

Small-quantity lipid-based nutrient supplements, together with malaria and diarrhea treatment, improve growth and development in young Burkinabe children

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Background

Preventive zinc supplementation reduces the incidence of diarrhea and increases growth¹⁻³

The impact of zinc is less certain when provided as fortified food or home fortification products⁴

The optimal dose of zinc in lipid-based nutrient supplements (SQ-LNS) remains to be determined

Objectives

To assess the appropriate dose of zinc in SQ-LNS by comparing biochemical and zinc-related functional responses among young children who receive different zinc doses in SQ-LNS or tablets

To compare the same outcomes among children who do or do not receive SQ-LNS and selected health services

Methods

- Community-based, partially double-blind, placebo-controlled, randomized clinical trial in rural Burkina Faso
- Inclusion criteria: 9 mo of age, parental consent
- Exclusion criteria: Hemoglobin (Hb) <50 g/L, weight/length <70% NCHS median, edema, chronic or congenital diseases, history of peanut allergy or anaphylactic reaction
- Cluster randomization of 34 communities to intervention (IC) or non-intervention (NIC) cohort
- Eligible children in IC were randomly assigned by concession to 1 of 4 intervention groups (Figure 1)
- Hemoglobin and rapid diagnostic test for malaria (RDT) at 9 mo
- Length and weight at 9 and 18 mo in both cohorts
- Plasma zinc at 9 and 18 mo in randomly selected sub-group in both cohorts
- Neurobehavioral development at 18 mo in a sub-group (4 grps only)

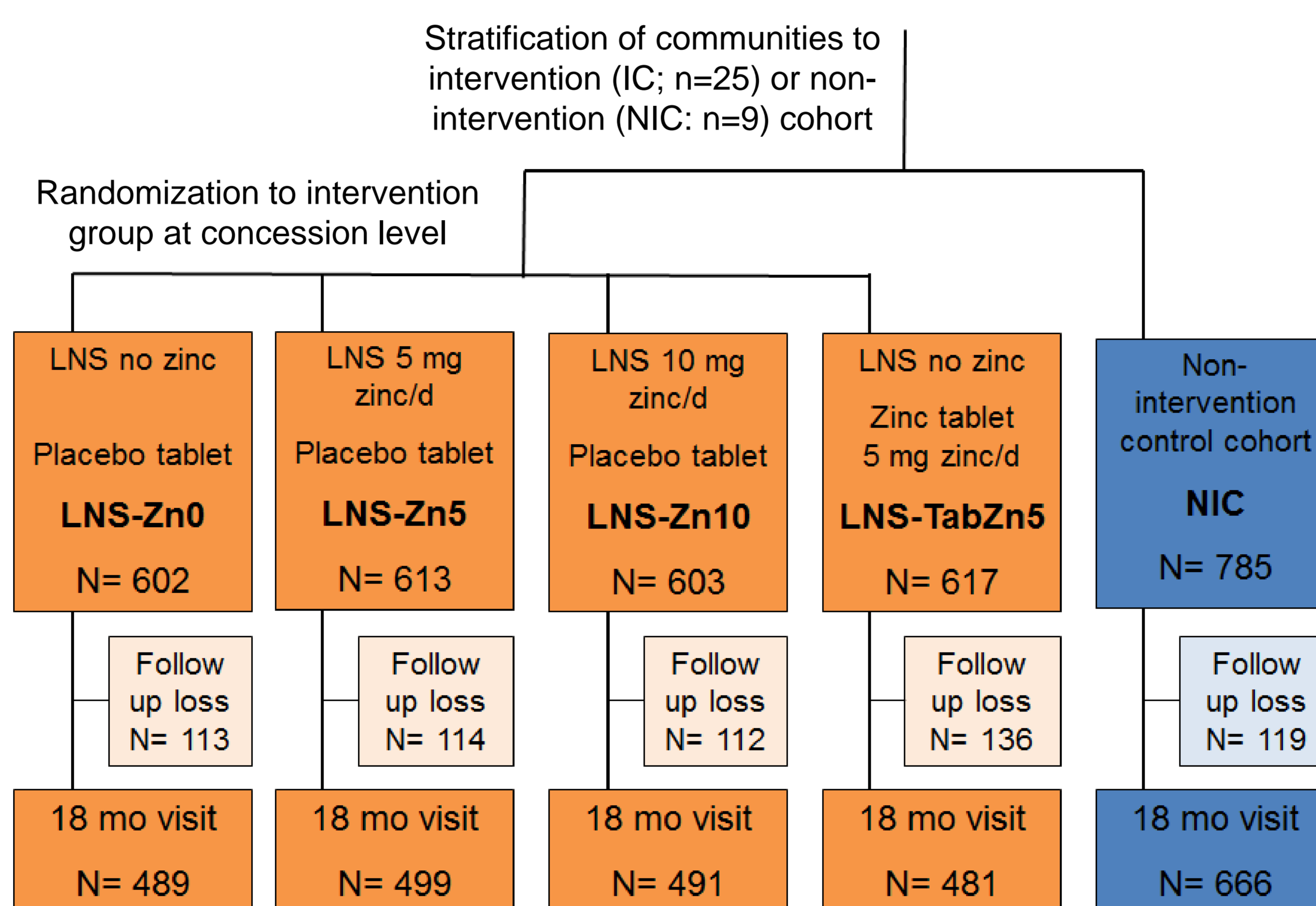
In intervention groups only:

