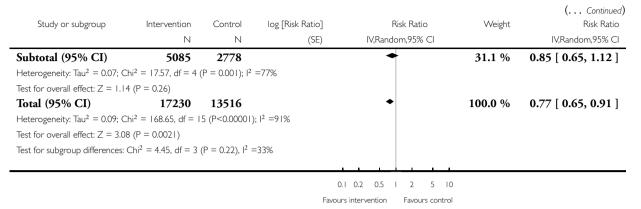
Analysis 3.3. Comparison 3 POU: water chlorination versus control, Outcome 3 Diarrhoea: cluster-RCTs; subgrouped by adherence.

Comparison: 3 POU: water chlorination versus control

Outcome: 3 Diarrhoea: cluster-RCTs; subgrouped by adherence



(Continued . . .)

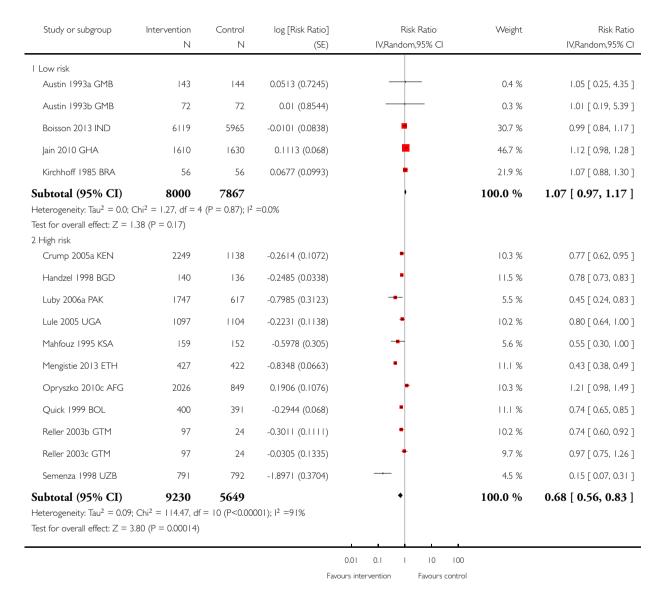


- (1) Handzel 1998 BGD: Free chlorine was measureable in 77% of samples Unclear whether testing was testing was during unannounced visits
- (2) Crump 2005-i KEN: Free chlorine residuals > 0.1 mg/L in 85% of samples during scheduled visits and 61% of samples during unnanounced visits
- (3) Jain 2010 GHA: Free chlorine residuals > 0.2 mg/L in 74-89% of samples Unclear whether testing was during unannounced visits
- (4) Mengistie 2013 ETH: Free chlorine residuals > 0.2 mg/L in 76-77% of samples Testing was during unannounced visits
- (5) Quick 1999 BOL: The proportion of stored water samples with detectable levels of total chlorine increased from 71 % at the time of the first observation to 95% at the final virit
- (6) Semenza 1998 UZB: Chlorine was detected in 73% of household samples at the end of the study.
- (7) Boisson 2013 IND: Free chlorine was measureable in 32% of samples Unclear whether testing was testing was during unannounced visits
- (8) Reller 2003-ii GUA: Participants had free chlorine >0.1 mg/mL in 36% of samples Testing during unannounced visits
- (9) Reller 2003-iii GUA: Participants had free chlorine >0.1 mg/mL in 44% of samples Testing during unannounced visit
- (10) Kirchhoff 1985 BRA: The chlorination was performed daily by blinded health staff.
- (11) Lule 2005 UGA: Compliance not reported
- (12) Mahfouz 1995 KSA: The average free residual chlorine is reported as 0.13 ppm
- (13) Opryszko 2010-iii AFG: Self reported use of Chlorine in the previous two weeks was 82%

Analysis 3.4. Comparison 3 POU: water chlorination versus control, Outcome 4 Diarrhoea: cluster-RCTs by risk of bias by blinding of participants.

Comparison: 3 POU: water chlorination versus control

Outcome: 4 Diarrhoea: cluster-RCTs by risk of bias by blinding of participants



Analysis 3.5. Comparison 3 POU: water chlorination versus control, Outcome 5 Diarrhoea: cluster-RCTs; subgrouped by additional water storage intervention.

Comparison: 3 POU: water chlorination versus control

Outcome: 5 Diarrhoea: cluster-RCTs; subgrouped by additional water storage intervention

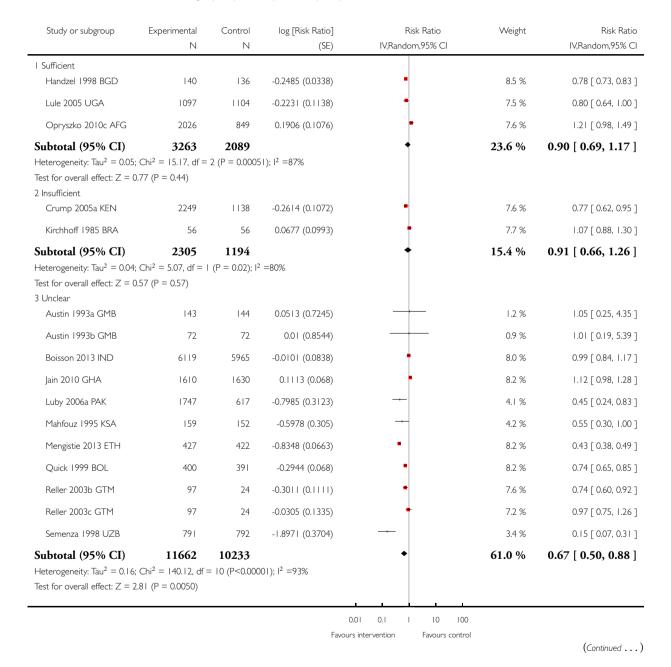
| Study or subgroup | log [Risk Ratio] (SE) | Risk Ratio IV,Random,95% CI | Weight | Risk Ratio IV,Random,95% CI |
|---|---|--|-------------------|--------------------------------|
| I Chlorination kit alone | | | | |
| Austin 1993a GMB | 0.0513 (0.7245) | | 1.2 % | 1.05 [0.25, 4.35] |
| Austin 1993b GMB | 0.01 (0.8544) | | 0.9 % | 1.01 [0.19, 5.39] |
| Boisson 2013 IND (1) | -0.0101 (0.0838) | + | 8.0 % | 0.99 [0.84, 1.17] |
| Crump 2005a KEN (2) | -0.2614 (0.1072) | - | 7.6 % | 0.77 [0.62, 0.95] |
| Kirchhoff 1985 BRA (3) | 0.0677 (0.0993) | + | 7.7 % | 1.07 [0.88, 1.30] |
| Mahfouz 1995 KSA (4) | -0.5978 (0.305) | | 4.2 % | 0.55 [0.30, 1.00] |
| Mengistie 2013 ETH (5) | -0.8348 (0.0663) | | 8.2 % | 0.43 [0.38, 0.49] |
| Reller 2003b GTM (6) | -0.3011 (0.1111) | - | 7.6 % | 0.74 [0.60, 0.92] |
| Subtotal (95% CI) Heterogeneity: $Tau^2 = 0.16$; $Chi^2 = 0.16$ for overall effect: $Z = 1.70$ (P = 2 Chlorination kit plus water storage | 0.090) | 2% | 45.4 % | 0.75 [0.54, 1.05] |
| Handzel 1998 BGD (7) | -0.2485 (0.0338) | • | 8.5 % | 0.78 [0.73, 0.83] |
| Jain 2010 GHA (8) | 0.1113 (0.068) | - | 8.2 % | 1.12 [0.98, 1.28] |
| Luby 2006a PAK | -0.7985 (0.3123) | | 4.1 % | 0.45 [0.24, 0.83] |
| Lule 2005 UGA (9) | -0.2231 (0.1138) | - | 7.5 % | 0.80 [0.64, 1.00] |
| Opryszko 2010c AFG (10) | 0.1906 (0.1076) | - | 7.6 % | 1.21 [0.98, 1.49] |
| Quick 1999 BOL (11) | -0.2944 (0.068) | • | 8.2 % | 0.74 [0.65, 0.85] |
| Reller 2003c GTM (12) | -0.0305 (0.1335) | + | 7.2 % | 0.97 [0.75, 1.26] |
| Semenza 1998 UZB (13) | -1.8971 (0.3704) | | 3.4 % | 0.15 [0.07, 0.31] |
| Subtotal (95% CI) Heterogeneity: Tau² = 0.06; Chi² = Test for overall effect: Z = 2.25 (P = Total (95% CI) Heterogeneity: Tau² = 0.09; Chi² = Test for overall effect: Z = 3.08 (P = | 0.024) 168.65, df = 15 (P<0.00001); l ² | • | 54.6 % 100.0 % | 0.80 [0.66, 0.97] |
| Test for subgroup differences: Chi ² = | | 0.1 0.2 0.5 1 2 5 10 avours intervention Favours control | | |

- (1) Boisson 2013 IND: Free chlorine was measureable in 32% of samples Unclear whether testing was testing was during unannounced visits
- (2) Crump 2005-i KEN: Free chlorine residuals > 0.1 mg/L in 85% of samples during scheduled visits and 61% of samples during unnanounced visits
- (3) Kirchhoff 1985 BRA: The chlorination was performed daily by blinded health staff.
- (4) Mahfouz 1995 KSA: The average free residual chlorine is reported as 0.13 ppm
- (5) Mengistie 2013 ETH: Free chlorine residuals > 0.2 mg/L in 76-77% of samples Testing was during unannounced visits
- (6) Reller 2003-ii GUA: Participants had free chlorine >0.1 mg/mL in 36% of samples Testing during unannounced visits
- (7) Handzel 1998 BGD: Free chlorine was measureable in 77% of samples Unclear whether testing was testing was during unannounced visits
- (8) Jain 2010 GHA: Free chlorine residuals > 0.2 mg/L in 74-89% of samples Unclear whether testing was during unannounced visits
- (9) Lule 2005 UGA: Compliance not reported
- (10) Opryszko 2010-iii AFG: Self reported use of Chlorine in the previous two weeks was 82%
- (11) Quick 1999 BOL: The proportion of stored water samples with detectable levels of total chlorine increased from 71 % at the time of the first observation to 95% at the final visit
- (12) Reller 2003-iii GUA: Participants had free chlorine >0.1 mg/mL in 44% of samples Testing during unannounced visit
- (13) Semenza 1998 UZB: Chlorine was detected in 73% of household samples at the end of the study.

Analysis 3.6. Comparison 3 POU: water chlorination versus control, Outcome 6 Diarrhoea: cluster-RCTs: subgrouped by sufficiency of water quantity.

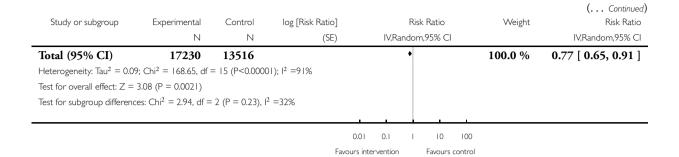
Comparison: 3 POU: water chlorination versus control

Outcome: 6 Diarrhoea: cluster-RCTs: subgrouped by sufficiency of water quantity



Interventions to improve water quality for preventing diarrhoea (Review)

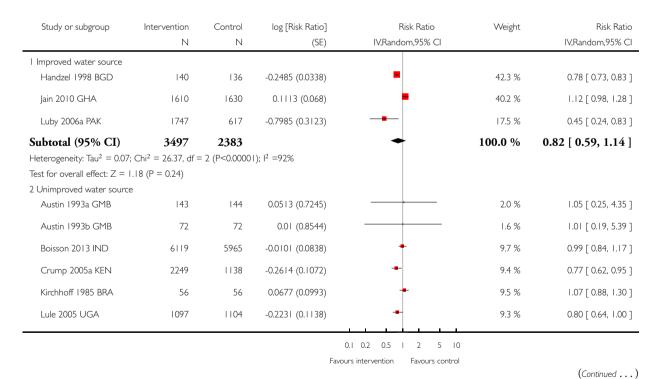
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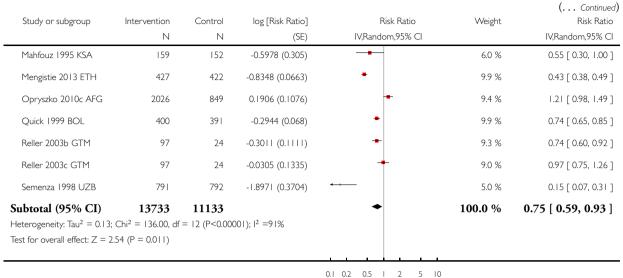


Analysis 3.7. Comparison 3 POU: water chlorination versus control, Outcome 7 Diarrhoea: cluster-RCTs: subgrouped by water source.

Comparison: 3 POU: water chlorination versus control

Outcome: 7 Diarrhoea: cluster-RCTs: subgrouped by water source



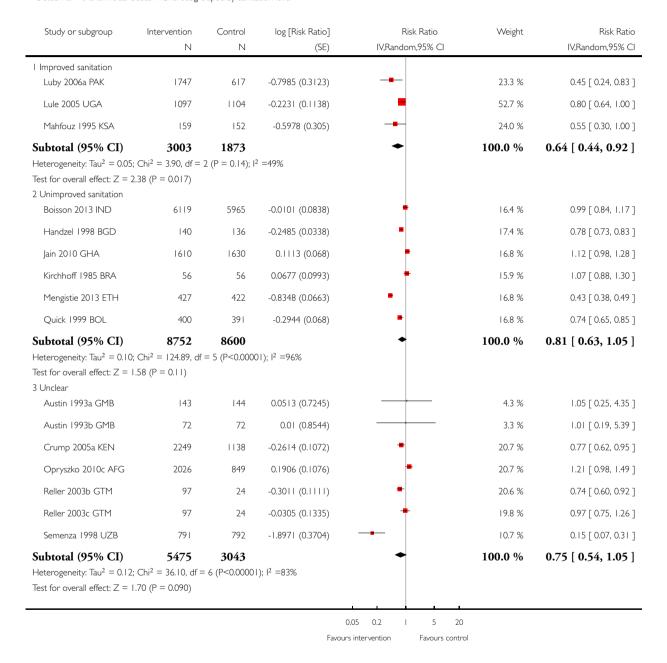


Favours intervention Favours control

Analysis 3.8. Comparison 3 POU: water chlorination versus control, Outcome 8 Diarrhoea: cluster-RCTs: subgrouped by sanitation level.

Comparison: 3 POU: water chlorination versus control

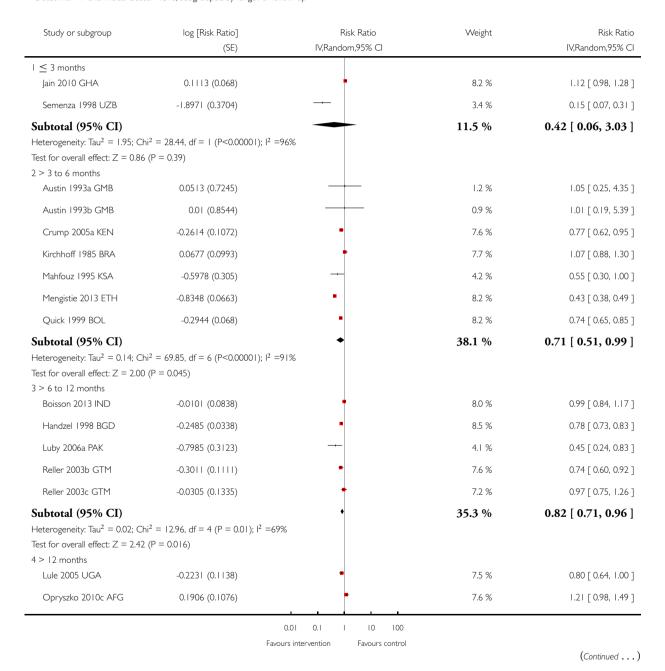
Outcome: 8 Diarrhoea: cluster-RCTs: subgrouped by sanitation level



Analysis 3.9. Comparison 3 POU: water chlorination versus control, Outcome 9 Diarrhoea: cluster-RCTs; subgrouped by length of follow-up.

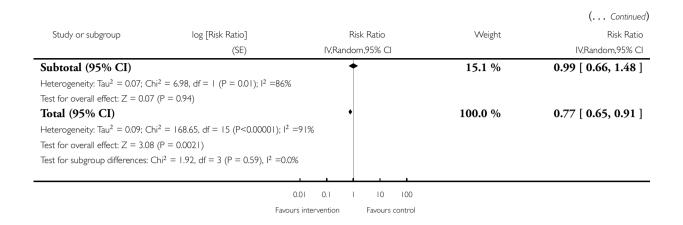
Comparison: 3 POU: water chlorination versus control

Outcome: 9 Diarrhoea: cluster-RCTs; subgrouped by length of follow-up



Interventions to improve water quality for preventing diarrhoea (Review)

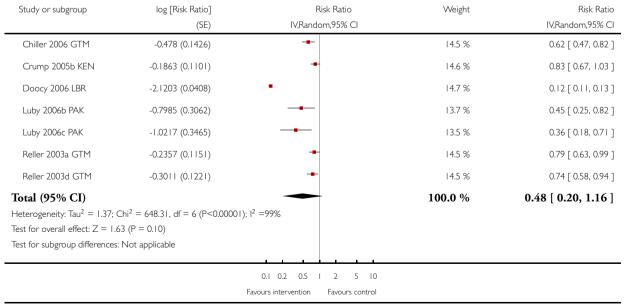
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Analysis 4.1. Comparison 4 POU: flocculation and disinfection versus control, Outcome 1 Diarrhoea: cluster-RCTs.

Comparison: 4 POU: flocculation and disinfection versus control

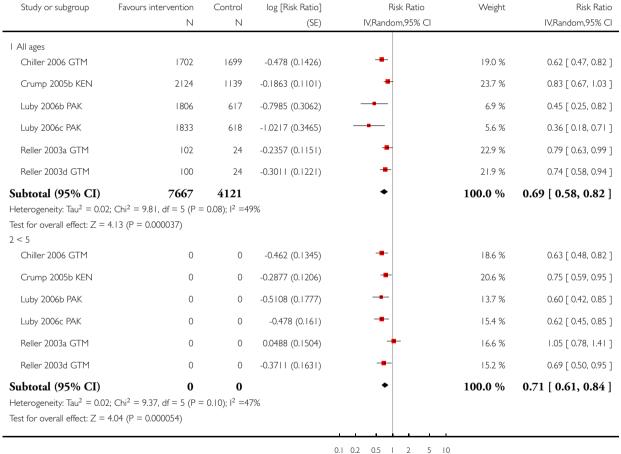
Outcome: I Diarrhoea: cluster-RCTs



Analysis 4.2. Comparison 4 POU: flocculation and disinfection versus control, Outcome 2 Diarrhoea: cluster-RCTs: subgrouped by age; excluding Doocy 2006 LBR.

