9.3 Composition of Parenteral Nutrition: Zinc (either alone or in combination with other antioxidants) March 2013

There are no new randomized controlled trials since the 2009 update and hence there are no changes to the following summary of evidence.

Recommendation: There are insufficient data to make a recommendation regarding IV/PN zinc supplementation in critically ill patients.

Discussion: The committee noted the potentially large treatment effect of zinc enriched PN with respect to a reduction in mortality. The wide confidence intervals weaken this estimate. Safety, cost and feasibility issues were considered to be favourable. The committee noted that in some sub populations of critical illness with high zinc losses (GI fistula, burns, etc) there may be some benefit to zinc supplementation but data are lacking to support a recommendation.

Values	Definition	2009 Score (0.1.2.3)	2013 Score (0.1.2.3)
Effect size	Magnitude of the absolute risk reduction attributable to the intervention listeda higher score indicates a larger effect size	3 (mortality)	3
Confidence interval	95% confidence interval around the point estimate of the absolute risk reduction, or the pooled estimate (if more than one trial)a higher score indicates a smaller confidence interval	2*	0*
Validity	Refers to internal validity of the study (or studies) as measured by the presence of concealed randomization, blinded outcome adjudication, an intention to treat analysis, and an explicit definition of outcomesa higher score indicates presence of more of these features in the trials appraised	3*	2*
Homogeneity or Reproducibility	Similar direction of findings among trialsa higher score indicates greater similarity of direction of findings among trials	1	1
Adequacy of control group	Extent to which the control group presented standard of care (large dissimilarities=1, minor dissimilarities=2, usual care=3)	3	3
Biological Plausibility	Consistent with understanding of mechanistic and previous clinical work (large inconsistencies=1, minimal consistencies=2, very consistent=3)	2	2
Generalizability	Likelihood of trial findings being replicated in other settings (low likelihood i.e. single centre=1, moderate likelihood i.e. multicentre with limited patient population or practice setting=2, high likelihood i.e. multicentre, heterogenous patients, diverse practice settings=3)	1	1
Low cost	Estimated cost of implementing the intervention listeda higher score indicates a lower cost to implement the intervention in an average ICU	3	3
Feasible	Ease of implementing the intervention listeda higher score indicates greater ease of implementing the intervention in an average ICU	3	3
Safety	Estimated probability of avoiding any significant harm that may be associated with the intervention listeda higher score indicates a lower probability of harm	3	3

Semi Quantitative Scoring

*The 2009 scoring for confidence intervals and validity were corrected in December 2012

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Question: Does zinc supplementation (via IV/PN) given either alone or in combination with other nutrients result in improved outcomes in the critically ill patient?

Summary of evidence: There were 4 level 2 studies reviewed, one that compared a higher dose of parenteral zinc to a lower dose in ventilated head injured patients (Porter), both groups progressing to oral zinc (higher vs. lower). The other three studies compared IV zinc in combination with other antioxidants (selenium, α tocopherol and/or copper) to placebo.

Mortality: When all three studies were aggregated, zinc supplementation was associated with a trend a reduction in mortality (RR 0.58, 95% CI 0.23, 1.44, p=0.24; figure 1).

Infections: Only reported in two studies, one reported number of infections per patient (Young), hence unable to do a meta-analysis. The other study reported no differences in infectious complications between the two groups (Berger 2001).

Hospital/ICU length of stay, ventilator days: There were no statistical differences between the groups (figures 3 and 4).

Cost, other complications: Only one study reported the number of patients with organ failure, which was the same in the group receiving zinc supplementation and none (Berger 2001)

Conclusion:

Zinc supplementation given IV/PN (either alone or in combination with other antioxidants) may be associated with a trend towards a reduction in mortality in critically ill patients.

Level 1 study: if all of the following are fulfilled: concealed randomization, blinded outcome adjudication and an intention to treat analysis. *Level 2 study:* If any one of the above characteristics are unfulfilled.

Study	Population	Methods	Intervention	Mortalit	y # (%)†	Infectior	ns # (%)‡
		(score)		Experimental	Control	Experimental	Control
1) Young 1996	Severely head injured patients, ventilated N=68	C.Random: not sure ITT: yes Blinding: double (12)	12 mg elemental zinc via PN, then progressing to oral zinc vs. 2.5 mg elemental zinc, then progressing to oral placebo	4/33 (12)	9/35 (26)	NR	NR
2) Berger 1998	Burns > 30 % TBSA N=20	C.Random: not sure ITT: yes Blinding: double blind (11)	IV Copper (40.4 μmol), selenium (2.0 μmol), zinc (406 μmol) + standard trace elements vs. standard trace elements elements (Copper 20 μmol, selenium 0.4μmol, zinc 100 μmol) X 8 days, all received early EN.	1/10 (10)	0/10 (0)	1.9 ± 0.9 (1-4) per patient	3.1 ± 1.1 (2-5) per patient
3) Berger 2001*	Trauma patients, surgical ICU N=32	C.Random: not sure ITT: no Blinding: single (7)	IV Selenium + α tocopherol + zinc vs placebo (All groups received enteral nutrition)	0/11 (0)	1/11 (9)	3/11 (27)	3/11 (27)
4) Berger 2007	Burns > 20 % BSA N=21	C.Random: not sure ITT: yes Blinding: no (8)	IV 100 mls of Copper (59 μmol) + Selenium (375 μgm + zinc (574 μmol) vs NaCl (0.9%) from admission for 5-15 days. Both groups were on EN.	1/11 (9)	1/10 (10)	2.1 ± 1.0 per patient	$3.6 \pm per patient$
5) Berger 2008	Mixed ICU N=200	C.Random: not sure ITT: yes Blinding: no (10)	IV Selenium supplementation loading dose 540 μg/day + zinc (60 mg) + Vit C 2700 mg + Vit B 305 mg + Vit E enteral 600 mg + Vit E 12.8 mg IV for 2 days followed by half the dose of all vs. standard vitamins. (All groups received EN or PN).	ICU 8/102 (8) Hospital 14/102 (14) 3 month 14/602 (14)	ICU 5/98 (5) Hospital 9/98 (11) 3 month 11/98 (11)	36/102 (35)	34/98 (35)