- Weekly home visits for morbidity surveillance and distribution of LNS and tablets
- Free treatment of reported diarrhea, fever and confirmed malaria

References

- Brown et al, Food Nutr Bull 2009
- Imdad et al, BMC Public Health, 2011
- Yakoob et al, BMC Public Health, 2011
- Das JK, et al. Systematic Reviews 2013

Results Figure 1: Flow diagram for the study participants



Results

Table 2: Hemoglobin and plasma zinc

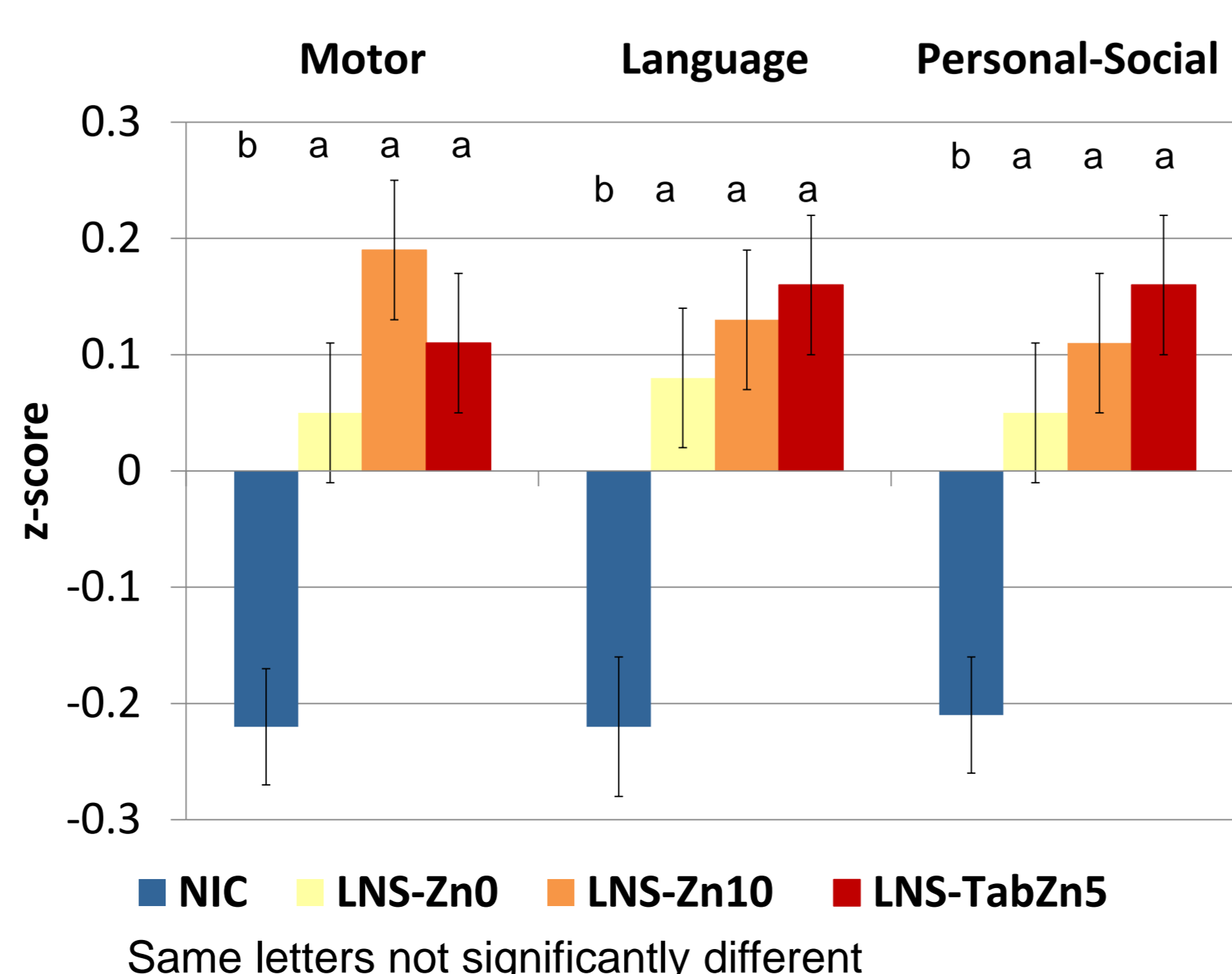
	IC	NIC	P-value
Hemoglobin concentration (g/L) ¹			
N	2435	785	
At 9 mo	89 ±15	88 ±16	0.417
at 18 mo	97 ±15	88 ±16	<0.0001
Adjusted plasma zinc concentration (µg/dL) ^{1,2}			
N	310	93	
at 9 mo	69.6 ±1.2	67.0 ±1.2	0.090
at 18 mo	64.6 ±1.2	64.1 ±1.2	0.659