Comparison: 4 POU: flocculation and disinfection versus control

Outcome: 2 Diarrhoea: cluster-RCTs: subgrouped by age; excluding Doocy 2006 LBR



0.1 0.2 0.3 1 2 3 10

Favours intervention Favours control

Analysis 4.3. Comparison 4 POU: flocculation and disinfection versus control, Outcome 3 Diarrhoea: cluster-RCTs: subgrouped by adherence.

Comparison: 4 POU: flocculation and disinfection versus control

Outcome: 3 Diarrhoea: cluster-RCTs: subgrouped by adherence

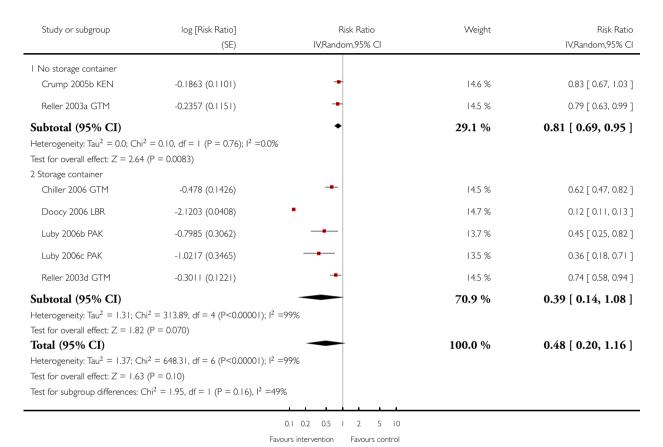
| Study or subgroup | Intervention N | Control N | log [Risk Ratio] (SE) | Risk Ratio IV,Random,95% CI | Weight | Risk Ratio IV,Random,95% CI |
|--|------------------------|-------------------|--------------------------|--------------------------------|---------|--------------------------------|
| 2 Residual chlorine 51 to 85 | 5% | | | | | |
| Doocy 2006 LBR | 1138 | 1053 | -2.1203 (0.0408) | - | 100.0 % | 0.12 [0.11, 0.13] |
| Subtotal (95% CI) | 1138 | 1053 | | • | 100.0 % | 0.12 [0.11, 0.13] |
| Heterogeneity: not applicab | ble | | | | | |
| Test for overall effect: $Z = 5$ | 51.97 (P < 0.00001) | | | | | |
| 3 Residual chlorine < 50% | | | | | | |
| Chiller 2006 GTM | 1702 | 1699 | -0.478 (0.1426) | - | 17.9 % | 0.62 [0.47, 0.82] |
| Crump 2005b KEN | 2124 | 1139 | -0.1863 (0.1101) | = | 30.1 % | 0.83 [0.67, 1.03] |
| Reller 2003a GTM | 102 | 24 | -0.2357 (0.1151) | - | 27.5 % | 0.79 [0.63, 0.99] |
| Reller 2003d GTM | 100 | 24 | -0.3011 (0.1221) | - | 24.5 % | 0.74 [0.58, 0.94] |
| Subtotal (95% CI) | 4028 | 2886 | | • | 100.0 % | 0.76 [0.67, 0.85] |
| Heterogeneity: $Tau^2 = 0.0$; | $Chi^2 = 2.83, df = 3$ | $(P = 0.42); I^2$ | =0.0% | | | |
| Test for overall effect: $Z = \frac{1}{2}$ | 4.64 (P < 0.00001) | | | | | |
| 4 Residual chlorine not mea | asured | | | | | |
| Luby 2006b PAK | 1806 | 617 | -0.7985 (0.3062) | - | 56.2 % | 0.45 [0.25, 0.82] |
| Luby 2006c PAK | 1833 | 618 | -1.0217 (0.3465) | | 43.8 % | 0.36 [0.18, 0.71] |
| Subtotal (95% CI) | 3639 | 1235 | | • | 100.0 % | 0.41 [0.26, 0.64] |
| Heterogeneity: $Tau^2 = 0.0$; | $Chi^2 = 0.23, df = 1$ | $(P = 0.63); I^2$ | =0.0% | | | |
| Test for overall effect: $Z = 3$ | 3.91 (P = 0.000094) | | | | | |
| | | | | | | |
| | | | | 0.1 0.2 0.5 1 2 5 10 |) | |

Favours intervention Favours control

Analysis 4.4. Comparison 4 POU: flocculation and disinfection versus control, Outcome 4 Diarrhoea: cluster-RCTs: subgrouped by additional storage container.

Comparison: 4 POU: flocculation and disinfection versus control

Outcome: 4 Diarrhoea: cluster-RCTs: subgrouped by additional storage container

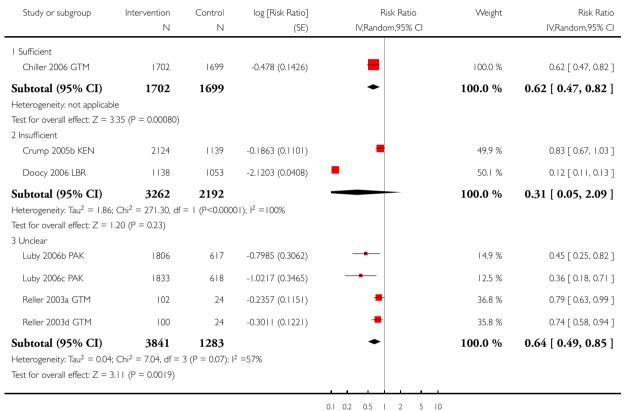


Interventions to improve water quality for preventing diarrhoea (Review)

Analysis 4.5. Comparison 4 POU: flocculation and disinfection versus control, Outcome 5 Diarrhoea: cluster-RCTs: subgrouped by sufficiency of water quantity.

Comparison: 4 POU: flocculation and disinfection versus control

Outcome: 5 Diarrhoea: cluster-RCTs: subgrouped by sufficiency of water quantity



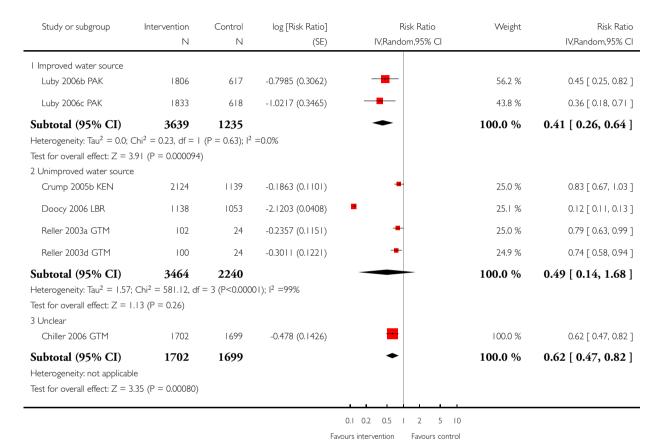
Favours intervention Favours control

Cochrane Collaboration.

Analysis 4.6. Comparison 4 POU: flocculation and disinfection versus control, Outcome 6 Diarrhoea: cluster-RCTs: subgrouped by water source.

Comparison: 4 POU: flocculation and disinfection versus control

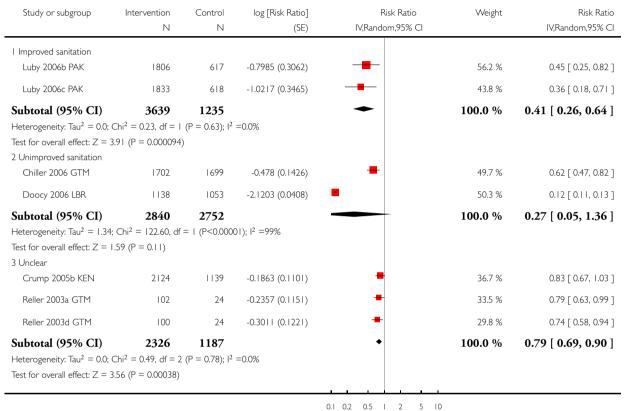
Outcome: 6 Diarrhoea: cluster-RCTs: subgrouped by water source



Analysis 4.7. Comparison 4 POU: flocculation and disinfection versus control, Outcome 7 Diarrhoea: cluster-RCTs: subgrouped by sanitation level.

Comparison: 4 POU: flocculation and disinfection versus control

Outcome: 7 Diarrhoea: cluster-RCTs: subgrouped by sanitation level

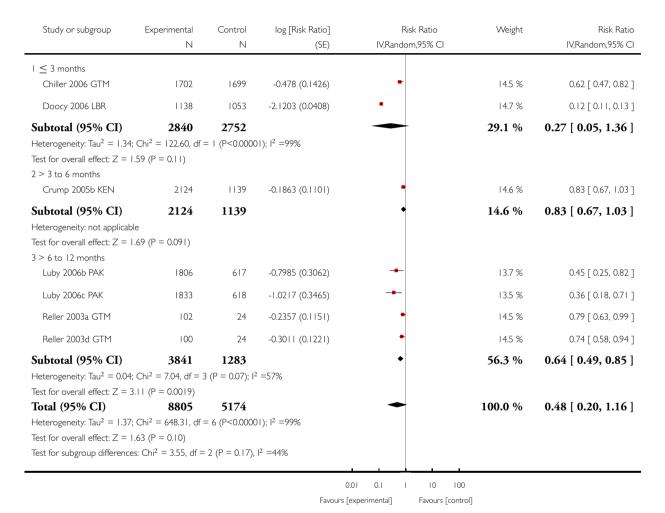


Favours intervention Favours control

Analysis 4.8. Comparison 4 POU: flocculation and disinfection versus control, Outcome 8 Diarrhoea: cluster-RCTs: subgrouped by length of follow-up.

Comparison: 4 POU: flocculation and disinfection versus control

Outcome: 8 Diarrhoea: cluster-RCTs: subgrouped by length of follow-up



Analysis 5.1. Comparison 5 POU: filtration versus control, Outcome 1 Diarrhoea: cluster-RCTs: subgrouped by age.

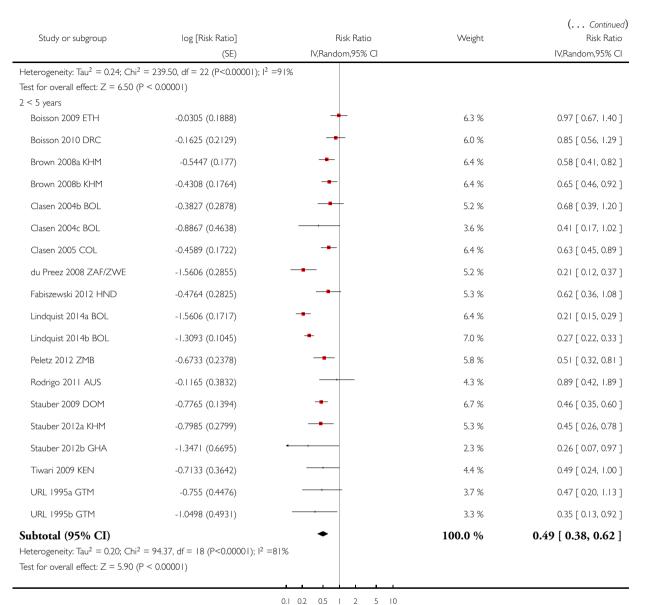
Comparison: 5 POU: filtration versus control

Outcome: I Diarrhoea: cluster-RCTs: subgrouped by age

| Study or subgroup | log [Risk Ratio] (SE) | Risk Ratio IV,Random,95% CI | Weight | Risk Ratio IV,Random,95% CI |
|-----------------------|--------------------------|--------------------------------|---------|--------------------------------|
| I All ages | | | | |
| Abebe 2014 ZAF | -1.5418 (0.0883) | • | 5.2 % | 0.21 [0.18, 0.25] |
| Boisson 2009 ETH | -0.2877 (0.1139) | - | 5.1 % | 0.75 [0.60, 0.94] |
| Boisson 2010 DRC | -0.1625 (0.1777) | + | 4.7 % | 0.85 [0.60, 1.20] |
| Brown 2008a KHM | -0.6733 (0.1114) | - | 5.1 % | 0.51 [0.41, 0.63] |
| Brown 2008b KHM | -0.5447 (0.1073) | - | 5.1 % | 0.58 [0.47, 0.72] |
| Clasen 2004b BOL | -0.6733 (0.3023) | | 3.9 % | 0.51 [0.28, 0.92] |
| Clasen 2004c BOL | -0.5852 (0.1332) | - | 5.0 % | 0.56 [0.43, 0.72] |
| Clasen 2005 COL | -0.803 (0.2132) | | 4.5 % | 0.45 [0.29, 0.68] |
| Colford 2002 USA | -0.6061 (0.1939) | | 4.6 % | 0.55 [0.37, 0.80] |
| Colford 2005 USA | -0.2399 (0.3853) | | 3.3 % | 0.79 [0.37, 1.67] |
| Colford 2009 USA | -0.1393 (0.0826) | • | 5.2 % | 0.87 [0.74, 1.02] |
| du Preez 2008 ZAF/ZWE | -1.5606 (0.2855) | | 4.0 % | 0.21 [0.12, 0.37] |
| Fabiszewski 2012 HND | -0.4748 (0.2905) | | 4.0 % | 0.62 [0.35, 1.10] |
| Lindquist 2014a BOL | -1.5606 (0.1717) | | 4.8 % | 0.21 [0.15, 0.29] |
| Lindquist 2014b BOL | -1.3093 (0.1045) | + | 5.1 % | 0.27 [0.22, 0.33] |
| Peletz 2012 ZMB | -0.7765 (0.2181) | | 4.5 % | 0.46 [0.30, 0.71] |
| Rodrigo 2011 AUS | -0.1625 (0.2039) | + | 4.6 % | 0.85 [0.57, 1.27] |
| Stauber 2009 DOM | -0.755 (0.1221) | | 5.0 % | 0.47 [0.37, 0.60] |
| Stauber 2012a KHM | -0.8916 (0.2732) | | 4.1 % | 0.41 [0.24, 0.70] |
| Stauber 2012b GHA | -0.8916 (0.42) | | 3.1 % | 0.41 [0.18, 0.93] |
| Tiwari 2009 KEN | -0.7765 (0.3763) | | 3.4 % | 0.46 [0.22, 0.96] |
| URL 1995a GTM | -0.755 (0.4476) | | 2.9 % | 0.47 [0.20, 1.13] |
| URL 1995b GTM | -1.0498 (0.4931) | | 2.7 % | 0.35 [0.13, 0.92] |
| Subtotal (95% CI) | | • | 100.0 % | 0.48 [0.38, 0.59] |

0.1 0.2 0.5 I 2 5 10 Favours intervention Favours control

(Continued \dots)



Favours intervention Favours control

Analysis 5.2. Comparison 5 POU: filtration versus control, Outcome 2 Diarrhoea: cluster-RCTs: subgrouped by type of filtration.

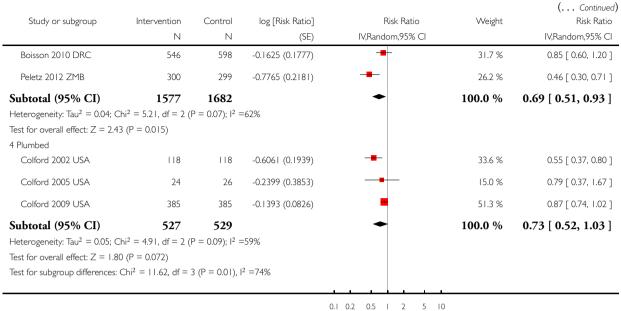
Comparison: 5 POU: filtration versus control

Outcome: 2 Diarrhoea: cluster-RCTs: subgrouped by type of filtration

| Study or subgroup | Intervention N | Control N | log [Risk Ratio] (SE) | Risk Ratio IV,Random,95% CI | Weight | Risk Ratio IV,Random,95% CI |
|---|----------------|--------------------|---|--------------------------------|---------|--------------------------------|
| I Ceramic filter | | | | | | |
| Abebe 2014 ZAF | 39 | 35 | -1.5418 (0.0883) | • | 9.9 % | 0.21 [0.18, 0.25] |
| Brown 2008a KHM | 395 | 203 | -0.6733 (0.1114) | - | 9.7 % | 0.51 [0.41, 0.63] |
| Brown 2008b KHM | 398 | 200 | -0.5447 (0.1073) | - | 9.8 % | 0.58 [0.47, 0.72] |
| Clasen 2004b BOL | 210 | 107 | -0.6733 (0.3023) | | 7.3 % | 0.51 [0.28, 0.92] |
| Clasen 2004c BOL | 140 | 140 | -0.5852 (0.1332) | | 9.5 % | 0.56 [0.43, 0.72] |
| Clasen 2005 COL | 415 | 265 | -0.803 (0.2132) | | 8.5 % | 0.45 [0.29, 0.68] |
| du Preez 2008 ZAF/ZWE | 60 | 55 | -1.5606 (0.2855) | | 7.5 % | 0.21 [0.12, 0.37] |
| Lindquist 2014a BOL | 330 | 140 | -1.5606 (0.1717) | | 9.1 % | 0.21 [0.15, 0.29] |
| Lindquist 2014b BOL | 285 | 139 | -1.3093 (0.1045) | - | 9.8 % | 0.27 [0.22, 0.33] |
| Rodrigo 2011 AUS | 698 | 654 | -0.1625 (0.2039) | - | 8.6 % | 0.85 [0.57, 1.27] |
| URL 1995a GTM | 289 | 134 | -0.755 (0.4476) | | 5.4 % | 0.47 [0.20, 1.13] |
| URL 1995b GTM | 297 | 135 | -1.0498 (0.4931) | | 4.9 % | 0.35 [0.13, 0.92] |
| Subtotal (95% CI) | 3556 | 2207 | | • | 100.0 % | 0.39 [0.29, 0.53] |
| Heterogeneity: Tau ² = 0.22; Chi Test for overall effect: Z = 6.22 2 Sand filtration Fabiszewski 2012 HND | | I (P<0.0000 488 | 1); l ² =91% -0.4748 (0.2905) | - | 11.3 % | 0.62 [0.35, 1.10] |
| Stauber 2009 DOM | 447 | 460 | -0.755 (0.1221) | - | 63.8 % | 0.47 [0.37, 0.60] |
| Stauber 2012a KHM | 546 | 601 | -0.8916 (0.2732) | | 12.8 % | 0.41 [0.24, 0.70] |
| Stauber 2012b GHA | 1012 | 1031 | -0.8916 (0.42) | | 5.4 % | 0.41 [0.18, 0.93] |
| Tiwari 2009 KEN | 206 | 181 | -0.7765 (0.3763) | | 6.7 % | 0.46 [0.22, 0.96] |
| Subtotal (95% CI) | 2743 | 2761 | | • | 100.0 % | 0.47 [0.39, 0.57] |
| Heterogeneity: $Tau^2 = 0.0$; Chi^2 Test for overall effect: $Z = 7.68$ | * | | 0.0% | | | [,, |
| 3 LifeStraw Boisson 2009 ETH | 731 | 785 | -0.2877 (0.1139) | _ | 42.1 % | 0.75 [0.60, 0.94] |

(Continued \dots)

Favours intervention Favours control



0.1 0.2 0.5 1 2 5 10

Favours intervention Favours control

Analysis 5.3. Comparison 5 POU: filtration versus control, Outcome 3 Diarrhoea: cluster-RCTs: subgrouped by blinding of participants.

