March 2021

4.7 Enteral Nutrition: Other Formulas: ß Hydroxyl Methyl Butyrate (HMB)

Question: Does the use of a formula supplemented with ß hydroxyl methyl butyrate (HMB) result in better outcomes in the critically ill

adult patient?

Summary of evidence: There was one level 2 study that studied the effect of supplementation of an enteral formula with ß hydroxyl methyl butyrate

(HMB), alone to standard enteral nutrition (EN) with an isonitrogenous isocaloric placebo in trauma patients (Kuhls 2007). One level 1 study compared

HMB in addition to arginine and glutamine to isocaloric standard EN in a mixed ICU population, with both groups receiving early rehabilitation with

electrical muscle stimulation (Nakamura 2019). The data pertaining to the second intervention from the Kuhls 2007 study comparing enteral nutrition

supplemented with ß hydroxyl methyl butyrate, arginine and glutamine (Juven®) to standard enteral nutrition alone, is described in section 4.1: Diets

supplemented with Arginine and select other nutrients.

Mortality: When the data from the two studies were aggregated, formula supplemented with HMB had no effect on mortality compared to standard

EN (RR 0.53, 95% CI 0.17, 1.59, p=0.26, test for heterogeneity  $I^2 = 0\%$ , figure 1).

**Infections:** Based on one study, formula supplemented with HMB had no effect on the number of infectious complications per patient (WMD 0.20,

95% CI -1.33, 1.73, p=0.80).

Length of Stay (LOS): HMB supplemented formula had no effect on ICU LOS (WMD 1.31, 95% CI -4.53, 7.16, p=0.66, test for heterogeneity I<sup>2</sup> =

47%, figure 2) or hospital LOS (WMD 4.63, 95% CI -11.36, 20.62, p=0.57, test for heterogeneity I<sup>2</sup> = 83%, figure 3), when compared to standard EN.

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**Ventilator days:** When the data from both studies were aggregated, HMB supplemented formula compared to standard EN, had no effect on the number of ventilator days (WMD -0.26, 95% CI -2.08, 1.57, p=0.78, test for heterogeneity I<sup>2</sup> =6%, figure 4).

**Other:** In the Kuhls 2007 study, there was no effect of HMB supplementation on nitrogen intake however nitrogen balance was significantly better in the HMB group (p=0.05). HMB supplementation did not inhibit femoral muscle loss compared to standard EN (11.4±8.1% vs. 14.4±7.1% respectively; p=0.18, Nakamura 2019).

## **Conclusions:**

## Compared to standard enteral nutrition,

- 1) Supplementation with ß hydroxyl methyl butyrate (HMB) has no effect on mortality or duration of mechanical ventilation.
- 2) Supplementation with ß hydroxyl methyl butyrate (HMB) has no effect on ICU length of stay.
- 3) Supplementation with ß hydroxyl methyl butyrate (HMB) has no effect on hospital length of stay.
- 4) Supplementation with ß hydroxyl methyl butyrate (HMB) may be associated with better nitrogen balance in trauma patients.
- 5) Supplementation with ß hydroxyl methyl butyrate (HMB) does not inhibit femoral muscle loss in heterogenous ICU 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.

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Table 1. Randomized studies evaluating other enteral formulas in critically ill patients

Study	Population	Methods (score)	Intervention	Mortality # (%)† HMB vs. Control	Infections # (%)‡ HMB vs. Control
1) Kuhls 2007*	Trauma patients in ICU Injusry Severity Score >18 N=100	C.Random: No/not sure ITT: No** Blinding: Double (10)	Standard EN + supplement of 3 gms ß hydroxyl methyl butyrate (HMB) vs. Standard EN + isonitrogenous placebo supplement Isonitrogenous/isocaloric 25kcal/kg/day, 1.5g pro/kg/day	0/28 (0) vs. 2/22 (9)	# infections per patient 4.8 ± 2.65 (28) vs. 4.6 ± 2.81 (22) WMD 0.20, 95% CI -1.33, 1.73, p=0.80
2) Nakamura 2020	Mix ICU population N=88	C.Random: Yes ITT: Yes Blinding: single (12) Level: 2	Daily 3 grams HMB+14 grams arginine+14 grams of glutamine vs. standard EN. Both groups received early rehabilitation with electrical muscle stimulation from day 2. Isocaloric 20-30 Kcal/kg	<b>28 day</b> 4/45 (8.7%) vs. 6/43 (13.6%); p=0.30	NR

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Table 1. Randomized studies evaluating other enteral formulas in critically ill patients (continued)

