ABSTRACT

Background: Zinc deficiency is prevalent in children in developing countries. Supplemental zinc provides therapeutic benefits in diarrhea.

Objective: We sought to measure the effect of supplemental zinc given with oral rehydration therapy during recovery from acute or persistent diarrhea.

Design: We conducted pooled analyses including all available published and unpublished randomized controlled trials of the effects of supplementary oral zinc in children aged <5 y with acute or persistent diarrhea. We used Cox survival regression analysis to evaluate the overall effect of zinc on continuation of diarrhea and possible differential effects in subgroups divided by sex, age, weight-for-height, and initial plasma zinc concentration. Dichotomous outcomes were analyzed by logistic regression. To assess the effects of excluding studies without original data from the pooled analyses, effect-size was estimated for all studies by using random-effects models.

Results: Zinc-supplemented children had a 15% lower probability of continuing diarrhea on a given day (95% CI: 5%, 24%) in the acute-diarrhea trials and a 24% lower probability of continuing diarrhea (95% CI: 9%, 37%) and a 42% lower rate of treatment failure or death (95% CI: 10%, 63%) in the persistent-diarrhea trials. In none of the subgroup analyses were the 2 subgroups of each pair significantly different from each other; however, in persistent diarrhea there tended to be a greater effect in subjects aged <12 mo, who were male, or who had wasting or lower baseline plasma zinc concentrations.


KEY WORDS Diarrhea, diarrheal disease, malnutrition, meta-analysis, randomized controlled trial, zinc, children, infants, developing countries, zinc supplementation, nutrition, zinc deficiency

INTRODUCTION

It is estimated that diarrheal diseases cause >3 million deaths of children in developing countries each year and contribute substantially to malnutrition in surviving children (1). Diarrheal episodes of longer duration, commonly called persistent diarrhea, have the greatest effect on these outcomes (2, 3). Treatment of acute diarrhea with oral rehydration solution has become widespread, resulting in reduced mortality from dehydrating diarrheas but no decrease in the duration of episodes or their consequences, such as malnutrition (4). Furthermore, adherence to recommendations regarding fluid therapy in children with diarrhea is poor because caregivers want to reduce the duration of illness and this often leads them to use antibiotics and other treatments of no proven value (5).

Two well-documented determinants of diarrheal duration are low weight-for-age and decreased cell-mediated immunity (6, 7). A common determinant of both of these factors is zinc deficiency (8, 9), thought to be prevalent in children in developing countries (10). Furthermore, zinc supplementation was shown to reduce the duration and severity of childhood diarrhea in randomized controlled trials (11–20).

1 From the Aga Khan University Medical Centre, Karachi, Pakistan; the Medical Research Council Biostatistics Unit, Cambridge, United Kingdom; the Johns Hopkins School of Public Health, Baltimore; the University of California, Davis; the University of the West Indies, Mona, Jamaica; Trisakti University, Jakarta, Indonesia; Dhaka Medical College, Dhaka, Bangladesh; Emory University, Atlanta; the National Institute of Nutrition, Hanoi, Vietnam; the Nutrition Research Institute, Lima, Peru; the National Institute of Nutrition, Mexico City; the International Centre for Diarrhoeal Disease Research, Dhaka, Bangladesh; the Institute of Nutrition of Central America and Panama, Guatemala City; and the All India Institute of Medical Sciences, New Delhi.

2 Supported by the Johns Hopkins Family Health and Child Survival Cooperative Agreement with the US Agency for International Development and the World Health Organization, Division of Child Health and Development, which provided support for the pooled analysis. REB drafted the manuscript while in residence at the Rockefeller Foundation Bellagio Study and Conference Center.

3 Address reprint requests to RE Black, Department of International Health, The Johns Hopkins School of Public Health, 615 North Wolfe Street, Baltimore, MD 21205. E-mail: rblack@jhsph.edu.