Comparison: 5 POU: filtration versus control

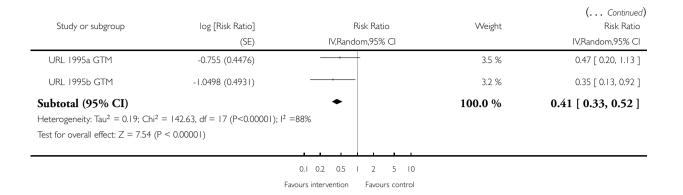
Outcome: 3 Diarrhoea: cluster-RCTs: subgrouped by blinding of participants

| Study or subgroup | log [Risk Ratio] (SE) | Risk Ratio IV,Random,95% CI | Weight | Risk Ratio IV,Random,95% CI |
|--|--------------------------|--------------------------------|---------|--------------------------------|
| I Low risk | | | | |
| Boisson 2010 DRC | -0.1625 (0.1777) | • | 17.7 % | 0.85 [0.60, 1.20] |
| Colford 2002 USA | -0.6061 (0.1939) | | 15.4 % | 0.55 [0.37, 0.80] |
| Colford 2005 USA | -0.2399 (0.3853) | | 4.4 % | 0.79 [0.37, 1.67] |
| Colford 2009 USA | -0.1393 (0.0826) | • | 48.3 % | 0.87 [0.74, 1.02] |
| Rodrigo 2011 AUS | -0.1625 (0.2039) | - | 14.1 % | 0.85 [0.57, 1.27] |
| Subtotal (95% CI) Heterogeneity: $Tau^2 = 0.01$; $Chi^2 = 0$ | | • | 100.0 % | 0.80 [0.68, 0.94] |
| Abebe 2014 ZAF | -1.5418 (0.0883) | + | 6.9 % | 0.21 [0.18, 0.25] |
| Boisson 2009 ETH | -0.2877 (0.1139) | | 6.8 % | 0.75 [0.60, 0.94] |
| Brown 2008a KHM | -0.6733 (0.1114) | - | 6.8 % | 0.51 [0.41, 0.63] |
| Brown 2008b KHM | -0.5447 (0.1073) | - | 6.8 % | 0.58 [0.47, 0.72] |
| Clasen 2004b BOL | -0.6733 (0.3023) | | 4.9 % | 0.51 [0.28, 0.92] |
| Clasen 2004c BOL | -0.5852 (0.1332) | | 6.6 % | 0.56 [0.43, 0.72] |
| Clasen 2005 COL | -0.803 (0.2132) | | 5.8 % | 0.45 [0.29, 0.68] |
| du Preez 2008 ZAF/ZWE | -1.5606 (0.2855) | | 5.1 % | 0.21 [0.12, 0.37] |
| Fabiszewski 2012 HND | -0.4748 (0.2905) | | 5.0 % | 0.62 [0.35, 1.10] |
| Lindquist 2014a BOL | -1.5606 (0.1717) | | 6.3 % | 0.21 [0.15, 0.29] |
| Lindquist 2014b BOL | -1.3093 (0.1045) | - | 6.8 % | 0.27 [0.22, 0.33] |
| Peletz 2012 ZMB | -0.7765 (0.2181) | | 5.8 % | 0.46 [0.30, 0.71] |
| Stauber 2009 DOM | -0.755 (0.1221) | + | 6.7 % | 0.47 [0.37, 0.60] |
| Stauber 2012a KHM | -0.8916 (0.2732) | | 5.2 % | 0.41 [0.24, 0.70] |
| Stauber 2012b GHA | -0.8916 (0.42) | | 3.7 % | 0.41 [0.18, 0.93] |
| Tiwari 2009 KEN | -0.7765 (0.3763) | | 4.1 % | 0.46 [0.22, 0.96] |

Favours intervention Favours control

0.1 0.2 0.5 1 2 5 10

(Continued \dots)



Analysis 5.4. Comparison 5 POU: filtration versus control, Outcome 4 Diarrhoea: ceramic filter studies subgrouped by water source.

Comparison: 5 POU: filtration versus control

Outcome: 4 Diarrhoea: ceramic filter studies subgrouped by water source

| Study or subgroup | Intervention | Control | log [Risk Ratio] | Risk Ratio | Weight | Risk Ratio |
|--|------------------------|--------------|---------------------|-----------------------------------|--------|---------------------|
| | Ν | Ν | (SE) | IV,Random,95% CI | | IV,Random,95% CI |
| I Improved water source | | | | | | _ |
| Abebe 2014 ZAF | 39 | 35 | -1.5418 (0.0883) | • | 9.9 % | 0.21 [0.18, 0.25] |
| Clasen 2004b BOL | 210 | 107 | -0.6733 (0.3023) | - | 7.3 % | 0.51 [0.28, 0.92] |
| du Preez 2008 ZAF/ZWE | 60 | 55 | -1.5606 (0.2855) | + | 7.5 % | 0.21 [0.12, 0.37] |
| Lindquist 2014a BOL | 330 | 140 | -1.5606 (0.1717) | • | 9.1 % | 0.21 [0.15, 0.29] |
| Lindquist 2014b BOL | 285 | 139 | -1.3093 (0.1045) | • | 9.8 % | 0.27 [0.22, 0.33] |
| Rodrigo 2011 AUS | 698 | 654 | -0.1625 (0.2039) | + | 8.6 % | 0.85 [0.57, 1.27] |
| URL 1995a GTM | 289 | 134 | -0.755 (0.4476) | | 5.4 % | 0.47 [0.20, 1.13] |
| URL 1995b GTM | 297 | 135 | -1.0498 (0.4931) | - | 4.9 % | 0.35 [0.13, 0.92] |
| Subtotal (95% CI) | 2208 | 1399 | | • | 62.5 % | 0.33 [0.23, 0.46] |
| Heterogeneity: Tau ² = 0.18; Ch | $i^2 = 47.69$, df = 7 | (P<0.00001); | I ² =85% | | | |
| Test for overall effect: $Z = 6.33$ | (P < 0.00001) | | | | | |
| | | | | | | |
| | | | | 0.001 0.01 0.1 1 10 100 1000 | 0 | |
| | | | Fav | ours intervention Favours control | | |
| | | | | | | (Continued) |

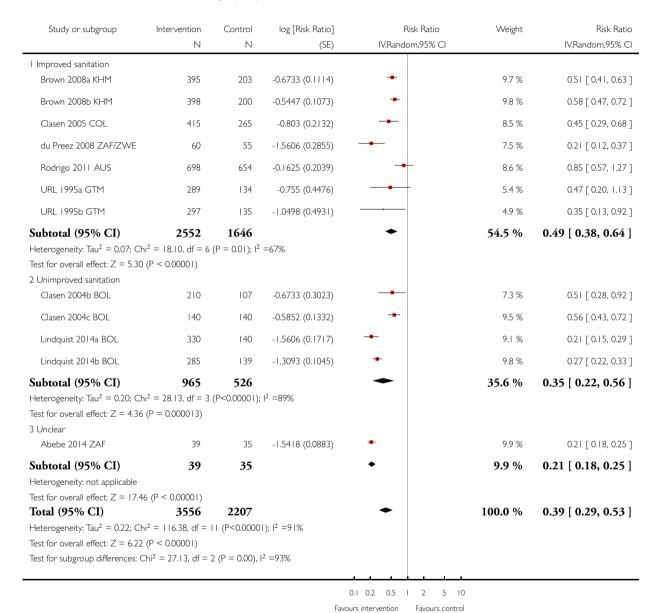
| Study or subgroup | Intervention | Control | log [Risk Ratio] | Risk Ratio | Weight | (Continued) Risk Ratio |
|---|--------------------------|---------------------------|------------------|------------------|---------|----------------------------|
| | N | N | (SE) | IV,Random,95% CI | | IV,Random,95% CI |
| 2 Unimproved water source | | | | | | |
| Brown 2008a KHM | 395 | 203 | -0.6733 (0.1114) | • | 9.7 % | 0.51 [0.41, 0.63] |
| Brown 2008b KHM | 398 | 200 | -0.5447 (0.1073) | • | 9.8 % | 0.58 [0.47, 0.72] |
| Clasen 2004c BOL | 140 | 140 | -0.5852 (0.1332) | - | 9.5 % | 0.56 [0.43, 0.72] |
| Clasen 2005 COL | 415 | 265 | -0.803 (0.2132) | • | 8.5 % | 0.45 [0.29, 0.68] |
| Subtotal (95% CI) | 1348 | 808 | | • | 37.5 % | 0.54 [0.48, 0.61] |
| Heterogeneity: Tau ² = 0.0; Ch | $i^2 = 1.53$, df = 3 (P | = 0.68); I ² = | 0.0% | | | |
| Test for overall effect: $Z = 9.7$ | I (P < 0.00001) | | | | | |
| Total (95% CI) | 3556 | 2207 | | • | 100.0 % | 0.39 [0.29, 0.53] |
| Heterogeneity: $Tau^2 = 0.22$; C | $hi^2 = 116.38, df = 1$ | I (P<0.0000 | 1); 12 =91% | | | |
| Test for overall effect: $Z = 6.2$ | 2 (P < 0.00001) | | | | | |
| Test for subgroup differences: | $Chi^2 = 7.04, df = 1$ | $(P = 0.01), I^2$ | =86% | | | |
| | | | , | | | |

0.001 0.01 0.1 I 10 100 1000 Favours intervention Favours control

Analysis 5.5. Comparison 5 POU: filtration versus control, Outcome 5 Diarrhoea: ceramic filter studies subgrouped by sanitation level.

Comparison: 5 POU: filtration versus control

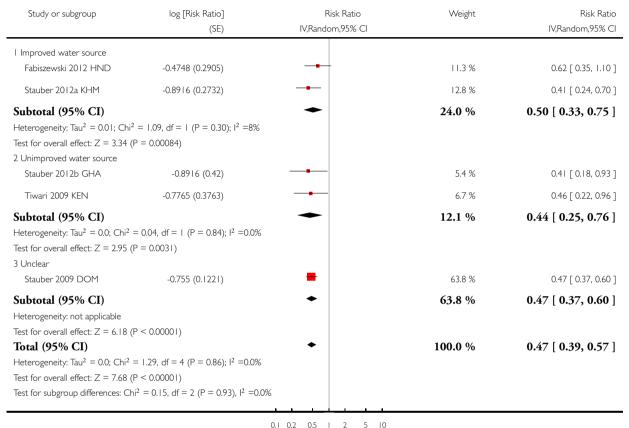
Outcome: 5 Diarrhoea: ceramic filter studies subgrouped by sanitation level



Analysis 5.6. Comparison 5 POU: filtration versus control, Outcome 6 Diarrhoea: sand filter studies: subgrouped by water source.

Comparison: 5 POU: filtration versus control

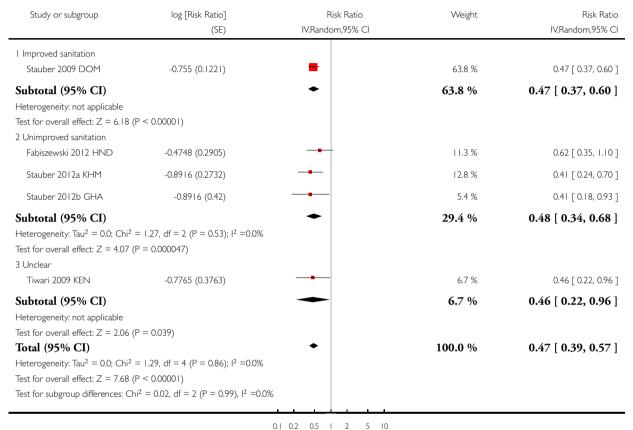
Outcome: 6 Diarrhoea: sand filter studies: subgrouped by water source



Analysis 5.7. Comparison 5 POU: filtration versus control, Outcome 7 Diarrhoea: sand filter studies: subgrouped by sanitation level.

Comparison: 5 POU: filtration versus control

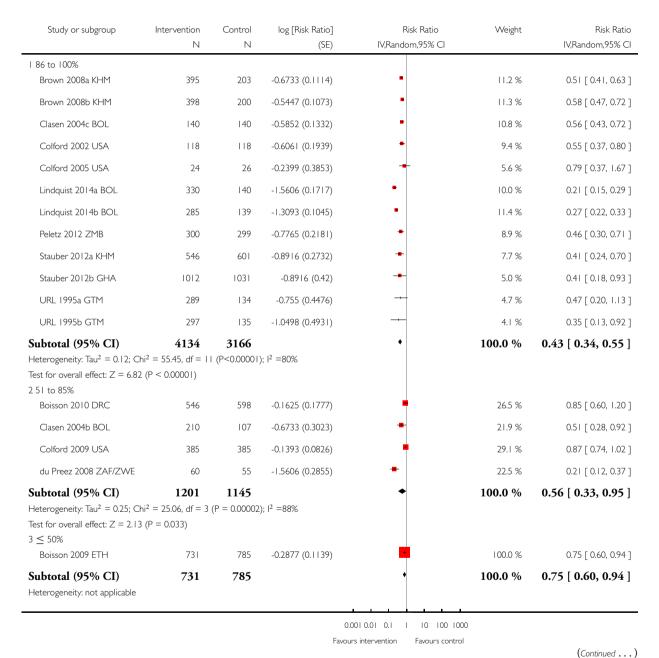
Outcome: 7 Diarrhoea: sand filter studies: subgrouped by sanitation level



Analysis 5.8. Comparison 5 POU: filtration versus control, Outcome 8 Diarrhoea: cluster-RCTs: subgrouped by adherence.

Comparison: 5 POU: filtration versus control

Outcome: 8 Diarrhoea: cluster-RCTs: subgrouped by adherence



(....

| Study or subgroup | Intervention | Control | log [Risk Ratio] | Risk Ratio | Weight | (Continued) Risk Ratio |
|--|----------------------------|--------------|------------------|------------------|---------|----------------------------|
| | N | N | (SE) | IV,Random,95% CI | | IV,Random,95% CI |
| Test for overall effect: $Z = 2.5$ | 3 (P = 0.012) | | | | | |
| 4 Not reported | | | | | | |
| Abebe 2014 ZAF | 39 | 35 | -1.5418 (0.0883) | • | 18.8 % | 0.21 [0.18, 0.25] |
| Clasen 2005 COL | 415 | 265 | -0.803 (0.2132) | - | 16.9 % | 0.45 [0.29, 0.68] |
| Fabiszewski 2012 HND | 532 | 488 | -0.4748 (0.2905) | - | 15.3 % | 0.62 [0.35, 1.10] |
| Rodrigo 2011 AUS | 698 | 654 | -0.1625 (0.2039) | + | 17.1 % | 0.85 [0.57, 1.27] |
| Stauber 2009 DOM | 447 | 460 | -0.755 (0.1221) | - | 18.4 % | 0.47 [0.37, 0.60] |
| Tiwari 2009 KEN | 206 | 181 | -0.7765 (0.3763) | | 13.4 % | 0.46 [0.22, 0.96] |
| Subtotal (95% CI) | 2337 | 2083 | | • | 100.0 % | 0.46 [0.28, 0.75] |
| Heterogeneity: $Tau^2 = 0.32$; C | $2hi^2 = 61.36$, $df = 5$ | (P<0.00001); | $1^2 = 92\%$ | | | |
| Test for overall effect: $Z = 3.0^{\circ}$ | 9 (P = 0.0020) | | | | | |
| | , , , , | | | | | |
| | | | | | | |

0.001 0.01 0.1 I 10 100 1000 Favours intervention Favours control

Analysis 5.9. Comparison 5 POU: filtration versus control, Outcome 9 Diarrhoea: cluster-RCTs: subgrouped by additional water storage intervention.