¹ Adjusted for baseline value and cluster randomization
² Also adjusted for presence of inflammation (CPR, AGP)

At 18 mo, change in Hb was greater in IC than NIC (+8 vs -1 g/L, p<0.0001), but 79% of IC were still anemic (vs. 91% in NIC).

Results

Figure 2: Neurobehavioral development score in randomly selected sub-sample at 18 mo



Results

Table 1: Baseline characteristics

N	3220
Age (mo)	9.4 ± 0.4
Boys (%)	50.3
LAZ	-1.21 ± 1.10
WAZ*	-1.42 ± 1.15
WLZ*	-0.99 ± 1.05
Hemoglobin (g/L)	89 ± 15
Malaria RDT (% positive)	61.4
Low plasma zinc (<65 µg/dL; %)	35.2

LAZ, length-for-age Z-score; WAZ, weight-for-age Z-score; WLZ, weight-for-length Z-score, RDT, rapid diagnostic test Results shown as mean ± SD or percentage (%)

* Significantly different by group

Results

Table 3: Length, weight and stunting and wasting prevalence at 18 mo

	LNS-Zn0	LNS-Zn5	LNS-Zn10	LNS-TabZn5	NIC	P-value
Length 18 mo (cm)	77.7 ^a ±3.0	77.7 ^a ±3.0	77.7 ^a ±3.1	77.7 ^a ±2.9	76.9 ^b ±3.4	P _I < 0.0001 P _G =0.974
Stunting (%)	29.9 ^a	30.1 ^a	32.8 ^a	24.4 ^a	39.3 ^b	P _I <0.0001 P _G =0.536
Weight 18 mo (kg)	9.29 ^a ±1.00	9.30 ^a ±1.09	9.29 ^a ±1.11	9.32 ^a ±1.05	9.02 ^b ±1.18	P _I < 0.0001 P _G =0.907
Wasting (%)	7.2 ^b	8.9 ^{ab}	13.2 ^a	5.4 ^b	13.5 ^a	P _I =0.0003 P _G =0.003

Same letters not significantly different
 Adjusted for cluster randomization (village, concession), baseline value, age and potential co-variables, when applicable.

- P_I: for comparison between **intervention** and **non-intervention** cohorts
- P_G: for comparison among **4 intervention groups**

Conclusion

SQ-LNS along with malaria and diarrhea treatment reduced stunting prevalence from 39% to 29%, decreased wasting and anemia prevalence and resulted in higher motor, language, and personal-social development scores compared with the non-intervention cohort.

Lack of impact on plasma zinc concentration suggests inadequate adherence to zinc tablet and/or inadequate absorption.

Unable to answer original research question on optimal dose of zinc because biochemical and functional responses to zinc were not detected.

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