9.2b Combined Parenteral and Enteral Glutamine Supplementation

There were no new randomized controlled trials since the 2015 update and hence there are no changes to the following summary of evidence.

Question: Compared to placebo, does combined enteral and parenteral glutamine-supplementation result in improved clinical outcomes in critically ill patients?

Summary of evidence: There was one level 1 study and 1 level 2 study on glutamine supplementation administered via both PN and EN that were included.

Mortality: Based on the single study that reported on this outcome, glutamine supplementation administered via both PN and EN was associated with a significant increase in hospital (RR 1.20, 95% CI 1.02, 1.40, p=0.02), 28-day (RR 1.19, 95% CI 1.00, 1.42, p=0.05), 3-month (RR 1.20, 95% CI 1.04, 1.38, p=0.01), and 6-month mortality (RR 1.19, 95% CI 1.03, 1.36, p=0.02); and was associated with a trend towards a increase in 14-day mortality (RR 1.21, 95% CI 0.99, 1.48, p=0.07).

Infections: Based on the single study that reported on this outcome, glutamine supplementation administered via both PN and EN had no effect on overall infectious complications (RR 1.10, 95% CI 0.92, 1.31, p=0.32) or ventilator associated pneumonia (RR 1.08, 95% CI 0.82, 1.43, p=0.59).

Length of Stay: Based on the single study that reported on this outcome, glutamine supplementation administered via both PN and EN was associated with a trend towards an increase in ICU length of stay (WMD 1.80, 95% CI -0.76, 4.36, p=0.17), but had no effect on hospital length of stay (WMD 1.30, 95% CI -4.05, 6.65, p=0.63).

Duration of ventilation: Based on the 2 studies, no effect in duration of ventilation was seen with (WMD 0.28, 95% CI -2.85, 3.41, p=0.86; figure 1).

Conclusions:

1) Combined parenteral and enteral glutamine supplementation is associated with an increase in hospital, 28-day, 3-month, and 6-month mortality, and may be associated with an increase in 14-day mortality.

2) Combined parenteral and enteral glutamine supplementation has no effect on overall infectious complications, ventilator associated pneumonia or duration of mechanical ventilation.

3) Combined parenteral and enteral glutamine supplementation may be associated with an increase in ICU length of stay but has no effect on hospital length of stay.

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	Mortality # (%	%)*	Infections # (%)†		
	Population	(score)	intervention	GLN PN+EN P	Placebo	GLN PN+EN	Placebo	
1) Heyland 2013	Multicentre mixed ICUs N=1218	C.Random: yes ITT: yes Blinding: double (12)	GLN supplementation (0.35 g/kg/day) parenterally vs placebo; additional GLN supplementation (30 g/day) enterally vs placebo	227/611 (37) 14 RR 1.20, 95% Cl 1.02, 1.44 14-day 157/611 (26) 1 RR 1.21, 95% Cl 0.99, 1.44 28-day 198/611 (32) 10 RR 1.19, 95% Cl 1.00, 1.43 3-month 252/611 (39) 20 RR 1.20, 95% Cl 1.04, 1.34 6-month	14-day 129/607 (21) 18, p=0.07 28-day 65/607 (27) 12, p=0.05 3-month 09/607 (32) 18, p=0.01 6-month 221/607 (37)	All 183/611 (30) RR 1.10, 95% CI 0.92 VAP 88/611 (14) RR 1.08, 95% CI 0.82	VAP 78/607 (13)	
2) Koksal 2014	Septic, malnourished ICU patients N=120	C.Random: yes ITT: other Blinding: single (outcomes)	15 g/day parenteral glutamine + 15 g/day enteral glutamine + EN vs	NR		NR		
		(9)	EN, no placebo, no supplemental glutamine					

Table 1. Randomized studies evaluating glutamine (PN + EN) in critically ill patients

Table 1. Randomized studies evaluating glutamine (PN + EN) in critically ill patients (continued)

Chudu	LOS	days‡	Ventilator days‡			
Study	GLN PN+EN	Placebo	GLN PN+EN	Placebo		
1) Heyland 2013	Hospital 31.0 ± 52.6 (611)	ICU 13.1±14.0 (607) CI -0.76, 4.36, p=0.17 Hospital 29.7 ± 42.1 (607) CI -4.05, 6.65, p=0.63	11.6 ± 26.3 (611)	9.8 ± 12.3 (607)		
2) Koksal 2014		NR	12.9±5.3	14.3±5.4		
NR: not reported VAP: ventilator associated GLN: glutamine	pneumonia	ITT: intent to treat ICU: intensive care unit PN: parenteral nutrition	١	C. Random: concealed randomization WMD: weighted mean difference; CI: EN: enteral nutrition		

onfidence interval

* presumed hospital mortality unless otherwise specified

† refers to the # of patients with infections unless specified

‡ LOS and ventilation statistics calculated using all patients who were discharged; for patients who died, death date was substituted for discharge date.

Critical Care Nutrition: Systematic Reviews March 2021

Figure 1: Duration of Mechanical Ventilation

	EN+PN Glutamine			Control			Mean Difference		Mean Difference		
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% CI	Year	r IV, Random, 95% Cl	
Heyland	11.6	26.3	611	9.8	12.3	607	52.6%	1.80 [-0.50, 4.10]	2013	3 📮	
Koksal	12.9	5.3	30	14.3	5.4	30	47.4%	-1.40 [-4.11, 1.31]	2014	4 📕	
Total (95% CI)			641			637	100.0%	0.28 [-2.85, 3.41]			
Heterogeneity: Tau² = 3.48; Chi² = 3.11, df = 1 (P = 0.08); l² = 68% Test for overall effect: Z = 0.18 (P = 0.86)										-100 -50 0 50 10 Favours EN+PN glutamine Favours control	00

References

Included Articles

- 1. Heyland D, Muscedere J, Wischmeyer PE, Cook D, Jones G, Albert M, Elke G, Berger MM, Day AG for the Canadian Critical Care Trials Group. A Randomized Trial of Glutamine and Antioxidants in Critically III Patients. N Engl J Med 2013;368(16):1487-95.
- 2. Koksal GM, Erbabacan E, Tunali Y, Karaoren G, Vehid S, Oz H. The effects of intravenous, enteral and combined administration of glutamine on malnutrition in sepsis: a randomized clinical trial. Asia Pac J Clin Nutr. 2014;23(1):34-40.

Excluded Articles

No other articles were found.