Comparison: 5 POU: filtration versus control

Outcome: 9 Diarrhoea: cluster-RCTs: subgrouped by additional water storage intervention

| Study or subgroup | log [Risk Ratio] (SE) | Risk Ratio IV,Random,95% Cl | Weight | Risk Ratio IV,Random,95% CI |
|---|--|--------------------------------|---------|--------------------------------|
| I Filtration alone | | | | |
| Boisson 2009 ETH | -0.2877 (0.1139) | • | 29.1 % | 0.75 [0.60, 0.94] |
| Boisson 2010 DRC | -0.1625 (0.1777) | - | 21.0 % | 0.85 [0.60, 1.20] |
| Fabiszewski 2012 HND | -0.4748 (0.2905) | - | 11.7 % | 0.62 [0.35, 1.10] |
| Stauber 2012a KHM | -0.8916 (0.2732) | + | 12.7 % | 0.41 [0.24, 0.70] |
| Stauber 2012b GHA | -0.8916 (0.42) | | 6.6 % | 0.41 [0.18, 0.93] |
| Tiwari 2009 KEN | -0.7765 (0.3763) | | 7.9 % | 0.46 [0.22, 0.96] |
| URL 1995a GTM | -0.755 (0.4476) | - | 5.9 % | 0.47 [0.20, 1.13] |
| URL 1995b GTM | -1.0498 (0.4931) | - | 5.0 % | 0.35 [0.13, 0.92] |
| Subtotal (95% CI) | | • | 100.0 % | 0.60 [0.48, 0.76] |
| Heterogeneity: Tau ² = 0.03; Chi ² = | = 10.79, df = 7 (P = 0.15); $I^2 = 35\%$ | Ś | | |
| Test for overall effect: $Z = 4.27$ (P | = 0.000019) | | | |
| 2 Filtration plus storage | | | | |
| Abebe 2014 ZAF | -1.5418 (0.0883) | • | 10.2 % | 0.21 [0.18, 0.25] |
| Brown 2008a KHM | -0.6733 (0.1114) | • | 9.9 % | 0.51 [0.41, 0.63] |
| Brown 2008b KHM | -0.5447 (0.1073) | • | 10.0 % | 0.58 [0.47, 0.72] |
| Clasen 2004b BOL | -0.6733 (0.3023) | - | 7.0 % | 0.51 [0.28, 0.92] |
| Clasen 2004c BOL | -0.5852 (0.1332) | - | 9.7 % | 0.56 [0.43, 0.72] |
| Clasen 2005 COL | -0.803 (0.2132) | • | 8.5 % | 0.45 [0.29, 0.68] |
| du Preez 2008 ZAF/ZWE | -1.5606 (0.2855) | - | 7.3 % | 0.21 [0.12, 0.37] |
| Lindquist 2014a BOL | -1.5606 (0.1717) | • | 9.1 % | 0.21 [0.15, 0.29] |
| Lindquist 2014b BOL | -1.3093 (0.1045) | • | 10.0 % | 0.27 [0.22, 0.33] |
| Peletz 2012 ZMB | -0.7765 (0.2181) | | 8.4 % | 0.46 [0.30, 0.71] |
| Stauber 2009 DOM | -0.755 (0.1221) | • | 9.8 % | 0.47 [0.37, 0.60] |
| Subtotal (95% CI) Heterogeneity: Tau ² = 0.18; Chi ² = | , , | =90% | 100.0 % | 0.38 [0.29, 0.49] |
| Test for overall effect: $Z = 7.11$ (P | < 0.00001) | | | |

0.001 0.01 0.1 I 10 100 1000 Favours intervention Favours control

Analysis 5.10. Comparison 5 POU: filtration versus control, Outcome 10 Diarrhoea: cluster-RCTs; subgrouped by length of follow-up.

Comparison: 5 POU: filtration versus control

Outcome: 10 Diarrhoea: cluster-RCTs; subgrouped by length of follow-up

| Study or subgroup | log [Risk Ratio] (SE) | Risk Ratio IV,Random,95% CI | Weight | Risk Ratio IV,Random,95% CI |
|---|--------------------------|---|---------------|--------------------------------|
| I ≤ 3 months | | | | |
| Lindquist 2014a BOL | -1.5606 (0.1717) | + | 4.8 % | 0.21 [0.15, 0.29] |
| Lindquist 2014b BOL | -1.3093 (0.1045) | • | 5.1 % | 0.27 [0.22, 0.33] |
| Stauber 2012b GHA | -0.8916 (0.42) | | 3.1 % | 0.41 [0.18, 0.93] |
| Subtotal (95% CI) Heterogeneity: $Tau^2 = 0.01$; $Chi^2 = Test$ for overall effect: $Z = 11.18$ (Fig. 2 > 3 to 6 months | , | =30% | 13.0 % | 0.26 [0.20, 0.33] |
| Boisson 2009 ETH | -0.2877 (0.1139) | - | 5.1 % | 0.75 [0.60, 0.94] |
| Brown 2008a KHM | -0.6733 (0.1114) | • | 5.1 % | 0.51 [0.41, 0.63] |
| Brown 2008b KHM | -0.5447 (0.1073) | - | 5.1 % | 0.58 [0.47, 0.72] |
| Clasen 2004c BOL | -0.5852 (0.1332) | + | 5.0 % | 0.56 [0.43, 0.72] |
| Clasen 2005 COL | -0.803 (0.2132) | - | 4.5 % | 0.45 [0.29, 0.68] |
| Colford 2002 USA | -0.6061 (0.1939) | + | 4.6 % | 0.55 [0.37, 0.80] |
| du Preez 2008 ZAF/ZWE | -1.5606 (0.2855) | - | 4.0 % | 0.21 [0.12, 0.37] |
| Fabiszewski 2012 HND | -0.4748 (0.2905) | -+ | 4.0 % | 0.62 [0.35, 1.10] |
| Stauber 2009 DOM | -0.755 (0.1221) | * | 5.0 % | 0.47 [0.37, 0.60] |
| Stauber 2012a KHM | -0.8916 (0.2732) | | 4.1 % | 0.41 [0.24, 0.70] |
| Tiwari 2009 KEN | -0.7765 (0.3763) | | 3.4 % | 0.46 [0.22, 0.96] |
| Subtotal (95% CI) Heterogeneity: Tau ² = 0.03; Chi ² = Test for overall effect: Z = 8.46 (P 3 > 6 to 12 months Abebe 2014 ZAF | , , | • | 50.0 % | 0.52 [0.44, 0.60] |
| Boisson 2010 DRC | -0.1625 (0.1777) | | 4.7 % | 0.85 [0.60, 1.20] |
| Clasen 2004b BOL | -0.1623 (0.1777) | | 3.9 % | 0.51 [0.28, 0.92] |
| | , , | | | - |
| Colford 2005 USA | -0.2399 (0.3853) | 0.01 0.1 I 10 100 Favours Intervention Favours Control | 3.3 % | 0.79 [0.37, 1.67] |

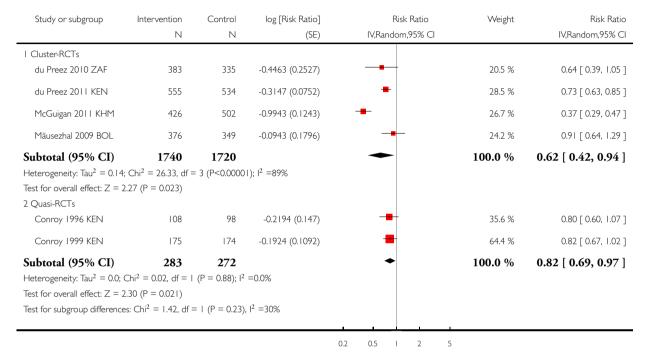
(Continued \dots)

| Study or subgroup | log [Risk Ratio] (SE) | Risk Ratio IV,Random,95% CI | Weight | (Continued) Risk Ratio IV,Random,95% CI |
|---|--------------------------|--|---------------|--|
| Peletz 2012 ZMB | -0.7765 (0.2181) | + | 4.5 % | 0.46 [0.30, 0.71] |
| Rodrigo 2011 AUS | -0.1625 (0.2039) | + | 4.6 % | 0.85 [0.57, 1.27] |
| URL 1995a GTM | -0.755 (0.4476) | | 2.9 % | 0.47 [0.20, 1.13] |
| URL 1995b GTM | -1.0498 (0.4931) | | 2.7 % | 0.35 [0.13, 0.92] |
| Subtotal (95% CI) Heterogeneity: Tau ² = 0.50; Chi ² = Test for overall effect: Z = 2.46 (P 4 > 12 months Colford 2009 USA | . , | 2% | 31.8 % | 0.51 [0.30 , 0.87] |
| Subtotal (95% CI) Heterogeneity: not applicable Test for overall effect: Z = 1.69 (P | , , , | • | 5.2 % | 0.87 [0.74, 1.02] |
| Total (95% CI) Heterogeneity: $Tau^2 = 0.24$; $Chi^2 = 0.24$; $Chi^2 = 0.50$ (P Test for overall effect: $Z = 6.50$ (P Test for subgroup differences: Chi^2 | < 0.00001) | | 100.0 % | 0.48 [0.38, 0.59] |
| | Fave | 0.01 0.1 1 10 100 ours Intervention Favours Control | | |

Analysis 6.1. Comparison 6 POU: solar disinfection versus control, Outcome 1 Diarrhoea: subgrouped by study design.

Comparison: 6 POU: solar disinfection versus control

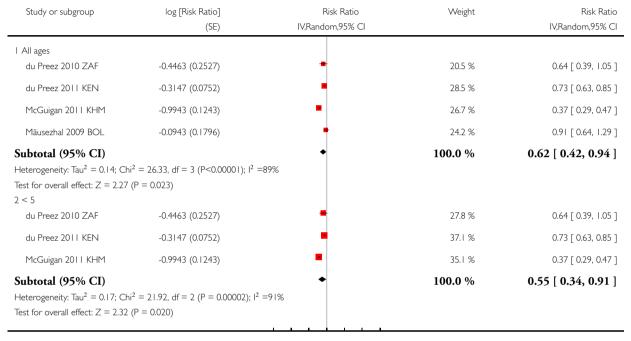
Outcome: 1 Diarrhoea: subgrouped by study design



Analysis 6.2. Comparison 6 POU: solar disinfection versus control, Outcome 2 Diarrhoea: cluster-RCTs; subgrouped by age.

Comparison: 6 POU: solar disinfection versus control

Outcome: 2 Diarrhoea: cluster-RCTs; subgrouped by age



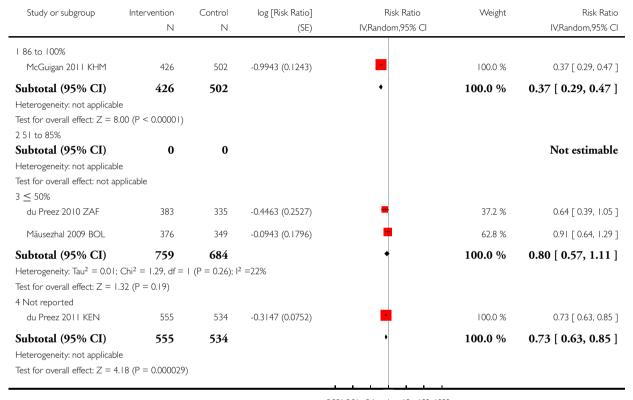
0.001 0.01 0.1 1 10 100 1000

Favours intervention Favours control

Analysis 6.3. Comparison 6 POU: solar disinfection versus control, Outcome 3 Diarrhoea: cluster-RCTs; subgrouped by adherence.

Comparison: 6 POU: solar disinfection versus control

Outcome: 3 Diarrhoea: cluster-RCTs; subgrouped by adherence

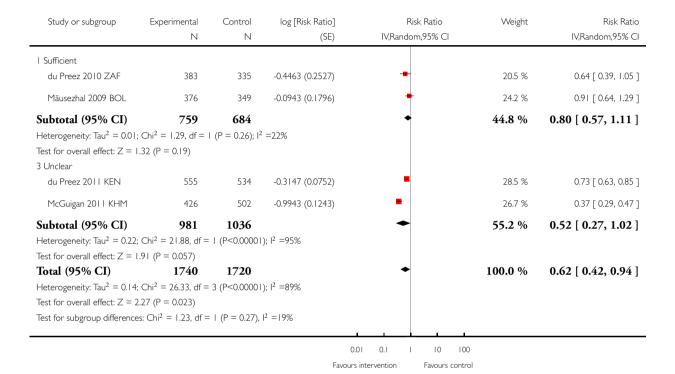


0.001 0.01 0.1 1 10 100 1000 Favours intervention Favours control

Analysis 6.4. Comparison 6 POU: solar disinfection versus control, Outcome 4 Diarrhoea: cluster-RCTs; subgrouped by sufficiency of water supply level.

Comparison: 6 POU: solar disinfection versus control

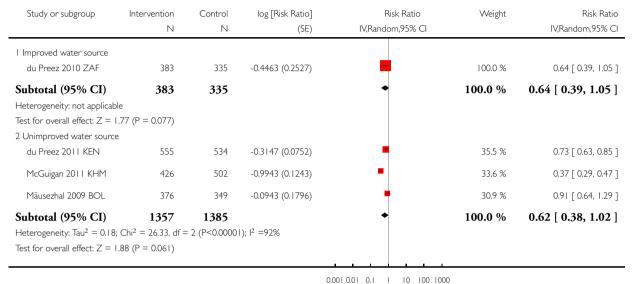
Outcome: 4 Diarrhoea: cluster-RCTs; subgrouped by sufficiency of water supply level



Analysis 6.5. Comparison 6 POU: solar disinfection versus control, Outcome 5 Diarrhoea: cluster-RCTs; subgrouped by water source.

Comparison: 6 POU: solar disinfection versus control

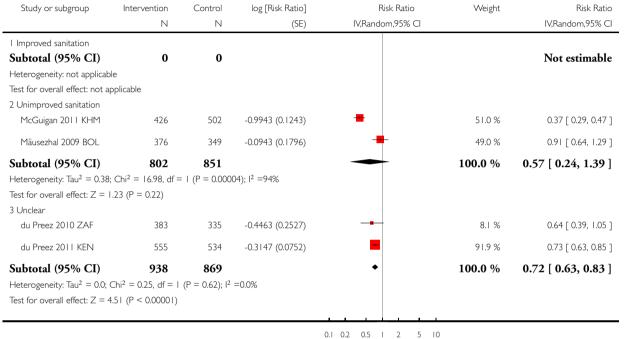
Outcome: 5 Diarrhoea: cluster-RCTs; subgrouped by water source



Analysis 6.6. Comparison 6 POU: solar disinfection versus control, Outcome 6 Diarrhoea: cluster-RCTs; subgrouped by sanitation level.

Comparison: 6 POU: solar disinfection versus control

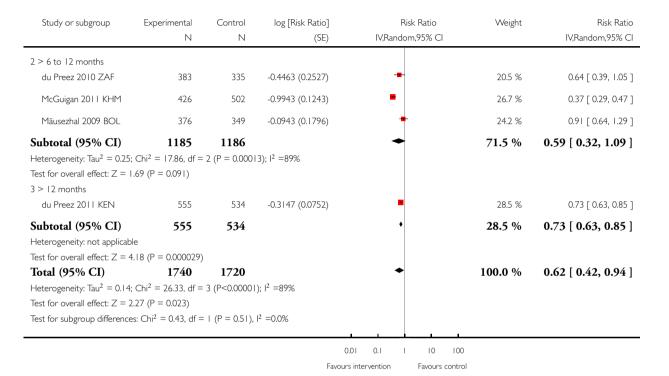
Outcome: 6 Diarrhoea: cluster-RCTs; subgrouped by sanitation level



Analysis 6.7. Comparison 6 POU: solar disinfection versus control, Outcome 7 Diarrhoea: cluster-RCTs; subgrouped by length of follow-up.

Comparison: 6 POU: solar disinfection versus control

Outcome: 7 Diarrhoea: cluster-RCTs; subgrouped by length of follow-up

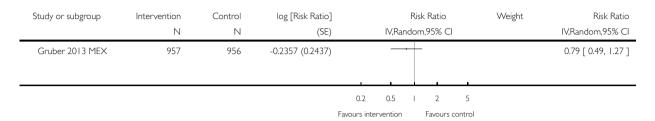


Analysis 7.1. Comparison 7 POU: UV disinfection versus control, Outcome I Diarrhoea: cluster-RCT.

Review: Interventions to improve water quality for preventing diarrhoea

Comparison: 7 POU: UV disinfection versus control

Outcome: I Diarrhoea: cluster-RCT

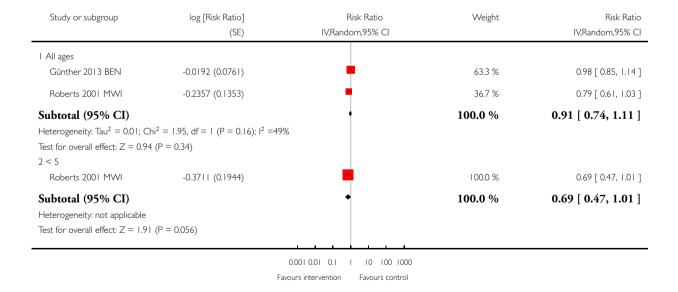


Analysis 8.1. Comparison 8 POU: improved storage versus control, Outcome I Diarrhoea: cluster-RCTs: subgrouped by age.

Review: Interventions to improve water quality for preventing diarrhoea

Comparison: 8 POU: improved storage versus control

Outcome: | Diarrhoea: cluster-RCTs: subgrouped by age



ADDITIONAL TABLES

Table 1. Water quality indicators post-intervention

| Trial | Water quality indicator | Water quality post-intervention: Intervention group | Water quality post intervention: Control group | |
|--|---|---|--|--|
| Abebe 2014 ZAF | CFUs/100 mL | 0 | 80% of control HHs had 10 to 10000 | |
| Austin 1993a GMB | Geometric mean CFUs/100 mL | 178 | 3020 | |
| Austin 1993b GMB | Geometric mean CFUs/100 mL | 42 | 3020 | |
| Boisson 2009 ETH | Arithmetic mean TTC/100 mL (95% CI) | 0 | 725.7 (621.0 to 830.4) | |
| Boisson 2010 DRC | n 2010 DRC Geometric mean TTC/100 mL (95% CI) | | 173.7 (136.6 to 220.9) | |
| Boisson 2013 IND | Geometric mean TTC/100 mL (95% CI) | 50 (44 to 57) | 122 (107 to 139) | |
| Brown 2008a KHM | Geometric mean <i>E. coli</i> /100 mL | 17 | 600 | |
| Brown 2008b KHM | Geometric mean E. coli /100 mL | 15 | 600 | |
| Clasen 2004b BOL | Mean TTC/100 mL | 0.13 | 108 | |
| Clasen 2004c BOL | Arithmetic mean TTC/100 mL | 100% of intervention households: 0 | 16% of control households: 0 66% > 10, 34% > 100, and 11% > 1000 | |
| Clasen 2005 COL | Arithmetic mean TTC/100 mL (95% CI) | 37.3 (6.3 to 48.3) | 150.6 (34.8 to 166.4) | |
| Colford 2002 USA; Colford 2005 USA; Colford 2009 USA | All water met FDA requirements | Not measured because of high water quality | Not measured because of high water quality | |
| Crump 2005a KEN | Samples met WHO guidelines for water quality | 82% | 14% | |
| Crump 2005b KEN | Samples met WHO guidelines for water quality | 78% | 14% | |