Table 1. Randomized s	studies evaluating	zinc supple	mentation in o	critically ill p	atients
				···· · · · · · · · · · · · · · ·	

Study	LOS	days	Ventilat	tor days	Сс	ost	Other	
otaaj	Experimental	Control	Experimental	Control	Experimental	Control	Experimental Control	
1) Young 1996	NR	NR	NR	NR	NR	NR	NR	
2) Berger 1998	ICU 30 ± 12 (10) Hospital 54 ± 27 (10)	ICU 39 ± 13 (10) Hospital 66 ± 31 (10)	9 ± 10 (10)	12 ± 9 (10)	NR	NR	NR	
3) Berger 2001*	ICU 5.8 ± 4.4 (11) Hospital 60 ± 48 (11)	ICU 6.1 ± 6.0 (11) Hospital 59 ± 37 (11)	4.1 ± 3.6 (11)	4.2 ± 5.2 (11)	NR	NR	Organ failure 3/11 (27) 4/11 (36)	
4) Berger 2007	ICU 35 ± 27 (11)	ICU 47 ± 37 (10)	7.6 ± 6 (11)	12.6 ± 6 (10)	NR	NR	NR	
5) Berger 2008	ICU 5.8 ± 5.4 (102) Hospital 23 ± 20 (102)	ICU 5.4 ± 5.7 (98) Hospital 26 ± 20 (98)	Vent free days 26.1 ± 5.7	Vent free days 26.6 ± 5.2	NR	NR	NR	

Table 1. Randomized studies evaluating zinc supplementation in critically ill patients (continued)

C.Random: concealed randomization

ITT: intent to treat

NR: not reported

‡ refers to the # of patients with infections unless specified † presumed hospital mortality unless otherwise specified

* only data pertaining to the selenium + α tocopherol + zinc vs placebo groups reported here **RR (CI): Relative risk (95 % confidence intervals)

Figure 1. Mortal Review: Parentera Comparison: 01 Parent Outcome: 01 Mortalit	lity I Zinc eral Zinc vs control ty						
Study	Parenteral zinc	Control		RR (random)	Weight	RR (random)	
or sub-category	nΛ	nN		95% CI	%	95% CI	Year
Young 1996	4/33	9/35		_	70.94	0.47 [0.16, 1.38]	1996
Berger 1998	1/10	0/10				3.00 [0.14, 65.90]	1998
Berger 2001	0/11	1/11		-	- 8.58	0.33 [0.02, 7.39]	2001
Berger 2007	1/11	1/10			11.85	0.91 [0.07, 12.69]	2007
Total (95% Cl)	65	66			100.00	0.58 [0.23, 1.44]	
Total events: 6 (Parenteral :	zinc), 11 (Control)			-			
Test for heterogeneity: Chi ²	² = 1.47, df = 3 (P = 0.69), l ² = 0%						
Test for overall effect: Z =	1.18 (P = 0.24)						
			0.1 0.1	2 0.5 1 2	5 10		
			F	avours zinc Favours con	ntrol		

Figure 2. Hospital LOS

Review: Comparison: Outcome:	Parenteral Zinc 01 Parenteral Zi 02 Hospital Lenj	nc vs contr gth of Stay	ol										
Study or sub-categor	у	N	^p arenteral zinc Mean (SD)	N	Control Mean (SD)		١	MMD (rando 95% Cl	m)	Weight %		WMD (random) 95% Cl	Year
Berger 1998		10	54.00(27.00)	10	66.00(31.00)		_			66.40	-12.00	[-37.48, 13.48]	1998
Berger 2001		11	60.00(48.00)	11	59.00(37.00)		-	-+		33.60	1.00	[-34.81, 36.81]	2001
Total (95% Cl) Test for hetero Test for overall	geneity: Chi² = 0.3 I effect: Z = 0.72 (I	21 4, df = 1 (P P = 0.47)	= 0.56), I² = 0%	21				+		100.00	-7.63	[-28.39, 13.13]	
						-100	-50	ó	50	100			
							Favours	zine Fav	ours cont	rol			

Figure 3. ICU LOS

Review: Comparison: Outcome:	v: Parenteral Zinc rison: 01 Parenteral Zinc vs control ne: 03 ICU Length of stay									
Study or sub-category	У	F	Parenteral zinc Mean (SD)	N	Control Mean (SD)		VVMD (random) 95% Cl	Weight %	WMD (random) 95% Cl	Year
Berger 1998		10	30.00(12.00)	10	39.00(13.00)	+		25.13	-9.00 [-19.97, 1.97]	1998
Berger 2001		11	5.80(4.40)	11	6.10(6.00)		_	70.04	-0.30 [-4.70, 4.10]	2001
Berger 2007		11	35.00(27.00)	10	47.00(37.00)			4.84	-12.00 [-39.94, 15.94]	2007
Total (95% CI)		32		31				100.00	-3.05 [-9.34, 3.24]	
Test for heterog	geneity: Chi ² = 2.61,	df = 2 (P :	= 0.27), I² = 23.5%				_			
Test for overall	l effect: Z = 0.95 (P =	= 0.34)								
						-10	-5 0 5	10		
						-	F			

Favours zinc Favours control