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We conducted pooled analyses of available randomized controlled trials that evaluated the effects of supplementary oral zinc given as an adjunct to other therapy in children with acute or persistent diarrhea. We also attempted to identify any differential effects in subgroups of children. Finding these differential effects might allow for targeting of this therapy toward children who would derive the greatest benefit. For the pooled analyses, we formed a group of investigators who had conducted trials of zinc supplementation. To reduce possible publication bias, we attempted to include all investigators of published and unpublished trials. The use of original trial data from individual study children allowed standardization of outcome and subgroup definitions, use of information not included in the publications, and use of proper statistical methods. We present here the pooled analyses of a total of 7 trials of zinc supplementation; there were 3 trials in children with acute diarrhea and 4 trials in children with persistent diarrhea. We also performed a meta-analysis of the effect of zinc on diarrheal duration; we included all known trials to ensure that our results were not biased because we excluded from the pooled analyses 3 trials for which original data were not available.

SUBJECTS AND METHODS

We attempted to locate all published and unpublished randomized controlled trials of oral zinc supplementation in preschool children in developing countries. We systematically searched MEDLINE, SCI-SCIMATE, CURRENT CONTENTS, Cochrane Clinical Trials Register, and references from articles. Potential funding agencies were also contacted to identify trials. International agencies such as the World Health Organization (WHO) and UNICEF were contacted. The principal investigators of known trials and researchers in the micronutrient field were asked to identify trials.

Studies eligible for inclusion were randomized, controlled, and masked trials that assessed the adjunctive therapeutic benefit of zinc supplements containing ≥50% of the US recommended dietary allowance (RDA) per day in children aged <5 y. The children had acute (<14 d pre-enrollment duration) or persistent (≥14 d pre-enrollment duration) diarrhea and resided in a developing country. Subgroups for the analyses were defined a priori as follows: sex, age (<12 mo or ≥12 mo), and weight-for-height z score (<−2 or ≥−2 compared with the National Center for Health Statistics reference) (21). Subgroups for baseline plasma zinc concentration were defined as those subjects below or above the median plasma zinc concentration for that study. The subgroups were selected because some original trial reports suggested differential effects by sex, age, nutritional status, or baseline plasma zinc concentrations.

From the search, 26 zinc supplementation trials were identified and 10 of these were therapeutic trials that met the inclusion criteria (11–20). Of the 5 trials of zinc therapy for acute diarrhea, only 3 were incorporated into the pooled analysis because original, individual case data were no longer available for one trial (11) and the investigator did not provide data for another (16). Of the 5 trials on persistent diarrhea, 1 could not be included because original data were not available (12). The published data from the 3 trials that could not be included in the pooled analysis were incorporated in other analyses of effect size for all known trials. All trials used standard fluid and dietary case management of diarrhea as recommended by the WHO (4).

Principal investigators of the trials included in the pooled analysis agreed to join the Zinc Investigators’ Collaborative (ZINC) Group, which also included 2 external advisors (SMB and RM) selected by the WHO Programme on Child Health and Development and 1 advisor selected by the coordinators of the ZINC Group (KHB). An initial meeting with some of the investigators and correspondence with the others resulted in consensus on trial inclusion criteria, subgroup and outcome definitions, and procedures for the pooled analyses. After a preliminary analysis, the ZINC Group met to consider conclusions and implications and to evaluate the need for additional analyses. Final analyses and draft manuscripts were approved by all investigators who had trials included and by the advisors.

For the acute-diarrhea trials, diarrhea was defined as 3 or 4 loose stools (depending on the original trial definition) in a 24-h period. The final day of diarrhea was defined as the last day meeting the above definition followed by 48 h without diarrhea. For the persistent-diarrhea trials, the definitions of diarrhea and recovery used in the original trials were retained, as was the definition of treatment failure (generally an increase in diarrheal severity, occurrence of dehydration, or continued diarrhea for >7 d or >14 d). These trials differed in their withdrawal criteria. In the pooled analysis of recovery after enrollment, data from children who withdrew from the study, who were declared a treatment failure, or who died were included up to the time that they dropped out of the study. Children who had no diarrhea after enrollment in the study or who had an unknown pre-enrollment duration were excluded from the analyses of diarrheal recovery. In the acute-diarrhea analysis of 2446 children, 1 did not have a pre-enrollment duration and 30 had no diarrhea after enrollment. In the persistent-diarrhea analysis of 640 children, 3 did not have a pre-enrollment duration and 84 had no diarrhea after enrollment.

Individual subject data were provided from each trial with outcomes redefined as necessary, along with descriptive information on the trial methods and study populations. For the Indonesian trial that included >1 diarrhea episode per child, only the first episode for each child was used.