Table 1. Water quality indicators post-intervention (Continued)

| du Preez 2008 ZAF/ZWE | Samples met WHO guidelines for water quality | 57% | 30% | | |
|-----------------------|--|--|---|--|--|
| du Preez 2010 ZAF | E. coli in concentrations/100 mL | 62% | "No significant difference be- tween intervention and control groups" | | |
| du Preez 2011 KEN | E. coli ln concentrations/100 mL | Storage containers: 0.723 SODIS bottles: -0.727 | Not reported | | |
| Fabiszewski 2012 HND | Geometric mean <i>E. coli</i> counts per 100 mL (95% CI) | 23.4 (20.2 to 27.0) | 45.4 (38.6 to 53.4) | | |
| Gasana 2002 RWA | Total coliforms/100 mL | Range: 3 to 43 | Range: 4 to 1100 | | |
| Gruber 2013 MEX | Samples with detectable E. coli | 43% | 59% | | |
| Günther 2013 BEN | E. coli contamination > 1000 CFU/100 mL | Not reported specifically; findings imply a 70% reduction in <i>coli</i> incidence for intervention households | | | |
| Handzel 1998 BGD | Stored water samples with <i>E. coli</i> 100 MPN/100 mL | 3% | 16% | | |
| Jain 2010 GHA | Samples with E. coli | 8% | 54% | | |
| Jensen 2003 PAK | Geometric mean E. coli /100 mL | 3 | 49 | | |
| Kirchhoff 1985 BRA | Mean number of faecal coliforms/dL in the samples | 70 | 16000 | | |
| Kremer 2011 KEN | Average reduction in log <i>E. coli</i> | -1.07, corresponding to a 66% | reduction | | |
| Lule 2005 UGA | Median <i>E. coli</i> CFU/100 mL | 23 | 59 | | |
| McGuigan 2011 KHM | Geometric mean CFU/100 mL | 6.8 | 48 | | |
| Mengistie 2013 ETH | Mean E. coli | 0 | 60 | | |
| Peletz 2012 ZMB | Geometric mean TTC/100 mL | Stored water: 3 | Stored water: 181 | | |
| Quick 1999 BOL | Median E. coli /100 mL | 0 | 6400 | | |
| Quick 2002 ZMB | Median E. coli /100 mL | 0 | 3 | | |

Table 1. Water quality indicators post-intervention (Continued)

| Reller 2003a GTM | Samples with < 1 <i>E. coli</i> /100 mL (flocculant/disinfectant) | 40% | 7% |
|-------------------|---|---|--------------------|
| Reller 2003b GTM | Samples with < 1 <i>E. coli</i> /100 mL (flocculant/disinfectant+ vessel) | 57% | 7% |
| Reller 2003c GTM | Samples with < 1 <i>E. coli</i> /100 mL (bleach) | 51% | 7% |
| Reller 2003d GTM | Samples with < 1 <i>E. coli</i> /100 mL (bleach + vessel) | 61% | 7% |
| Semenza 1998 UZB | Faecal colonies/100 mL | 47 | 52 |
| Stauber 2009 DOM | E. coli MPN/100 mL | 11 | 19 |
| Stauber 2012a KHM | E. coli CFU/100 mL | 2.9 | 19.7 |
| Stauber 2012b GHA | Geometric mean <i>E. coli</i> MPN/ 100 mL (95% CI) | Direct filtrate 16 (13 to 20) Stored filtrate: 76 (62 to 91) | 490 (426 to 549) |
| Tiwari 2009 KEN | Geometric mean faecal coliforms/100 mL (95% CI) | 30.0 (21.3 to 42.1) | 88.9 (58.7 to 135) |
| URL 1995a GTM | Samples with fecal coliforms | 91% had 0 fecal coliforms | Not reported |
| URL 1995b GTM | Samples with fecal coliforms | 91% had 0 fecal coliforms | Not reported |

Abbreviations: E. coli: Escherichia coli; FC: faecal coliform.

Table 2. Studies reporting deaths

| Study ID | Intervention | | Control | | P value | Comment |
|---------------------|--------------|--------------|---------|--------------|---------|---------|
| | Deaths | Participants | Deaths | Participants | | |
| Boisson 2010 DRC | 12 | 546 | 8 | 598 | 0.27 | - |
| Colford 2009 USA | 7 | 385 | 6 | 385 | > 0.05 | - |
| Crump 2005a KEN | 17 | 2249 | 28 | 2277 | 0.108 | - |

Table 2. Studies reporting deaths (Continued)

| Crump 2005b KEN | 14 | 2124 | 28 | 2277 | 0.052 | - |
|----------------------|----|------|----|------|--------|--|
| du Preez 2011 KEN | 3 | 555 | 3 | 534 | > 0.05 | - |
| Peletz 2012 ZMB | 3 | 300 | 6 | 299 | 0.28 | - |
| Boisson 2013 IND | ? | 6119 | ? | 5965 | - | Only reports total deaths (46) |
| du Preez 2010 ZAF | ? | 383 | ? | 335 | - | Only reports total deaths (7) |
| Kremer 2011 KEN | ? | - | ? | - | - | Reports recording deaths but does not state how many |
| Boisson 2009 ETH | ? | 731 | ? | 785 | - | Reports recording deaths but does not state how many |

Table 3. Summary of findings: improved water source

Improved water source compared with no intervention for preventing diarrhoea in rural settings in low- and middle-income countries

Patient or population: adults and children

Settings: low- and middle-income countries in rural areas

Intervention: water source improvement

Comparison: no intervention

| Outcomes | Illustrative compara | ntive risks* (95% CI) | Relative effect (95% CI) | pants | Quality of the evidence (GRADE) | |
|---------------------------------------|--|--|-----------------------------|---------------------|-----------------------------------|--|
| | Assumed risk | Corresponding risk | | (studies) | | |
| | No intervention Water source improvement | | | | | |
| Diarrhoea episodes Cluster-RCTs | 3 episodes per per- son per year | 3.7 episodes per person per year (2. 9 to 4.7) | | 3266 (1 trial) | ⊕○○○ very low ^{1,2,3} | |
| Diarrhoea episodes CBA studies | - | - | - | 5895 (5 studies) | ⊕○○○ very low ^{1,4,5} | |

The basis for the **assumed risk** (for example, 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).

CI: confidence interval; RR: risk ratio.

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 assumed risk is based on 2.9 episodes/child year in 2010 (Fischer Walker 2012).

Table 4. Improved water source: description of the interventions

| Study ID | Study design | Setting | In- cidence of di- | Intervention areas | | Control areas | |
|-----------------------|--------------|----------------|--|--|---|-------------------------------------|-----------------------------|
| | | | arrhoea in the control group | Water source intervention | Health promotion activities | Water source | Health promotion activities |
| Opryszko 2010b AFG | Cluster-RCT | Rural villages | 3.1 episodes per person per year | One well per 25 households provid- ing 25 litres/ person/day | None | 35% used unprotected hand dug wells | None |
| Alam 1989 BGD | СВА | Rural villages | 4. 1 episodes per child per year | pump per 4-6 households (3 times as | Female health visitors visited peoples homes and or- ganised group discussion and demonstra- tions to pro- | low, hand-dug wells; some | None described |

¹Downgraded by 1 for serious risk of bias: as diarrhoea episodes were reported by participants this outcome is susceptible to bias from lack of blinding. None of these studies blinded participants and outcome assessors to the treatment allocation.

²No serious inconsistency.

³Downgraded by 2 for serious indirectness: this single RCT from Afghanistan evaluated the provision of protected wells. It is not possible to make broad generalizations to other settings.

⁴Downgraded by 1 for serious inconsistency: statistical heterogeneity was very high (I² statistic = 98%), such that the data could not be pooled. Some large and statistically significant effects were seen in some individual trials, but not others.

⁵Downgraded by 1 for serious indirectness: these studies are from a variety of low- and middle-income countries (Bangladesh, Rwanda, Pakistan, South Africa, China). However, as only single trials evaluated each intervention it is not possible to make broad generalizations.

Table 4. Improved water source: description of the interventions (Continued)

| | | | | | mote hygienic prac- tices for hand pump use, wa- ter storage, child faeces disposal, hand washing | | |
|--------------------|-----|----------------|--|---|--|---|-------------------|
| Gasana 2002 RWA | СВА | Rural villages | 3 episodes per child per year | Site A: Sedimentation tank/ Katadyn filter with communal tap Site B: Gravelsand-charcoal filter on existing water spring Site C: Protective fence around an existing water spring | None described | An existing water spring | None described |
| Jensen 2003 PAK | СВА | Rural villages | 2.8 episodes per person per year | Chlorination of public water supply | None described | Unchlo- rinated poorly functioning sand filter sys- tem | None described |
| Majuru 2011 ZAF | СВА | Rural villages | 0.6 episodes per person per year | Provision of intermittently operated small community water systems distributing potable water to multiple taps throughout the community | None described | Untreated water from a river and its tributaries | |
| Xiao 1997 CHN | СВА | Rural villages | Not reported | | Hygiene edu- cation | Not reported | None described |

Table 4. Improved water source: description of the interventions (Continued)

| tural improve- ments to wells |
|----------------------------------|
|----------------------------------|

Table 5. Improved water source: primary drinking water supply and sanitation facilities

| Trial | Description | Source ¹ | Access to source | Quantity available ³ | Ambient water quality | Sanitation ⁴ |
|--------------------|---|-------------------------|------------------|---------------------------------|--|-------------------------|
| Alam 1989 BGD | Shallow, hand- dug wells; some hand pumps | Unimproved | Unclear | Unclear | Not tested | Unclear |
| Gasana 2002 RWA | Spring | Unimproved | Unclear | Unclear | Baseline range 4 to 1100 total co- liforms/100 mL | Unimproved |
| Jensen 2003 PAK | Some slow sand filters in poor condition; some house- hold taps; major- ity used ground water | Improved | Unclear | Unclear | Baseline geometric mean in intervention village: 13. 3 <i>E. coli</i> CFU/100 mL; control villages: 137/100 mL | Unclear |
| Majuru 2011 ZAF | Surface water, boreholes, water tankers | Improved and unimproved | Unclear | Unclear | Not tested | Unclear |
| Opryszko 2010 | 35% use unprotected dug wells | Unimproved | Sufficient | Sufficient | Not tested | Unclear |
| Xiao 1997 CHN | Well water | Unimproved | Unclear | Unclear | Not tested | Unclear |

¹ Improved' includes household connection, public standpipe, borehole, protected dug well, protected spring, rainwater collection; 'unimproved' includes unprotected well, unprotected spring, vendor-provided water, bottled water; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

²'Sufficient' means located within 500 m, queuing no more than 15 minutes, no more than three minutes to fill 20 L container, and maintained so available consistently; 'insufficient' means that it does not meet any of above; and 'unclear' means unclear or not reported; definition based minimum standards established by The Sphere Project 2011.

³'Sufficient' means a minimum of 15 L/day/person; 'insufficient' means less than 15 L/day/person; and 'unclear' means unclear or not reported; definition based on minimum standards established by The Sphere Project 2011.

⁴'Improved' means connection to a public sewer or septic system, pour flush latrine, simple pit latrine, or ventilated improved pit latrine; 'unimproved' means service or bucket latrine, public latrines, open latrines; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

Table 6. POU chlorination: description of the intervention

| Trial | Study design | Chlorination product? | Distributed free? | Frequency of distribution? | Storage container also distributed? | Compliance | Additional hygiene pro- motion |
|-----------------------|--------------|---|-------------------|----------------------------|-------------------------------------|---|--|
| Austin 1993a GMB | Cluster-RCT | Sodium hypochlorite solution | Yes | Fortnightly | No | 40% compliance measured by residual chlorine | None |
| Austin 1993b GMB | Cluster-RCT | Sodium hypochlorite solution | Yes | Fortnightly | No | 59% compliance measured by residual chlorine | None |
| Boisson 2013 IND | Cluster-RCT | Sodim dichloro- isocyanurate tablets | Yes | Bimonthly | No | 32% compliance measured by residual chlorine | None |
| Crump 2005a KEN | Cluster-RCT | 1% sodium hypochlorite | Yes | Weekly | No | 61% compliance during unannounced weekly visits measured by residual chlorine | Use of ORS, treatment seeking for di- arrhoea |
| Handzel 1998 BGD | Cluster-RCT | 0.25% to 0.3% chlorine solution | Yes | Weekly | Yes | 90% compliance based on residual chlorine measurements | and sanitation |
| Jain 2010 GHA | Cluster-RCT | Sodim dichloro- isocyanurate tablets | Yes | Twice weekly | Yes | 74% to 89% com- pliance mea- sured by chlo- rine residual | ORS provided to those with diarrhoea |
| Kirchhoff 1985 BRA | Cluster-RCT | 10% sodium hypochlorite | Yes | Daily | No | Not reported | Chlorination preformed by study staff |
| Luby 2006a PAK | Cluster-RCT | Sodium hypochlorite solution | Yes | Unclear | Yes | Yes, though rate unclear | Encouraged to only drink treated water |
| Lule 2005 UGA | Cluster-RCT | 0.5% sodium hypochlorite | Yes | Weekly | Yes | Not reported | hygiene edu- cation |

Table 6. POU chlorination: description of the intervention (Continued)

| | | | | | | · | |
|-----------------------|-------------|---|-----|---------|-----|--|--|
| Mahfouz 1995 KSA | Cluster-RCT | Packets of 50 g calcium hypochloride 70%. | Yes | Unclear | No | Some residual chlorine in all intervention samples | None |
| Mengistie 2013 ETH | Cluster-RCT | 1. 25% sodium hypochlorite solution | Yes | Weekly | No | 80% compliance measured by chlorine residual | None |
| Opryszko 2010c AFG | Cluster-RCT | 0. 05% sodium hypochlorite solution | Yes | Monthly | Yes | 78% compliance measured by previous 2 weeks self-report use of chlorine | None |
| Quick 1999 BOL | Cluster-RCT | MIOX unit electrolytically produced disinfectant with 3% brine solution, hypochlorite, chlorine dioxide, ozone, peroxide and other oxidants | Yes | Weekly | Yes | by | munity health volunteers re- inforced mes- sages |
| Reller 2003b GTM | Cluster-RCT | Sodium hypochlorite solution (50, 000 ppm) | Yes | Monthly | No | 36% compliance measure by residual chlorine > 0.1 mg/L on unannounced visits | Motiva- tional and ed- ucational mes- sages about chlori- nation, use of ORS, care seeking for di- arrhoea |
| Reller 2003c GTM | Cluster-RCT | Sodium hypochlorite solution (50, 000 ppm) | Yes | Monthly | Yes | 44% compliance measure by residual chlorine > 0.1 mg/L on unannounced visits | Motiva- tional and ed- ucational mes- sages about chlori- nation, use of ORS, care seeking for di- arrhoea |

 Table 6. POU chlorination: description of the intervention (Continued)

| Semenza 1998 UZB | Cluster-RCT | 1.5% chlorine solution | Yes | Unclear but households were visited twice weekly | Yes | 73% based on residual chlo- rine levels at time of visit | Only drink chlo- rinated water and wash all fruit and veg- eta- bles with chlo- rinated water |
|---------------------|-------------|------------------------------------|-----|---|------------------------|---|---|
| Luby 2004a PAK | CBA | Bleach (sodium hypochlorite) | Yes | Study workers visited weekly and re- supplied the house- holds with di- lute bleach | Yes | Not reported | Encouraged regular treat- ment of drink- ing water |
| Luby 2004b PAK | СВА | Bleach (sodium hypochlorite) | Yes | Study workers visited weekly and re- supplied the house- holds with di- lute bleach | Yes | Not reported | Encouraged regular treat- ment of drink- ing water |
| Quick 2002 ZMB | СВА | 0.5% sodium hypochlorite | Yes | Unclear but house- holds were vis- ited once every two weeks | HHs paid for vessel | 72% compliance measured by water in vessel with chlorine residual | munity volun- |

Table 7. POU chlorination: primary drinking water supply and sanitation facilities

| Trial | Description | Source ¹ | Access to source | Quantity available ³ | Ambient water quality | Sanitation ⁴ |
|-------------|-------------|---------------------|------------------|---------------------------------|---|-------------------------|
| Austin 1993 | Open wells | Unimproved | Sufficient | Unclear | Mean 1871 FC/ 100 mL in wells; among stored water samples: | Unclear |

Table 7. POU chlorination: primary drinking water supply and sanitation facilities (Continued)

| | | | | | mean 3358 FC/ 100 mL in rainy season, 1014 FC/100 mL in dry season | |
|-----------------------|--|-------------------------|------------|--------------|--|---------------------------------------|
| Boisson 2013 IND | 62% unprotected dug well, 17% tube- well, 14% tap, 5% surface water | Unimproved | Unlcear | Unclear | Baseline not reported. Control households: Geometric mean 122 TTC/100 mL | Unimproved |
| Crump 2005 | 50% ponds, 49% rivers | Unimproved | Unclear | Insufficient | Baseline mean 98 <i>E. coli </i> 100 mL | Unclear; 33% defecate on ground |
| Handzel 1998 BGD | 48% tap, 52% tube- well; 61% paid for drinking wa- ter | Improved | Sufficient | Sufficient | Baseline geometric mean 138. 1 faecal coloform counts/100 mL | Unimproved |
| Jain 2010 GHA | 95% of house-holds use tap, 84% surface water, 46% wells, 35% rainwater, 25% borehole | Improved and unimproved | Unclear | Unclear | Baseline: median E. coli MPN 93/ 100 mL | Unimproved |
| Kirchhoff 1985 BRA | Pond water stored in clay pots after filtering with cloth | Unimproved | Unclear | Insufficient | Source water: mean 970 faecal coliforms/ 100 mL | Unimproved |
| Luby 2004 | Tanker trucks, munici- pal taps (house- hold and com- munity level) | Mostly unimproved | Unclear | Unclear | Baseline: approximately 60% of stored drinking water samples were free of <i>E. coli</i> | Improved |
| Luby 2006 | Tanker trucks, munici- pal taps (house- hold and com- mu- nity level), water bearer, boreholes | Mostly improved | Unclear | Unclear | Not tested | Improved |

Table 7. POU chlorination: primary drinking water supply and sanitation facilities (Continued)

| Lule 2005 UGA | 16% surface or shallow wells, 50% protected springs, 49% boreholes or taps | Unimproved | Sufficient | Sufficient | Source mean E. coli counts: 11/100 mL | Improved |
|-----------------------|--|------------|------------|------------|--|--|
| Mahfouz 1995 KSA | Shallow wells | Unimproved | Unclear | Unclear | Source: 92% positive with <i>E. coli</i> ; precise level not reported | Improved |
| Mengistie 2013 ETH | 50% well, 41% spring, 9% river | Unimproved | Unclear | Unclear | Baseline: E. coli MPN 70/ 100 mL | Unimproved |
| Opryszko 2010 | 35% use unprotected dug wells | Unimproved | Sufficient | Sufficient | Not tested | Unclear |
| Quick 1999 BOL | Shallow uncovered wells; 38% treated water | Unimproved | Unclear | Unclear | Source water: median colony count <i>E. coli</i> : 57, 050/100 mL | Unim- proved, but 47% used latrine |
| Quick 2002 ZMB | Shallow wells; some boiling | Unimproved | Unclear | Unclear | Source water: median colony count <i>E. coli</i> : 34/ 100 mL | Unclear |
| Reller 2003 | Surface water from shallow wells, rivers and springs | Unimproved | Unclear | Unclear | Baseline drinking water: median colony count <i>E. coli</i> 63/100 mL | Unclear |
| Semenza 1998 UZB | Households with- out piped water (procured from street tap, neigh- bour tap, well, vendor, or river) | Unimproved | Unclear | Unclear | Source water: 54 coliform colonies/100 mL | Unclear |

¹'Improved' includes household connection, public standpipe, borehole, protected dug well, protected spring, rainwater collection; 'unimproved' includes unprotected well, unprotected spring, vendor-provided water, bottled water; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

²'Sufficient' means located within 500 m, queuing no more than 15 minutes, no more than three minutes to fill 20 L container, and maintained so available consistently; 'insufficient' means that it does not meet any of above; and 'unclear' means unclear or not reported; definition based minimum standards established by The Sphere Project 2011.

Table 8. Summary of findings: POU chlorination

POU chlorination compared with no intervention for preventing diarrhoea

Patient or population: adults and children **Settings:** low- and middle-income countries

Intervention: distribution of chlorine for POU water treatment and instruction on use

Comparison: no intervention

| Outcomes | Illustrative compara | ntive risks* (95% CI) | Relative effect (95% CI) | Number of partici- | dence | |
|--|-------------------------------------|--|-------------------------------|-----------------------|-------------------------------------|--|
| | Assumed risk | Corresponding risk | | (studies) | (GRADE) | |
| | No intervention | POU Chlorination | | | | |
| Diarrhoea episodes cluster-RCTs | | 2.3 episodes per year (2.0 to 2.7) | RR 0.77 (0.65 to 0.91) | 30,746 (14 trials) | ⊕⊕⊖⊖ low ^{1,2,3,4} | |
| Diarrhoea episodes CBA studies | 3 episodes per per- son per year | 1.5 episodes per year (1.0 to 2.3) | RR 0.51 (0.34 to 0.75) | 3948 (2 studies) | ⊕○○○ very low ^{5,6,7,8} | |

The basis for the **assumed risk** is provided in the 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).

CI: confidence interval; RR: risk ratio.

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.

³'Sufficient' means a minimum of 15 L/day/person; 'insufficient' means less than 15 L/day/person; and 'unclear' means unclear or not reported; definition based on minimum standards established by The Sphere Project 2011.

⁴'Improved' means connection to a public sewer or septic system, pour flush latrine, simple pit latrine, or ventilated improved pit latrine; 'unimproved' means service or bucket latrine, public latrines, open latrines; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

The assumed risk is based on 2.9 episodes/child year in 2010 (Fischer Walker 2012).

¹Downgraded by 1 for serious risk of bias: as diarrhoea episodes were reported by participants this outcome is susceptible to bias from lack of blinding. Only two of these studies blinded participants and outcome assessors to the treatment allocation, and these two studies found no evidence of an effect with chlorination.

²Downgraded by 1 for serious inconsistency: statistical heterogeneity was very high (I² statistic = 91%). In a subgroup analysis by compliance with the intervention (assessed by measurements of residual chlorine in drinking water) found larger effects in the studies with better compliance.

Table 9. POU flocculation/disinfection: description of the interventions

| Study ID | Study design | Setting | Intervention a | reas | | Control areas | |
|---------------------|--------------|-------------------------|---|-----------------------------|---|--|-----------------------------|
| | | | Water quality intervention | Health promotion activities | Compliance | Water source | Health promotion activities |
| Chiller 2006 GTM | Cluster-RCT | Rural villages | Provided households with a large spoon and a widemouthed bucket for mixing, a narrowtopped vessel with a lid for storing treated water and provided households with sachets of the flocculant-disinfectant every week | None | - | 31% tap, 40% river or spring and 25% well. | None |
| Crump 2005b KEN | Cluster-RCT | Rural villages | Each week households were given sa- chets of the flocculant- disinfectant | None | 44% compliance during unannounced weekly visits measured by residual chlorine | 50% pond, 49% river and 2% spring | None |
| Doocy 2006 LBR | Cluster-RCT | Liberian camps for dis- | House- holds received | None | 85% compli- ance based on | Received a funnel and an | None |

³No serious indirectness: these studies are mainly from low- and middle-income countries (the Gambia, India, Kenya, Bangladesh, Ghana, Brazil, Pakistan, Uganda, Saudi Arabia, Ethiopia, Afghanistan, Bolivia, Guatemala, and Uzbekistan). The interventions consisted of free distribution of chlorine (every one to four weeks) plus instructions on how to use it. In some cases, the intervention included hygiene education and storage containers in which to treat and store water.

⁴No serious imprecision: the average effect suggests POU chlorination may reduce diarrhoea episodes by about a quarter. The analysis is adequately powered to detect this effect.

⁵Downgraded by 1 for serious risk of bias: as diarrhoea episodes were reported by participants this outcome is susceptible to bias from lack of blinding. None of these studies blinded participants and outcome assessors to the treatment allocation.

⁶Downgraded by 1 for serious inconsistency: statistical heterogeneity was very high (I² statistic = 63%).

⁷Downgraded by 1 for serious indirectness: there are only two studies (three comparisons) from Pakistan and Zambia.

⁸No serious imprecision.

Table 9. POU flocculation/disinfection: description of the interventions (Continued)

| | | placed persons | a bucket and large mixing spoon for preparation, a decanting cloth, a funnel and a storage container with a narrow opening and lid. Each household received a maximum of 21 flocculation-disinfectant packets per week | | residual chlo- rine sampling | identical storage container | |
|---------------------|-------------|-------------------------|--|--|---------------------------------|--|------|
| Luby 2006b PAK | Cluster-RCT | Squatter settlements | Provided households with floccu- lant-disinfec- tant sachets, a water ves- sel and soap. Weekly distri- butions of sa- chets | Field workers educated neighbour-hoods about health problems resulting from hand and water contamination and instructed households on how and when to wash hands | Yes, though rate unclear | Municipal supply at household (33%), at community tap (37%), tanker truck (12%), water bearer (13%) and tube well (5%) | None |
| Luby 2006c PAK | Cluster-RCT | Squatter settlements | | educated | Yes, though rate unclear | Municipal supply at household (33%), at community tap (37%), tanker truck (12%), water bearer (13%) and tube well (5%) | None |
| Reller 2003a GTM | Cluster-RCT | Rural villages | Weekly distri- bution of floc- culant-dis- | Field workers discussed the impor- | 27% compliance measure by | 33% tap, 46% river or spring, 21% well. | None |

Table 9. POU flocculation/disinfection: description of the interventions (Continued)

| | | | gave 2 cloths initially, | tance of water treatment and demonstrated the water prepara- tion process | rine > 0.1 mg/ L on unan- | | |
|---------------------|-------------|----------------|--|--|--|---|------|
| Reller 2003d GTM | Cluster-RCT | Rural villages | culant-dis- infectant and gave 2 cloths initially, which could | discussed the importance of water treatment and demonstrated the water prepara- | ance measure by residual chlo- rine > 0.1 mg/ | - | None |

Table 10. POU flocculation/disinfection: primary drinking water supply and sanitation facilities

| Trial | Description | Source ¹ | Access to source 2 | Quantity available ³ | Ambient H ₂ O quality | Sanitation ⁴ |
|---------------------|---|---------------------|--------------------|---------------------------------|---|---------------------------------------|
| Chiller 2006 GTM | Rivers, springs, taps, and wells | Unclear | Unclear | Sufficient | 98% of source samples contained <i>E. coli</i> ; precise level not reported | Mostly unimproved |
| Crump 2005b KEN | 50% ponds, 49% rivers | Unimproved | Unclear | Insufficient | Baseline mean 98 <i>E. coli </i> 100 mL | Unclear; 33% defecate on ground |
| Doocy 2006 LBR | Surface sources and some tap stands | Unimproved | Unclear | Insufficient | Source water: 88% samples tested positive for faecal contamination; pre- | Unimproved |

Table 10. POU flocculation/disinfection: primary drinking water supply and sanitation facilities (Continued)

| | | | | | cise level not reported | |
|---------------------|--|--------------------|---------|---------|---|----------|
| Luby 2006b PAK | Tanker trucks, munici- pal taps (house- hold and com- mu- nity level), water bearer, boreholes | Mostly improved | Unclear | Unclear | Not tested | Improved |
| Reller 2003a GTM | Surface water from shallow wells, rivers and springs | Unimproved | Unclear | Unclear | Baseline drinking water: median colony count <i>E. coli</i> 63/100 mL | Unclear |

¹'Improved' includes household connection, public standpipe, borehole, protected dug well, protected spring, rainwater collection; 'unimproved' includes unprotected well, unprotected spring, vendor-provided water, bottled water; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

Table 11. Summary of findings: POU flocculation and disinfection

| POU water flocculation and disinfection compared with no intervention for preventing diarrhoea | | | | | | | | |
|--|--------------------------|-------------------------------------|-----------------------------|------------------------|---------|--|--|--|
| Patient or popular | tion: adults and childre | en | | | | | | |
| U | middle-income countr | | | | | | | |
| | | bining water flocculation | n and disinfection and | l instructions on use | | | | |
| Comparison: no in | ntervention | | | | | | | |
| Outcomes | Illustrative compa | rative risks* (95% CI) | Relative effect (95% CI) | Number of participants | dence | | | |
| | Assumed risk | Corresponding risk | | (studies) | (GRADE) | | | |
| | No intervention | Water flocculation and disinfection | | | | | | |

²'Sufficient' means located within 500 m, queuing no more than 15 minutes, no more than three minutes to fill 20 L container, and maintained so available consistently; 'insufficient' means that it does not meet any of above; and 'unclear' means unclear or not reported; definition based minimum standards established by The Sphere Project 2011.

³'Sufficient' means a minimum of 15 L/day/person; 'insufficient' means less than 15 L/day/person; and 'unclear' means unclear or not reported; definition based on minimum standards established by The Sphere Project 2011.

⁴'Improved' means connection to a public sewer or septic system, pour flush latrine, simple pit latrine, or ventilated improved pit latrine; 'unimproved' means service or bucket latrine, public latrines, open latrines; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

Table 11. Summary of findings: POU flocculation and disinfection (Continued)

| Diarrhoea | 3 episodes per per- | 2.1 episodes per | RR 0.69 | 11,788 | ⊕⊕⊕⊜ |
|--------------|---------------------|------------------|----------------|------------|-----------------------------|
| episodes | son per year | person per year | (0.58 to 0.82) | (4 trials) | moderate ^{1,2,3,4} |
| Cluster-RCTs | | (1.7 to 2.5) | | | |

The basis for the **assumed risk** is provided in the 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).

CI: confidence interval; RR: risk ratio.

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 assumed risk is based on 2.9 episodes/child year in 2010 (Fischer Walker 2012).

¹Downgraded by 1 for serious risk of bias: as diarrhoea episodes were reported by participants this outcome is susceptible to bias from lack of blinding. None of these studies blinded participants and outcome assessors to the treatment allocation.

²No serious inconsistency: In the complete analysis of five trials statistical heterogeneity was very high (I² statistic = 99%). However, this heterogeneity was related to a single trial showing very large effects conducted in an emergency setting in Liberia possibly due to epidemic diarrhoea. When this trial was removed as an outlier, there was a smaller, but more consistent effect.

³No serious indirectness: the studies were conducted in rural areas in Guatemala (two studies), and Kenya (one study), one trial was from a camp for displaced persons in Liberia and one from squatter settlements in Pakistan. Sanitation was improved in only one of these studies.

⁴No serious imprecision: all five studies found benefits with flocculation. The 95% CI of the pooled effect includes the possibility of no effect, but this imprecision is a result of the heterogeneity between studies.

Table 12. POU filtration: description of interventions

| Study ID | Interven- tion sub- group | - Study ub- design | , | Intervention areas | | | Control areas | |
|-------------------|---------------------------------|-----------------------|-------|--|--|-----------------|-----------------|--|
| | | | | _ | Health promotion activities | _ | Water source | Health promotion activities |
| Abebe 2014 ZAF | Ceramic filter | Cluster- RCT | Rural | pregnated with silver nanoparti- cles with safe | cation about safe wa- ter and hy- giene and in- | Not reported | , com- | usual clini- cal care in- cluding edu- cation about |

Table 12. POU filtration: description of interventions (Continued)

| Brown 2008a KHM | Ceramic filter | Cluster- RCT | Rural | CWP (Cam- bodian Ce- ramic Water Purifier) in- cluding safe storage con- tainer | None | 98% compliance measured by self-report | Surface water (55%) and ground water (48%) during the dry sea- son and sur- face water (45%) , ground water (48%) and rain wa- ter (73%) dur- ing the rainy season | None |
|-----------------------|---------------------|-----------------|------------------------------------|--|------|---|--|------|
| Brown 2008b KHM | Ceramic filter | Cluster- RCT | Rural | CWP- Fe (iron-rich ceramic wa- ter purifier) in- cluding safe storage con- tainer | None | 98% compliance measured by self-report | Surface water (55%) and ground water (48%) during the dry sea- son and sur- face water (45%) , ground water (48%) and rain wa- ter (73%) dur- ing the rainy season | None |
| Clasen 2004b BOL | Ceramic filter | Cluster- RCT | Rural | Ceramic fil- ters in- cluding im- proved stor- age | None | 67% of households had filters in regular use | 68% had taps and 11% boiled water. | None |
| Clasen 2004c BOL | Ceramic fil- ter | Cluster- RCT | Rural | Ceramic fil- ters in- cluding im- proved stor- age | None | 100% of intervention households' water free of TTC | Water from canal (52%) , river (35%) or rainwater (4%) | None |
| Clasen 2005 COL | Ceramic fil- ter | Cluster- RCT | Ru- ral and ur- ban affected | Ceramic water filter system includ- | None | Not reported | River (27. 6%), | None |

Table 12. POU filtration: description of interventions (Continued)

| | | | by conflict | ing im- proved stor- age | | | rainwater (12.1%), yard tap (67. 2%). 70.7% claimed to treat water | |
|------------------------------|----------------|-----------------|-------------|---|---|---|--|--|
| du Preez 2008 ZAF/ ZWE | Ceramic filter | Cluster- RCT | Rural | Ceramic fil- ters in- cluding im- proved stor- age | None | 55% compliance measured by water quality (approximate compliance across intervention households in Zimbabwe and South Africa) | Protected water source (53.8%) and unprotected water source (46.2%) | None |
| Lindquist 2014a BOL | Ceramic filter | Cluster- RCT | Peri-urban | Received a PointONE Filter and a 30 L bucket (with lid) | Participants were in- structed on diar- rhoeal trans- mission (bi- ological ver- sus cul- tural beliefs- based), pre- vention and treatment | 97% compliance based on re- ported use | 83% used water from tanker trucks and 12% from water cool- ers | Received weekly messages on life skills and attitudes. Also were instructed on diarrhoeal transmission, prevention and treatment |
| Lindquist 2014b BOL | Ceramic filter | Cluster- RCT | Peri-urban | | | 90% compliance based on re- ported use | water from tanker trucks and | skills and attitudes. Also |

Table 12. POU filtration: description of interventions (Continued)

| | | | | | filtration), vitamin A, hygienic food prepa- ration and cleaning, and parasite prevention. | | | |
|-------------------------|-----------------|-----------------|-------|--|--|--|--|----------------------|
| URL 1995a GTM | Ceramic filter | Cluster- RCT | Rural | Handmade ceramic wa- ter filter | None | 87% to 93% use of filter by children | Major- ity of house- holds col- lected water from house- hold tap (not chlori- nated) | None |
| URL 1995b GTM | Ceramic filter | Cluster- RCT | Rural | Handmade ceramic wa- ter filter | Education on nutrition (ORS, basic nutri- tion and ma- ternal and child nutri- tion), health (hygiene) and family values | As above | Major- ity of house- holds col- lected water from house- hold tap (not chlori- nated) | None |
| Fabiszewski 2012 HND | Sand filtration | Cluster- RCT | Rural | Hydraid plastic- housing BioSand fil- ter (BSF) + 20 L water jug | Training for the use and mainte- nance of the BSF and general edu- cation about hygiene and sanitation | Not reported | Among all study participants- the main source of drinking water were: protected water sources (49% to 69% house-holds per month), protected sources (24% to 50% per month) | nance of the BSF and |

Table 12. POU filtration: description of interventions (Continued)

| | | | | | | | , piped water (1% to 11% per month), and rainwater (0% to 2% per month). | |
|-------------------------|----------------------|-----------------|-------------------------|---|--|---|--|--|
| Stauber 2009 DOM | Sand filtration | Cluster- RCT | Semi-rural and urban | Received a biosand fil- ter and safe storage con- tainer | Nothing | Water quality test- ing, however no interven- tion house- hold level compliance reported | ported treat- ing drinking | None |
| Stauber 2012a KHM | Sand filtra- tion | Cluster- RCT | Rural | Plas- tic biosand filter. HHs were asked to pay USD 10 for the filter | Health and hygiene ed- ucation ses- sions | 89% compliance measured by household-reported use at least 3 times per week | Improved water sources during the dry season (7.1%) and during the rainy season (88.9%) . 49.5% reported boiling drinking water | Health and hygiene ed- ucation ses- sions |
| Stauber 2012b GHA | Sand filtra- tion | Cluster- RCT | Rural | Plastic biosand fil- ter | Not specified | 97% compliance measured by household- reported use | Use surface water during dry season (95%) and use surface water during rainy season (70.6%). 96.5% reported sieving drinking water through cloth | nothing |
| Tiwari 2009 KEN | Sand filtra- tion | Cluster- RCT | Rural | Provided with the concrete | At each visit, three oral re- hydra- | | All control houses reported | At each visit, three oral re- hydra- |

Table 12. POU filtration: description of interventions (Continued)

| | | | | BioSand Filter | tion packets and instruc- tions were provided | | drinking river or unprotected spring water; drink rainwater (96.6%), drink improved source (24. 1%). 34.5% reported boiling drinking water | tion packets and instruc- tions were provided |
|---------------------|------------------------|-----------------|------------|--|--|---|--|--|
| Boisson 2009 ETH | LifeStraw® Personal | Cluster- RCT | Rural | A LifeS-traw® personal pipe-style water treatment device was given to each member of the household >6 months and encouraged to use it at home and away from home | None | 13% report use today | The primary drinking water source for 84% was from spring, 12% from rivers, 2.5% from hand dug wells and 4% from communal taps | None |
| Boisson 2010 DRC | LifeStraw® Family | Cluster- RCT | Rural | Households received a LifeS- traw® Fam- ily filters | None | 76% compliance measured by self-re-port use to-day or yesterday (at 14 month follow-up) | Received a placebo fil- ter. | None |
| Peletz 2012 ZMB | LifeStraw® Family | Cluster- RCT | Peri-urban | Households received a LifeS- traw® Fam- ily filter and two 5 L safe | None | 87% compliance measured by improved water quality | 46% use unprotected dug wells, 19% boreholes, 17% public | None |