A detailed methodologic assessment and scoring system (available on request) for assessment of study quality was used. This system incorporated standard randomized controlled trial quality assessments and items specific to these trials, such as definitions of recovery and degree of co-intervention. Trials were independently evaluated by REB and SS and any disagreements were resolved by further review of the methods and consensus. At a meeting of the ZINC Group, the methodologic scores were reviewed by the principal investigators of the trials and any errors and inconsistencies were corrected.

For the pooled analysis of trials with available individual child data, the therapeutic effect of zinc on the duration of diarrhea was analyzed by using Cox survival regression models stratified by individual trial (22, 23) using the exact method for handling ties (23, 24). In the simplest of these, continuation of the episode after enrollment was modeled as the dependent variable and treatment group and pre-enrollment duration were independent variables. Additional analyses were performed with nutritional status, sex, and age added as covariates. The analyses were performed with SAS (version 8.0; SAS Institute, Cary, NC). These models permitted calculation of the relative hazard (RH) for continuation of the episode and its 95% CI. The control group was coded as 1 and the zinc group as 0, resulting in an RH of <1 for a beneficial effect, consistent with the beneficial effect
expressed by the odds ratio (OR). The dichotomous dependent outcomes, which were duration ≤ 7 d compared with > 7 d postenrollment in acute-diarrhea trials and treatment failure or death in persistent-diarrhea trials, were analyzed by using logistic regression models. These models were also stratified by trial, with treatment group, subgroup categories, and potential interaction terms as independent variables, to calculate the ORs and 95% CIs (25). To evaluate a possible clustering effect in the survival analysis, we used random-effects extensions of the Cox models (26); the robust correlation matrix and SEs were calculated with use of the PHREG procedure in SAS version 8.0 (27).

To estimate summary effects, including data from the 3 eligible trials for which original data were not available, we performed a meta-analysis of effect size. Means and SDs of diarrheal duration in the zinc and control groups were used to estimate the effect size. By using Bayesian methods, we estimated a joint posterior probability distribution for the variable of interest (28). From these joint distributions, effect size and 95% CI were calculated for each study. By using a random-effects hierarchical model (28), the summary effect size and the 95% CI were estimated. The confidence profile method was also used to estimate the ORs and CIs for effects on episodes lasting > 7 d in acute-diarrhea trials and effects on treatment failure or death in the persistent-diarrhea trials; summary estimates were determined by using random-effects hierarchical models (28). Of the 3 trials not included in the pooled analysis, only 1 (16) reported data on these outcomes and so was incorporated into the analysis. Finally, to evaluate heterogeneity across studies, we determined the chi-square for heterogeneity.

RESULTS

The 3 acute-diarrhea trials, conducted in Indonesia (18), India (13), and Bangladesh (14), were similar in terms of the age of the children and the dose of zinc (Table 1). The study in Bangladesh enrolled only children who were underweight. In part because of this, the nutritional status of children in the 3 trials differed (Table 2). In the trial for which plasma zinc concentrations before and after treatment were available, the mean plasma zinc concentration increased in the zinc group after treatment but did not increase in the control group; this difference between the zinc and control groups was not statistically significant.

In the acute-diarrhea trials (Table 3), zinc-supplemented children had a 15% lower probability of continuing diarrhea on a given day (95% CI: 8%, 22%) than did the children in the control group. The random-effects extension of survival analysis yielded similar estimates (Table 3).

In the effect-size analysis, which used data from all 5 acute-diarrhea trials, mean duration of diarrhea was lower in the zinc-supplemented group in all 5 studies, significantly so in 2 studies (Table 4). The effect size for reduction in the mean duration of the diarrheal episode in individual trials ranged from 10% to 24%. The summary estimate of the effect size for reduction in duration was 16% (95% CI: 7%, 26%). The results of these 5 trials were not significantly heterogenous (chi square = 2.99, P = 0.56). For the analysis of the effect on episodes lasting ≥ 7 d, zinc-supplemented children had a 27% lower rate of prolonged episodes than did control children (OR = 0.73; 95% CI: 0.55, 0.98). These results were also not significantly heterogenous (chi square = 1.39, P = 0.71).