Table 12. POU filtration: description of interventions (Continued)

| | | | | storage containers | | | stand- pipes, 12% protected dug well, 5% piped into home or yard and 2% surface water | |
|---------------------|-----------------------------------|-----------------|-------|--|-------------------------|--|---|---|
| Colford 2002 USA | Plumbed in filter | Cluster- RCT | Urban | Installation of water treatment devices to 1 tap in HH that include: a 1-micron absolute prefilter cartridge and a UV lamp | None | 96% compliance measured by not dropping out of study (plumbedin unit) | Sham device | None |
| Colford 2005 USA | Plumbed in filter | Cluster- RCT | Urban | micron filter | the current CDC safe | pliance measured by not | Sham device | All participants received the current CDC safe drinking water guidelines for immuno-compromised persons |
| Colford 2009 USA | Plumbed in filter | Cluster- RCT | Urban | Installation of filter (1- micron filter and a UV lamp) to main faucet of household | None | 83% compliance measured by not dropping out of study (filter attached to kitchen sink) | Sham device | None |
| Rodrigo 2011 AUS | Ceramic fil- ter/plumbed in | | Urban | Bench-top silver im- pregnated ceramic wa- ter treat- | None | Not reported | Sham water treatment unit | None |

Table 12. POU filtration: description of interventions (Continued)

| quired participants to use fill it but then house- holds that had rainwa- ter piped into kitchen were offered an under sink unit |
|--|
|--|

Table 13. POU filtration: primary drinking water supply and sanitation facilities

| Trial | Description | Source ¹ | Access to source | Quantity available ³ | Ambient H ₂ O quality | Sanitation ⁴ |
|---------------------|---|------------------------------------|------------------|---------------------------------|---|-------------------------|
| Abebe 2014 ZAF | In-home taps or community taps | Improved | Sufficient | Unclear | 80% of house- holds had con- tamina- tion between 10 to 10000 CFUs/ 100 mL | Unclear |
| Brown 2008 | 62% households rely on surface water during dry sea- son and 55% rely on surface water during rainy sea- son | Unimproved | Unlcear | Unclear | Baseline not reported. Control households: Geometric mean 600 <i>E. coli</i> /100 mL | Improved |
| Clasen 2004b BOL | 80% yard taps supplied by untreated sur- face source, 20% directly from un- treated surface sources | 80% improved, 20% unimproved | Sufficient | Sufficient | Baseline arithmetic mean 86 TTC/100 mL | Unimproved |
| Clasen 2004c BOL | Irrigation canals and other surface sources | Unimproved | Sufficient | Sufficient | Baseline arithmetic mean 797 TTC/100 mL | Unimproved |

Table 13. POU filtration: primary drinking water supply and sanitation facilities (Continued)

| Clasen 2005 COL | 67% yard tap from municipal- ity (not treated), 28% river, 12% rainwater | Unimproved | Unclear | Unclear | Baseline not reported. Control house-holds: arithmetic mean 151 TTC/100 mL | · · · · · · · · · · · · · · · · · · · |
|--------------------------|---|-------------------------|------------|---------|--|---------------------------------------|
| du Preez 2008 ZAF/ZWE | Protected wells | Improved | Sufficient | Unclear | Baseline not reported. Control house-holds: 30% samples post-intervention met WHO guidelines for water quality | Improved |
| Lindquist 2014 | Municipal sup- ply | Improved | Sufficient | Unclear | Not tested | Unimproved |
| URL 1995 | House- hold tap (27%), public tap (21%) , well (23%) | Improved | Unclear | Unclear | Range 5 to 260; average 106 fae- cal co- liforms/100 mL across three sites | Improved |
| Fabiszewski 2012 HND | 49% to 69% households use unprotected sources, 24% to 50% use protected sources, 1% to 11% piped water, 0% to 2 % rainwater | Improved and unimproved | Unclear | Unclear | Geometric mean <i>E. coli</i> concentrations of both unprotected and protected sources were > 100 MPN/100 mL | Unimproved |
| Stauber 2009 DOM | Unclear | Unclear | Unclear | Unclear | Baseline: geometric mean 21 MPN <i>E. coli </i> 100 mL | Improved |
| Stauber 2012a KHM | 77% used improved water source during dry season, 89% during rainy season | Improved | Unclear | Unclear | Baseline: geo- metric mean 27. 5 CFU/100 mL | Unimproved |

Table 13. POU filtration: primary drinking water supply and sanitation facilities (Continued)

| Stauber 2012b GHA | Surface water 70% in dry season, 95% in rainy season | Unimproved | Unclear | Unclear | Baseline: geometric mean 792 or 832 <i>E. coli</i> /100 mL for control and in- terven- tion households, respectively | Unimproved |
|----------------------|---|-------------------------|------------|------------|--|------------|
| Tiwari 2009 KEN | Primar- ily river water; 27% drink pro- tected sources | Unimproved | Unclear | Unclear | Baseline not reported. Control households: 88.9 faecal coliforms/ | Unclear |
| Boisson 2009 ETH | 84% springs, 12% river, 2% handdug well, 4% com- munal tap | Unimproved | Unclear | Unclear | Baseline arithmetic mean 449 TTC/100 mL | Unimproved |
| Boisson 2010 DRC | 97% surface water, 38% rainwater, 16% springs | Unimproved | Unclear | Unclear | Source drinking water: 75% of household sam- ples > 1000 TTC/ 100 mL | Unimproved |
| Peletz 2012 ZMB | 46% unprotected dug wells, 22% taps, 16% borehole or protected dug well, 2% surface water | Improved and unimproved | Unclear | Unclear | Unfiltered water: Geometric mean 190 TTC/100 mL | Unimproved |
| Colford 2002 USA | Household taps supplied by municipal water treatment | Improved | Sufficient | Sufficient | Data from water treatment plant: met US fed- eral and Califor- nia drinking wa- ter standards | Improved |
| Colford 2005 USA | Household taps supplied by municipal water treatment | Improved | Sufficient | Sufficent | Data from water treatment plant: met US fed- eral drinking wa- ter standards | Improved |

Table 13. POU filtration: primary drinking water supply and sanitation facilities (Continued)

| Colford 2009 USA | Household taps supplied by municipal water treatment | Improved | Sufficient | Sufficient | Data from water treatment plant: met US fed- eral drinking wa- ter standards | Improved |
|---------------------|---|----------|------------|------------|--|----------|
| Rodrigo 2011 AUS | Untreated rain- water | Improved | Sufficient | Sufficient | Not tested | Improved |

Abbreviations: TTC: thermotolerant coliforms, MPN: most probable number, CFU: colony-forming units

Table 14. Summary of findings: POU filtration

| POU filtration compared with no intervention for preventing diarrhoea |
|---|
| |

Patient or population: adults and children Settings: low-, middle- and high-income countries

Intervention: distribution of water filters and instructions on use

Comparison: no intervention

| Outcomes | Illustrative compara | ntive risks* (95% CI) | Relative effect (95% CI) | Number of participants | Quality of the evidence | |
|---------------------------------|-------------------------------------|---|-------------------------------|------------------------|---|--|
| | Assumed risk | Corresponding risk | | (studies) | (GRADE) | |
| | No intervention | Water filtration | | | | |
| Diarrhoea | 3 episodes per per- | All filters | RR 0.48 | 15,582 | $\oplus \oplus \oplus \bigcirc$ | |
| episodes Cluster-RCTs | son per year | 1.4 episodes per person per year (1.1 to 1.8) | (0.38 to 0.59) | (18 trials) | 1.4 episodes per person per year (1.1 to 1.8) | |
| | 3 episodes per per- son per year | Ceramic filters | RR 0.39 (0.29 to 0.53) | 5763 (8 trials) | ⊕⊕⊕⊜ moderate ^{2,4,5,6} | |

¹'Improved' includes household connection, public standpipe, borehole, protected dug well, protected spring, rainwater collection; 'unimproved' includes unprotected well, unprotected spring, vendor-provided water, bottled water; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

²'Sufficient' means located within 500 m, queuing no more than 15 minutes, no more than three minutes to fill 20 L container, and maintained so available consistently; 'insufficient' means that it does not meet any of above; and 'unclear' means unclear or not reported; definition based minimum standards established by The Sphere Project 2011.

³'Sufficient' means a minimum of 15 L/day/person; 'insufficient' means less than 15 L/day/person; and 'unclear' means unclear or not reported; definition based on minimum standards established by The Sphere Project 2011.

⁴'Improved' means connection to a public sewer or septic system, pour flush latrine, simple pit latrine, or ventilated improved pit latrine; 'unimproved' means service or bucket latrine, public latrines, open latrines; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

Table 14. Summary of findings: POU filtration (Continued)

| 1.1 episo person per (0.8 to 1.5) | r year | | | 1.1 episodes per person per year (0.8 to 1.5) |
|---|------------|---------|------|---|
| Biosand fi | lters R | RR 0.47 | 5504 | 000 |
| 1.4 episo person per (1.2 to 1.7) | r year | | | 1.4 episodes per person per year (1.2 to 1.7) |
| LifeStraw | ®filters R | RR 0.69 | 3259 | $\oplus \oplus \bigcirc \bigcirc$ |
| 2.1 episo person per (1.5 to 2.8) | r year | | | 2.1 episodes per person per year (1.5 to 2.8) |
| Plumbed f | filters R | RR 0.73 | 1056 | $\oplus \oplus \oplus \bigcirc$ |
| 2.2 episo person per (1.6 to 3.1) | r year | | | 2.2 episodes per person per year (1.6 to 3.1) |

The basis for the **assumed risk** is provided in the 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).

CI: confidence interval; RR: risk ratio.

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 assumed risk is based on 2.9 episodes/child year in 2010 (Fischer Walker 2012).

¹Downgraded by 1 for serious risk of bias: as diarrhoea episodes were reported by participants, this outcome is susceptible to bias from lack of blinding. Only five studies blinded participants and outcome assessors to the treatment allocation and only one found an effect of the intervention.

²No serious inconsistency: statistical heterogeneity was very high, however there is consistency in the direction of the effect.

³No serious indirectness: these studies are from a variety of low-, middle-, and high-income countries (South Africa, Ethiopia, Democratic Republic of Congo, Cambodia, Bolivia, Colombia, USA, Australia, Honduras, Zimbabwe, Zambia, Dominican Republic, Ghana, Kenya and Guatemala).

⁴No serious imprecision.

⁵Downgraded by 1 for serious risk of bias: as diarrhoea episodes were reported by participants, this outcome is susceptible to bias from lack of blinding. Only one of these studies, Rodrigo 2011 AUS, blinded participants and outcome assessors to the treatment allocation. ⁶No serious indirectness: these studies are from a variety of low-, middle-, and high-income countries (South Africa, Cambodia, Bolivia, Colombia, Zimbabwe, Guatemala and Australia). The interventions consisted of distribution of water filters (which included a safe storage chamber) plus instructions on how to use them. In some cases, the intervention included hygiene education.

⁷Downgraded by 1 for serious risk of bias: as diarrhoea episodes were reported by participants, this outcome is susceptible to bias from lack of blinding. None these studies blinded participants and outcome assessors to the treatment allocation.

Table 15. POU solar disinfection (SODIS): description of the interventions

| Study ID | Study design | Setting | Intervention a | reas | | Control areas | |
|----------------------|--------------|------------|---|-----------------------------|---|--|-----------------------------|
| | | | Water quality intervention | Health promotion activities | Compliance | Water source | Health promotion activities |
| Conroy 1996 KEN | Quasi-RCT | Rural | Children were given two 1.5 L plastic bottles and told to keep the bottles on the roof of the hut throughout the day in full sunlight | None | dom checks by project work- ers un- | L plastic bot- tles and told to keep the bot- | None |
| Conroy 1999 KEN | Quasi-RCT | Rural | Mothers were given plastic bottles and told to keep the bottles on the roof of the hut throughout the day in full sunlight | None | Not reported | Mothers were given plas- tic bottles and told to keep the bottles in- doors | None |
| du Preez 2010 ZAF | Cluster-RCT | Peri urban | Received two 2 L polyethy- lene terephta- late (PET) bottles for each child. Car- ers were in- | None | 25% compliance measured by participants filling out diarrhoeal diaries at least 75% of the time | bottles and | None |

⁸No serious inconsistency: there was no statistical heterogeneity between studies, I² statistic = 0%.

⁹No serious indirectness: the studies were conducted in a variety of rural and urban settings in a variety of low- and middle-income countries (Honduras, Dominican Republic, Cambodia, Ghana and Kenya). The interventions consisted of distribution of water filters plus instructions on how to use them. In some cases, the intervention included hygiene education and a separate storage vessel.

¹⁰Downgraded by 1 for serious risk of bias: as diarrhoea episodes were reported by participants this outcome is susceptible to bias from lack of blinding. Only one of these studies, Boisson 2010 DRC, blinded participants and outcome assessors to the treatment allocation and found no evidence of effect of the filter.

¹¹Downgraded by 1 for some indirectness, the studies were only performed in three sub-Saharan African countries (Ethiopia, Democratic Republic of Congo, and Zambia).

¹²No serious risk of bias: the three studies blinded participants and outcome assessors to the treatment allocation.

¹³Downgraded by 1 for some indirectness, the three studies were only performed in the USA in water conditions that presumed to meet US EPA standards.

Table 15. POU solar disinfection (SODIS): description of the interventions (Continued)

| | | | structed to fill one bottle and place it in full, unobscured sunlight for a minimum of 6 h every day | | | | |
|-----------------------|-------------|-------------------------|---|--|--|--|------|
| du Preez 2011 KEN | Cluster-RCT | Peri urban and rural | Received two 2 L PET bottles for each child. Carers were instructed to fill one bottle and place it in full, unobscured sunlight for a minimum of 6 h every day | None | Not specified. | No SODIS bottles and maintain their usual practices | None |
| Mäusezhal 2009 BOL | Cluster-RCT | Rural | Households were supplied regularly with clean, PET bottles. They were instructed to expose the waterfilled bottles for at least 6 h to the sun | about the im- portance and benefits of drinking only treated water, the germ-dis- | 32% compliance measured by observation | ing water from | None |
| McGuigan 2011 KHM | Cluster-RCT | Rural | tic bottles for each child and a sheet of cor- rugated iron on which | verbal and written infor- | 90% (5% of chil-dren having < 10 months of follow-up and 2.3% having < 6 months) | of the house- holds (97%) obtained wa- ter from un- | None |

Table 15. POU solar disinfection (SODIS): description of the interventions (Continued)

| pose them to sunlight. Car- ers were in- structed to fill one bottle and place it in full, unobscured sunlight for a minimum of 6 | fection process and its effect on the micro- bial quality of their drinking water and sub- sequently the health of their children | drew water from shallow tube wells fitted with hand pumps. The remainder used unprotected wells or surface | |
|---|---|--|--|
| h every day | | or surface ponds | |

Table 16. POU solar disinfection (SODIS): primary drinking water supply and sanitation facilities

| Trial | Description | Source ¹ | Access to source | Quantity available ³ | Ambient H ₂ O quality | Sanitation ⁴ |
|----------------------|--|----------------------|------------------|---------------------------------|--|-------------------------|
| Conroy 1996 KEN | Open water holes, tank fed by untreated piped water sup- ply. | Unimproved | Unclear | Unclear | Source water: 10 ³ CFU/100 mL | Unclear |
| Conroy 1999 KEN | Open water holes, tank fed by untreated piped water sup- ply. | Unimproved | Unclear | Unclear | Source water: 10 ³ CFU/100 mL | Unclear |
| du Preez 2010 ZAF | 39% standpipes, 28% protected bore- hole, 10% un- protected bore- holes, protected springs | Mostly improved | Sufficient | Sufficient | Baseline not reported. Intervention households: 62% of samples met WHO guide- lines for water quality; no significant difference from control households | Unclear |
| du Preez 2011 KEN | Spring, protected and unprotected dug wells protected, canals, other | Mostly unimproved | Unclear | Unclear | 50% of samples from stored water had 10 CFU/100 mL or less; no signif- icant difference | Unclear |

Table 16. POU solar disinfection (SODIS): primary drinking water supply and sanitation facilities (Continued)

| | | | | | for intervention and controls | |
|-----------------------|--|------------|------------|------------|---|------------|
| Mäusezhal 2009 BOL | 48% spring, 52% tap, 22% river, 15% rain, 15% dug well | 1 | Sufficient | Sufficient | Not tested | Unimproved |
| McGuigan 2011 KHM | 97% households use unprotected sources: unpro- tected wells, sur- face ponds | Unimproved | Unclear | Unclear | Baseline not reported. Control households: geometric mean 48 CFU/100 mL | Unimproved |

¹ Improved' includes household connection, public standpipe, borehole, protected dug well, protected spring, rainwater collection; 'unimproved' includes unprotected well, unprotected spring, vendor-provided water, bottled water; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

Table 17. Summary of findings: POU solar disinfection (SODIS)

| POU solar disinfection (SODIS) of water compared with no intervention for preventing diarrhoea | | | | | | | | |
|--|------------------------|-----------------------|--------|--|--|--|--|--|
| | | | | | | | | |
| Patient or population: adults and children | | | | | | | | |
| Settings: low- and middle-income countries | | | | | | | | |
| Intervention: distribution of plastic bottles with instructions o | using them to treat wa | ter using the SODIS n | nethod | | | | | |
| Comparison: no intervention | | | | | | | | |
| Companion no metronico. | | | | | | | | |
| | | | | | | | | |

| Outcomes | Outcomes Illustrative comparative risks* (95% CI) | | | Number of participants | dence |
|---------------------------------------|---|---|-------------------------------|------------------------|-------------------------------------|
| | Assumed risk | Corresponding risk | | (studies) | (GRADE) |
| | No intervention | SODIS | | | |
| Diarrhoea episodes Cluster-RCTs | 3 episodes per per- son per year | 1.9 episodes per person per year (1.3 to 2.8) | RR 0.62 (0.42 to 0.94) | 3460 (4 trials) | ⊕⊕⊕⊜ moderate ^{1,2,3,4} |

²'Sufficient' means located within 500 m, queuing no more than 15 minutes, no more than three minutes to fill 20 L container, and maintained so available consistently; 'insufficient' means that it does not meet any of above; and 'unclear' means unclear or not reported; definition based minimum standards established by The Sphere Project 2011.