The 4 persistent-diarrhea trials in Peru (19), Bangladesh (15, 20), and Pakistan (17) were similar to each other and to the acute-diarrhea trials in terms of age group and dose of zinc (Table 1). One of the trials in Bangladesh (20) and the one in Pakistan had nutritional-status criteria for enrollment. The background characteristics of children in these 4 trials differed: the children in the Peru study were substantially better nourished than were children in the other studies in terms of weight, but not height (Table 2). This study also differed in that it enrolled children in the community rather than in a health facility. In 3 of the 4 trials, the plasma zinc concentrations showed a significant
response to zinc supplementation but did not change significantly in the control group (Table 2).

In a stratified analysis of the persistent-diarrhea trials (Table 3), zinc-supplemented children had a 24% lower probability of continuation of diarrhea on a given day (95% CI: 8%, 38%) than did control children. The random-effects survival analysis yielded similar results (Table 3). Zinc-supplemented children in these trials had a 42% lower rate of treatment failure or death (OR = 0.58; 95% CI: 0.37, 0.40) than did control children. For this outcome, the trials were significantly heterogeneous (chi-square = 8.6, \(P = 0.04\)), which was due mainly to the results of the study conducted in Pakistan. After exclusion of this study, there was no significant heterogeneity (chi-square = 3.2, \(P = 0.20\)). Random-effects analysis for this outcome yielded an OR of 0.61 (95% CI: 0.26, 1.46).

In the effect-size analysis, which used data from all 5 persistent-diarrhea trials including 1 trial (12) not included in pooled analysis, the mean duration of diarrhea tended to be lower in the zinc-supplemented group in all 5 studies, although this result was significant in only 1 study. The effect size for reduction in the mean duration of the diarrheal episode in individual trials ranged from 12% to 53%. The summary estimate of the effect size for reduction in duration was 29% (95% CI: 6%, 53%). The results of these 5 trials were not significantly heterogeneous (chi-square = 2.26, \(P = 0.69\)).

In the subgroup analyses in acute-diarrhea trials, the effect of zinc supplementation in each of the subgroups by age, wasting, and sex was significant (Figure 1). The subgroups did not differ from each other in terms of the magnitude of this effect. In persistent diarrhea, age <12 mo, wasting, and male sex were

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**TABLE 2**

Background characteristics and plasma zinc concentrations before and after supplementation in therapeutic zinc trials

<table>
<thead>
<tr>
<th>Trial</th>
<th>Percentage with illiterate mothers</th>
<th>Percentage with height-for-age z score &lt; -2</th>
<th>Percentage with weight-for-height z score &lt; -2</th>
<th>Plasma zinc concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>%</td>
<td>%</td>
<td>%</td>
<td>µmol/L</td>
</tr>
<tr>
<td><strong>Acute diarrhea</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Indonesia (18)</td>
<td>32</td>
<td>23</td>
<td>15</td>
<td>NA1</td>
</tr>
<tr>
<td>India (13)</td>
<td>81</td>
<td>67</td>
<td>21</td>
<td>10.0 ± 0.11</td>
</tr>
<tr>
<td>Bangladesh (14)</td>
<td>57</td>
<td>58</td>
<td>21.6 ± 0.8</td>
<td></td>
</tr>
<tr>
<td><strong>Persistent diarrhea</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Peru (19)</td>
<td>5</td>
<td>30</td>
<td>1</td>
<td>11.4 ± 0.3</td>
</tr>
<tr>
<td>Bangladesh (15)</td>
<td>40</td>
<td>34</td>
<td>42</td>
<td>13.4 ± 0.5</td>
</tr>
<tr>
<td>Bangladesh (20)</td>
<td>51</td>
<td>52</td>
<td>46</td>
<td>14.0 ± 1.0</td>
</tr>
<tr>
<td>Pakistan (17)</td>
<td>83</td>
<td>87</td>
<td>35</td>
<td>11.9 ± 0.8</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
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<td></td>
</tr>
</tbody>
</table>

1 NA, not available.

**TABLE 3**

Pooled analysis of the therapeutic effect of zinc supplementation on acute and persistent diarrhea