³'Sufficient' means a minimum of 15 L/day/person; 'insufficient' means less than 15 L/day/person; and 'unclear' means unclear or not reported; definition based on minimum standards established by The Sphere Project 2011.

⁴'Improved' means connection to a public sewer or septic system, pour flush latrine, simple pit latrine, or ventilated improved pit latrine; 'unimproved' means service or bucket latrine, public latrines, open latrines; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

Table 17. Summary of findings: POU solar disinfection (SODIS) (Continued)

| Diarrhoea episodes Ouasi-RCTs | 3 episodes per per- son per year | 2.5 episodes per person per year (2.1 to 2.9) | RR 0.82 (0.69 to 0.97) | 555 (2 studies) | ⊕⊕⊜⊝ low ^{1,5,6,7} |
|-------------------------------------|-------------------------------------|---|-------------------------------|--------------------|--------------------------------|
| Quasi ico is | | (2.1 to 2.)) | | | |

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.

Table 18. POU UV: description of the interventions

| Study ID | Study design | Setting | Intervention are | eas | Control areas | | |
|--------------------|--------------|---------|---|-----------------------------|---|--------------|----------------------------------|
| | | | Water quality intervention | Health promotion activities | Compliance | Water source | Health promo- tion activities |
| Gruber 2013 MEX | Cluster-RCT | Rural | Promotion of the UV Tube disinfection technology and safe storage | Unclear | 51% compliance measured by access to treatment device | Unclear | None |

The assumed risk is based on 2.9 episodes/child year in 2010 (Fischer Walker 2012).

¹Downgraded by 1 for serious risk of bias: as diarrhoea episodes were reported by participants this outcome is susceptible to bias from lack of blinding. None of these studies blinded participants and outcome assessors to the treatment allocation.

²No serious inconsistency: statistical heterogeneity was very high (I² statistic = 89%), however there is consistency in the direction of the effect. This heterogeneity may relate to differences in compliance across the studies, however compliance was not measured in the same way across studies.

³No serious indirectness: the studies were conducted in peri-urban South Africa (one study), peri-urban and rural Kenya (one study), rural Bolivia (one study) and rural Cambodia (one study).

⁴No serious imprecision: the average effect suggests that the intervention may reduce diarrhoea episodes by about one third.

⁵No serious inconsistency: statistical heterogeneity was low (I² statistic = 0%).

⁶Downgraded by 1 for serious indirectness: there are only two studies and both were conducted in the same province in Kenya (one study included children five to 16 years old and the other included children younger than six years old).

⁷No serious imprecision.

Table 19. POU UV: primary drinking water supply and sanitation facilities

| Trial | Description | Source ¹ | Access to source ² | Quantity available | Ambient H ₂ O quality | Sanitation ⁴ |
|--------------------|-------------|---------------------|-------------------------------|--------------------|---|-------------------------|
| Gruber 2013 MEX | Unclear | Unclear | Unclear | Unclear | Baseline: 60% of samples with detectable <i>E. coli</i> | Improved |

¹'Improved' includes household connection, public standpipe, borehole, protected dug well, protected spring, rainwater collection; 'unimproved' includes unprotected well, unprotected spring, vendor-provided water, bottled water; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

Table 20. POU Improved storage: description of the interventions

| Study ID | Study design | Setting | Intervention a | Intervention areas | | | |
|---------------------|--------------|---------|---|-----------------------------|--|--------------|---------------------------------------|
| | | | Water quality intervention | Health promotion activities | Compliance | Water source | Health pro- motion activ- ities |
| Günther 2013 BEN | Cluster-RCT | Rural | Provided house-holds with a new 30 L household water storage with a tap at the bottom, a new plastic container to transport water from the water source to the household and a sign attached to the transport and storage containers which empha- | None | Af- ter 7 months, 88% of house- holds were still using the im- proved storage containers | improved wa- | None |

²'Sufficient' means located within 500 m, queuing no more than 15 minutes, no more than three minutes to fill 20 L container, and maintained so available consistently; 'insufficient' means that it does not meet any of above; and 'unclear' means unclear or not reported; definition based minimum standards established by The Sphere Project 2011.

³'Sufficient' means a minimum of 15 L/day/person; 'insufficient' means less than 15 L/day/person; and 'unclear' means unclear or not reported; definition based on minimum standards established by The Sphere Project 2011.

⁴'Improved' means connection to a public sewer or septic system, pour flush latrine, simple pit latrine, or ventilated improved pit latrine; 'unimproved' means service or bucket latrine, public latrines, open latrines; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

Table 20. POU Improved storage: description of the interventions (Continued)

| | | | sized the im- por- tance of avoid- ing hand-con- tact with the water and to only use water from an im- proved water source | | | | |
|---------------------|-------------|--------------|--|------|---|---|------|
| Roberts 2001 MWI | Cluster-RCT | Refugee camp | All of the participating household's water collection vessels were exchanged for improved buckets (20 L with a narrow opening to limit hand entry). Households were offered 1 improved bucket in exchange for 1 vessel, 2 for 2, and 3 improved buckets for any number of containers > 2. Households were asked never to put their hands in the improved buckets and were shown how to rinse the bucket without hand entry | None | Intervention householders received buckets; actual use was not reported | Provided with 20 L standard ration bucket | None |

Table 21. POU Improved storage: primary drinking water supply and sanitation facilities

| Trial | Description | Source ¹ | Access to source ² | Quantity available ³ | Ambient H ₂ O quality | Sanitation ⁴ |
|---------------------|---|---------------------|-------------------------------|------------------------------------|--|-------------------------|
| Günther 2013 BEN | Public tap or pump | Improved | Sufficient | Unclear | 12% source water contaminated (≥ 1000 CFU per 100 mL) | Unclear |
| Roberts 2001 MWI | Traditional pots or standard ra- tion buckets filled at refugee camp water point | Improved | Unclear | Unclear | Source water: 71% of samples had ≤ 1 faecal coliform/ 100 mL | Unclear |

¹ Improved' includes household connection, public standpipe, borehole, protected dug well, protected spring, rainwater collection; 'unimproved' includes unprotected well, unprotected spring, vendor-provided water, bottled water; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

Table 22. Summary of findings: POU improved water storage

Cluster-RCTs

Improved water storage compared with no intervention for preventing diarrhoea

| Settings: areas with Intervention: distri | Patient or population: adults and children in sub-Saharan Africa Settings: areas with improved water sources Intervention: distribution of improved water containers Comparison: no intervention | | | | | | | |
|---|--|----------------------------------|--------------------------------|------------------------|--------------------------------|--|--|--|
| Outcomes | Outcomes Illustrative comparative risks* (95% CI) | | Relative effect (95% CI) | Number of participants | Quality of the evidence | | | |
| | Assumed risk | Corresponding risk | | (studies) | (GRADE) | | | |
| | No intervention | Water storage | | | | | | |
| Diarrhoea episodes | 3 episodes per per- son per year | 2.7 episodes per person per year | RR 0.91 (0.74 to 1. 11) | 1871 (2 trials) | ⊕⊕⊜⊜ low ^{1,2,3,4} | | | |

(2.2 to 3.3)

²'Sufficient' means located within 500 m, queuing no more than 15 minutes, no more than three minutes to fill 20 L container, and maintained so available consistently; 'insufficient' means that it does not meet any of above; and 'unclear' means unclear or not reported; definition based minimum standards established by The Sphere Project 2011.

³'Sufficient' means a minimum of 15 L/day/person; 'insufficient' means less than 15 L/day/person; and 'unclear' means unclear or not reported; definition based on minimum standards established by The Sphere Project 2011.

⁴'Improved' means connection to a public sewer or septic system, pour flush latrine, simple pit latrine, or ventilated improved pit latrine; 'unimproved' means service or bucket latrine, public latrines, open latrines; and 'unclear' means unclear or not reported; definition based on WHO/UNICEF 2015.

The basis for the **assumed risk** is the median control group risk across studies. 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).

CI: confidence interval; RR: risk ratio.

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.

Table 23. Estimates of household-level interventions after adjustment for non-blinding

| POU intervention | Number of comparisons | Not adjusted for non-blinding | | Adjusted for non-blinding | |
|-------------------------------|-----------------------|-------------------------------|----------------|---------------------------|----------------|
| | | RR | 95% CI | RR | 95% CI |
| All | 55 | 0.56 | (0.46 to 0.68) | 0.70 | (0.64 to 0.77) |
| Chlorination | 19 | 0.72 | (0.61 to 0.84) | 0.80 | (0.69 to 0.92) |
| Filtration | 23 | 0.48 | (0.38 to 0.59) | 0.62 | (0.55 to 0.70) |
| Flocculation and disinfection | 7 | 0.48 | (0.20 to 1.16) | 0.65 | (0.40 to 1.09) |
| SODIS | 6 | 0.68 | (0.53 to 0.89) | 0.80 | (0.60 to 1.01) |

Abbreviation: SODIS: solar disinfection; CI: confidence interval.

The assumed risk is based on 2.9 episodes/child year in 2010 (Fischer Walker 2012).

¹Downgraded by 1 for serious risk of bias: as diarrhoea episodes were reported by participants this outcome is susceptible to bias from lack of blinding. None of these studies blinded participants and outcome assessors to the treatment allocation.

²No serious inconsistency.

³Downgraded by 1 for indirectness: only 2 studies, from rural Benin and a refugee camp in Malawi, have been conducted to assess improved water storage.

⁴No serious imprecision.

Table 24. Potential reasons for finding of no-effect in trials with adequate blinding

| Study | Risk from ambient water quality | Compliance | Other issues |
|--------------------|--|--|--|
| Colford 2002 USA | Very low (USA) | High (Sham filter) | None |
| Colford 2005 USA | Very low (USA) | High (Sham filter) | None |
| Colford 2009 USA | Very low (USA) | High (Sham filter) | None |
| Rodrigo 2011 AUS | Very low (Australia) | Not reported | None |
| Jain 2010 GHA | Low (11 CFU/100 mL) | High (RFC) | Control group received jerry can; 13 week follow-up |
| Kirchhoff 1985 BRA | Very high (mean 16000 FC/dL) | Not reported | Only 112 persons from 16 households; 18 week trial |
| Austin 1993 | High (1871 FC/100 mL) | Low ("50% to 60%") | No test of blinding; not peer reviewed |
| Boisson 2010 DRC | High (75% of samples > 1000 TTC/100 mL) | High, but 73% of adults and 95% of children drank from untreated sources | |
| Boisson 2013 IND | Moderate (mean 122 TTC/100 mL) | Low and inconsistent (32% of samples positive for RFC) | None |

Abbreviations: TTC: thermotolerant coliforms, CFU: colony-forming units, FC: faecal coliforms, RFC: residual free chlorine.

APPENDICES

Appendix I. Search methods: detailed search strategies

| Search set | CIDG SR ^a | CENTRAL | $\mathbf{MEDLINE}^b$ | \mathbf{EMBASE}^b | LILACS ^b |
|------------|---|-------------------------|-------------------------|-------------------------|---|
| 1 | water | WATER PURIFICA- TION | WATER PURIFICA- TION | WATER PURIFICA- TION | water |
| 2 | purification OR treat- ment OR chlorina- tion OR decontami- | | WATER MICROBI- OLOGY | WATER MICROBI- OLOGY | purification OR treat- ment OR chlorina- tion OR decontami- |

| | nation OR filtration OR supply OR stor- age OR consumption | | | | nation OR filtration OR supply OR stor- age OR consumption |
|----|--|--|--|---|--|
| 3 | diarrhea | 1 OR 2 | 1 OR 2 | 1 OR 2 | diarrhea |
| 4 | 1 AND 2 AND 3 | water | water | water | 1 AND 2 AND 3 |
| 5 | - | purification OR treat- ment OR chlorina- tion OR decontami- nation OR filtration OR supply OR stor- age OR consumption OR drink* | purification OR treat- ment OR chlorina- tion OR decontami- nation OR filtration OR supply OR stor- age OR consumption OR drink* | purification OR treat- ment OR chlorina- tion OR decontami- nation OR filtration OR supply OR stor- age OR consumption OR drink\$ | - |
| 6 | - | 4 AND 5 | 4 AND 5 | 4 AND 5 | - |
| 7 | - | 3 OR 6 | 3 OR 6 | 3 OR 6 | - |
| 8 | - | DIARRHEA/ EPIDEMIOLOGY | DIARRHEA/ EPIDEMIOLOGY | DIARRHEA/ EPIDEMIOLOGY | - |
| 9 | - | DIARRHEA/ MICROBIOLOGY | DIARRHEA/ MICROBIOLOGY | DIARRHEA/ PREVENTION | F |
| 10 | - | DIARRHEA/ PREVENTION AND CONTROL | DIARRHEA/ PREVENTION AND CONTROL | waterborne infection\$ | - |
| 11 | - | waterborne infection* | waterborne infection* | cholera OR shigell\$ OR dysenter\$ OR cryptosporidi\$ OR giardia\$ OR Escherichia coli OR clostridium | - |
| 12 | - | INTESTINAL DIS- EASES | INTESTINAL DIS- EASES | ENTEROBACTE- RIACEAE | - |
| 13 | - | cholera OR shigell* OR dysenter* OR cryptosporidi* OR gi- ardia* OR Escherichia coli OR clostridium | cholera OR shigell* OR dysenter* OR cryptosporidi* OR gi- ardia* OR Escherichia coli OR clostridium | 8-12/OR | - |
| 14 | - | ENTEROBACTE- RIACEAE | ENTEROBACTE- RIACEAE | 7 AND 13 | - |

| 15 | - | 8-14/OR | 8-14/OR | LIMIT 14 TO HU- MAN | - |
|----|---|----------|------------------------|------------------------|---|
| 16 | - | 7 AND 15 | 7 AND 15 | - | - |
| 17 | - | - | LIMIT 16 TO HU- MAN | - | - |

^aCochrane Infectious Diseases Group Specialized Register.

Appendix 2. Data extracted from included studies

| Туре | Fields |
|----------------------------|--|
| Trial data | Country and setting (urban, rural) |
| | Number of participants/groups |
| | Unit of randomization, and whether measurement of effect adjusts for clustering where randomization is other than individual |
| | Definition and practices of control group |
| | Type and details of water quality intervention (filtration, flocculation, chemical disinfection, heat, or UV radiation) |
| | Other components of intervention (hygiene message, improved supply, improved sanitation, improved storage) |
| | Whether water protected to POU (i.e. by pipe, residual disinfection, or safe storage) |
| | Case definition of diarrhoea |
| | Method for diarrhoea assessment (self-reported, observed, or clinically confirmed) |
| | Where self reported, recall period used |
| | Study duration; Adherence rates |
| | Publication status |
| | Prescribed criteria of methodological quality |
| Individual characteristics | Age group |

^bSearch terms used in combination with the search strategy for retrieving trials developed by Cochrane (Higgins 2005); upper case: MeSH or EMTREE heading; lower case: free text term.

(Continued)

| | Type and description of water source | | | | |
|----------|--|--|--|--|--|
| | Level of faecal contamination of control water (low (< 100 thermotolerant coliforms (TTC)/ 100 mL), medium (100 to 1000 TTC/ 100 mL), and high (> 1000 TTC/ 100 mL) | | | | |
| | Causative agents identified (yes or no) | | | | |
| | Water collection, storage, and drawing practices | | | | |
| | Distance to and other constraints regarding water supply | | | | |
| | Sanitation facilities (improved or unimproved) | | | | |
| | Hygiene practices | | | | |
| Outcomes | Pre- and post-intervention faecal contamination of drinking water, and method of assessment (including indicator used) | | | | |
| | Diarrhoea morbidity and 95% CI for each age group reported | | | | |
| | Manner of measuring diarrhoea morbidity | | | | |
| | Mortality attributed to diarrhoea | | | | |
| | Rate of utilization of intervention and manner of assessing same | | | | |

Abbreviations: POU: point of use; CI: confidence interval; UV: ultraviolet.

WHAT'S NEW

| Date | Event | Description |
|-----------------|---------|------------------------------|
| 21 October 2015 | Amended | Amended author affiliations. |

HISTORY

| Date | Event | Description |
|-----------------|--|--|
| 15 October 2015 | New search has been performed | The review authors updated the review, and included several new studies, a 'Summary of findings' table, and 'Risk of bias' assessments |
| 15 October 2015 | New citation required and conclusions have changed | The review authors performed an updated literature search, reapplied the inclusion criteria, repeated data extraction, added new studies, and used the GRADE approach to assess the quality of the evidence. They also applied statistical methods to unify the measures of effect and applied additional criteria for subgrouping based on study design, setting, and length of follow-up |

CONTRIBUTIONS OF AUTHORS

TC and SC conceived the review. TC coordinated the review. TC, KA, SB, RP, HC, and SC designed the review. TC and authors of the initial review drafted the protocol. SB and Cochrane Infectious Diseases Group (CIDG) performed the search strategy. SB and RP screened search results. KA, SB, and RP retrieved papers. SB and RP applied inclusion criteria. KA, SB, and RP extracted data. KA, SB, RP, HC, and FM computed estimates of effect. KA, TC, FM, and DS applied quality criteria. KA contacted study authors for additional information. TC, KA, HC, DS, and CIDG addressed statistical issues. KA entered data into Review Manager (RevMan). TC, KA, and DS drafted the review. SB, RP, HC, and SC commented on the review. TC, KA, HC, FM, and DS prepared tables. KA prepared figures. TC is guarantor of this Cochrane Review.

DECLARATIONS OF INTEREST

TC, KA, SB, and SC have provided research or consulting services for Unilever, Ltd., Medentech, Ltd., DelAgua Health and Science, Ltd., and Vestergaard-Frandsen SA who manufacture or sell household-based water treatment devices.

SOURCES OF SUPPORT

Internal sources

• Liverpool School of Tropical Medicine, UK.

External sources

• Department for International Development (DFID), UK.

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

Risk of bias has been assessed using GRADE rather than the original methods expressed in the protocol. Statistical methods have been used to pool odds ratios, rate ratios, RRs and longitudinal prevalence ratios. Subgrouping has been done separately for each water quality intervention, and additional subgrouping has been conducted based on study design and length of follow up. Data has been provided on adjustment of studies for non-blinding.

INDEX TERMS

Medical Subject Headings (MeSH)

Controlled Before-After Studies; Diarrhea [*prevention & control]; Drinking Water [standards]; Randomized Controlled Trials as Topic; Water Purification [*methods; standards]; Water Supply [*standards]

MeSH check words

Adult; Child; Child, Preschool; Humans; Infant