<table>
<thead>
<tr>
<th>Trial</th>
<th>Effect on recovery</th>
<th></th>
<th></th>
<th>Effect on continuation for &gt;7 d, failure, or death1</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>Recovered</td>
<td>Censored2</td>
<td>Excluded3</td>
<td>Relative hazard (95% CI)</td>
</tr>
<tr>
<td></td>
<td>n</td>
<td>n</td>
<td>n</td>
<td></td>
</tr>
<tr>
<td></td>
<td>n (%)</td>
<td></td>
<td></td>
<td>Zinc Control</td>
</tr>
<tr>
<td>Acute diarrhea</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Indonesia (18)</td>
<td>1368</td>
<td>0</td>
<td>30</td>
<td>0.92 (0.83, 1.02)</td>
</tr>
<tr>
<td>India (13)</td>
<td>931</td>
<td>6</td>
<td>0</td>
<td>0.79 (0.69, 0.90)</td>
</tr>
<tr>
<td>Bangladesh (14)</td>
<td>101</td>
<td>9</td>
<td>1</td>
<td>0.85 (0.57, 1.28)</td>
</tr>
<tr>
<td>Pooled</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>0.85 (0.78, 0.92)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pooled multifactorial</td>
<td>0.85 (0.78, 0.92)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pooled random effect</td>
<td>0.85 (0.76, 0.95)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Persistent diarrhea</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Peru (19)</td>
<td>164</td>
<td>24</td>
<td>87</td>
<td>0.82 (0.60, 1.12)</td>
</tr>
<tr>
<td>Bangladesh (15)</td>
<td>138</td>
<td>52</td>
<td>0</td>
<td>0.85 (0.61, 1.19)</td>
</tr>
<tr>
<td>Bangladesh (20)</td>
<td>55</td>
<td>32</td>
<td>1</td>
<td>0.45 (0.26, 0.78)</td>
</tr>
<tr>
<td>Pakistan (17)</td>
<td>55</td>
<td>32</td>
<td>0</td>
<td>0.98 (0.57, 1.67)</td>
</tr>
<tr>
<td>Pooled</td>
<td>0.76 (0.62, 0.92)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pooled multifactorial</td>
<td>0.75 (0.62, 0.91)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pooled random effect</td>
<td>0.76 (0.63, 0.91)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

1 Effect on continuation for >7 d was used for acute-diarrhea trials; effect on treatment failure or death was used for persistent-diarrhea trials.

2 Includes withdrawal, treatment failure, and death.

3 Exclusions were because the postenrollment duration of diarrhea was zero, except for 4 cases for which pre-enrollment duration was missing.


**DISCUSSION**

These pooled analyses of data from trials of acute and persistent diarrhea in developing countries show that zinc, given in a daily dose of about twice the RDA, significantly reduces the duration of acute or persistent diarrhea. Three trials (11, 12, 16) that met the inclusion criteria could not be used in the pooled analysis but were included in meta-analyses of all eligible trials (Table 4). Note that one of the trials conducted in Bangladesh used a factorial design with zinc and vitamin A; the zinc effect on recovery from diarrhea was significant but there was no effect of vitamin A (16).

It is important to examine the effect of zinc supplementation on other measures of severity, such as diarrheal stool output, occurrence of dehydration, treatment failure, or death. The trials provided some information on these outcomes, but the different types of study data available precluded many pooled analyses. Three acute-diarrhea trials with appropriate outcome measures all found reductions in diarrheal severity in zinc-supplemented children compared with control children. Of the 2 trials conducted in India, one found 18% fewer diarrheal stools/d ($P < 0.1$) (11) and the other found 39% fewer watery stools/d ($P < 0.02$) (13). In the only hospital-based trial of acute diarrhea, zinc-supplemented children had a 28% lower measured diarrheal stool output/d ($P = 0.06$) (14). Of the 4 persistent-diarrhea trials that included severity measures, 2 found no significant difference between the groups (15, 19). The trial conducted in India reported a 21% lower diarrheal stool frequency ($P = 0.08$) (12) and a hospital-based trial in Bangladesh found a 37% lower measured stool output ($P < 0.02$) in zinc-supplemented children (20). Although there were too few

![Figure 1](image_url)

**FIGURE 1.** Therapeutic effect of zinc supplementation assessed by the relative hazard of continuation of diarrhea in subgroups of children with acute or persistent diarrhea (pooled analysis).
concentrations in different settings may have reflected method-
differences among the trials, we thought this approach was more appropriate than selecting a single plasma zinc concentra-
tion for use in all the trials. Furthermore, plasma zinc can be reduced by illness; therefore, some of the variability could reflect the severity of diarrhea or concomitant infections (33). In acute diarrhea, both of the subgroups with lower and higher baseline zinc concentrations showed a significant effect of supplemental zinc, but the effect tended to be larger in the group with lower baseline zinc concentrations. In children with persistent diarrhea, the group with lower baseline zinc concentrations showed a significant benefit of zinc supplementation and the group with higher baseline zinc did not, although the pooled effect estimate did suggest benefit. These effects suggest that zinc should be provided to all children with acute and persistent diarrhea in such settings.

The mechanisms of these effects of zinc on diarrhea are unclear. Zinc deficiency is associated with many immunologic deficits, and zinc supplementation was shown to improve immune function in children in developing countries (9, 34) and to reduce the incidence and prevalence of diarrhea (35). Other possible mechanisms include effects of zinc deficiency on intestinal permeability (36, 37), regulation of intestinal water and electrolyte transport (38), brush border enzymatic function (39, 40), and intestinal epithelial tissue repair (41, 42).

The use of zinc as adjunctive therapy has the potential to improve the management of diarrhea and increase survival in children, if it can be incorporated into diarrheal disease control programs in developing countries. Primary prevention of zinc deficiency would be expected to reduce infectious disease morbidity and improve the growth and development of children (8, 35, 43–45) and might also reduce the severity of diarrhea. Attention should now focus on the best means of providing zinc during diarrhea or on other ways to improve the zinc nutriture of children in developing countries.

Robert Black and Sunil Sazawal organized and conducted the pooled analy-
ses and drafted the manuscript. Zulfiquar Bhutta, Adi Hidayat, Farida Khatun, Mary Penny, Swapan Roy, and Sunil Sazawal were principal investigators of zinc therapeutic trials. Julie Meeks Gardner, Nguyen Ninh, Jorge Rosado, Marie Ruel, and Anu Shankar were principal investigators of zinc preventive trials and participated in meetings of the ZINC Group. Kenneth Brown, Sheila M Bird, and Reynaldo Martorell served as advisors. Members of the ZINC Group are listed in alphabetical order. Other investigators in the therapeutic trials in the pooled analyses were A Achadi, S Soedarmo, and Sunoto in Indonesia; MK Bhan, N Bhandari, RE Black, S Jalla, and A Sinha in India; S Akramuzzaman, R Behrens, G Fuchs, R Haider, D Mahulababis, M Abdul Malek, NR Sarkar, and A Tomkins in Bangladesh; RE Black, KH Brown, A Duran, CF Lanata, B Lönnrold, RM Marin, and JM Peerson in Peru; and Z Issani, S Nizzi, and S Nizami in Pakistan.

REFERENCES


### TABLE 5

<table>
<thead>
<tr>
<th>Trial</th>
<th>Baseline plasma zinc below median</th>
<th>Baseline plasma zinc above median</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acute diarrhea</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bangladesh (14)</td>
<td>0.90 (0.52, 1.59)</td>
<td>0.89 (0.57, 1.39)</td>
</tr>
<tr>
<td>Pooled</td>
<td>0.75 (0.63, 0.90)</td>
<td>0.83 (0.69, 0.99)</td>
</tr>
<tr>
<td>Persistent diarrhea</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bangladesh (15)</td>
<td>0.69 (0.43, 1.11)</td>
<td>1.01 (0.63, 1.61)</td>
</tr>
<tr>
<td>Bangladesh (20)</td>
<td>0.46 (0.24, 0.91)</td>
<td>0.43 (0.17, 1.11)</td>
</tr>
<tr>
<td>Pakistan (17)</td>
<td>0.82 (0.40, 1.70)</td>
<td>1.12 (0.48, 2.70)</td>
</tr>
<tr>
<td>Pooled</td>
<td>0.70 (0.53, 0.91)</td>
<td>0.88 (0.67, 1.17)</td>
</tr>
</tbody>
</table>

1 Relative hazard for continuation of diarrhea (95% CI).