Study	LOS days HMB vs. Control	Ventilator days HMB vs. Control	Other HMB vs. Control
1) ) Kuhls 2007*	ICU 28.9 ± 17.46 (28) vs. 22.4 ± 17.35 (22)  Hospital 44.4 ± 23.28 (28) vs. 30.3 ± 22.98 (22)	24.2 ± 12.70 (28) vs. 20.9 ± 12.66 (22)	# Patients with SIRS Score >3 or >4 Significantly less in HMB group on day 3 (p<0.01) and day 7 (p<0.02)  Average Nitrogen Balance -6.50 ± 6.35 vs9.0 ± 6.10  Change in Nitrogen Balance Comparing Week 1 to Week 2 Greater in HMB vs placebo (p<0.05)
2) Nakamura 2020	ICU 5.4 ±3.5 (45) vs. 5.8 ±3.8 (43); p=0.83  Hospital 21.9 ±8.8 (45) vs. 24.3 ±7.8 (43); p=0.18	4.8 ± 2.4 (45) vs. 5.3 ± 3.8 (43); p=0.45	Subgroup of femoral muscle analyses (n=50)  Total energy, kcal/d  1015±319 vs. 1098±462; p=0.90  Total protein, g/d  54.5±10.8 vs. 50.1±14.1; p=0.31  Femoral muscle loss, %  11.4±8.1 vs. 14.4±7.1; p=0.18

<sup>\*</sup> all "standard error" reported in the Kuhls 2007 study have been converted to "standard deviation"

ICU: Intensive care unit ITT: intent to treat C. Random: concealed randomization EN: enteral nutrition

SIRS: systemic inflammatory response syndrome WMD: weighted mean difference; CI: Confidence interval

Data pertaining to enteral nutrition supplemented with ß hydroxyl methyl butyrate, arginine and glutamine (Juven®) to standard enteral nutrition alone not shown here. Refer to section 4.1: Diets supplemented with Arginine and select other nutrients

<sup>\*\* 100</sup> pts randomized but only 72 reported on as 72 received at least 7 days of supplementation. Additional statistical exclusion criteria were established based on 50% treatment compliance, therefore 72 pts were used. † presumed hospital mortality unless otherwise specified

<sup>‡</sup> refers to the # of patients with infections unless specified

Figure 1. Mortality

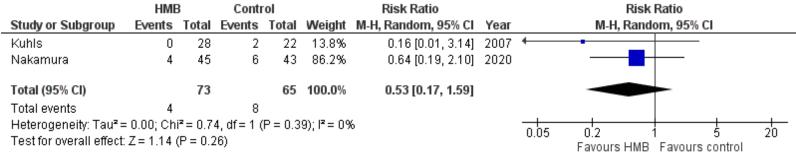


Figure 2. ICU Length of Stay

	HMB			C	ontrol		Mean Difference				Mean Difference			
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% CI	Year		IV, Random, 95% CI			
Kuhls	28.9	17.46	28	22.4	17.35	22	24.8%	6.50 [-3.22, 16.22]	2007			-		
Nakamura	5.4	3.5	45	5.8	3.8	43	75.2%	-0.40 [-1.93, 1.13]	2020					
Total (95% CI)			73			65	100.0%	1.31 [-4.53, 7.16]				<b>•</b>		
Heterogeneity: Tau² : Test for overall effect	-		-	= 1 (P =	0.17); l³	= 47%	ı			-100	-50 Favours HME	0 8 Favo	50 ours control	100

Figure 3. Hospital Length of Stay

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		HMB		(	ontrol			Mean Difference			Me	an Differen	ce	
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% CI	Year		IV, Random, 95% CI			
Kuhls	44.4	23.28	28	30.3	22.98	22	42.6%	14.10 [1.19, 27.01]	2007				-	
Nakamura	21.9	8.8	45	24.3	7.8	43	57.4%	-2.40 [-5.87, 1.07]	2020			•		
Total (95% CI)			73			65	100.0%	4.63 [-11.36, 20.62]				-		
Heterogeneity: Tau² : Test for overall effect		•		f=1 (P:	= 0.02);	l² = 83°	%			-100	-50 Favours	0 HMB Favo	50 urs control	100

Figure 4. Ventilator Days

_	- I	НМВ		0	ontrol		Mean Difference				Mean Difference			
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% CI	Year		IV, Random, 95% CI			
Kuhls	24.2	12.7	28	20.9	12.66	22	6.4%	3.30 [-3.78, 10.38]	2007			<u>+</u>		
Nakamura	4.8	2.4	45	5.3	3.8	43	93.6%	-0.50 [-1.83, 0.83]	2020			-		
Total (95% CI)			73			65	100.0%	-0.26 [-2.08, 1.57]				•		
Heterogeneity: Tau $^z$ = 0.47; Chi $^z$ = 1.07, df = 1 (P = 0.30); $I^z$ = 6% Test for overall effect: Z = 0.27 (P = 0.78)							-100	-50 Favours I	O HMB Favo	50 urs control	100			

## References:

## **Included Articles**

- 1. Kuhls DA, Rathmacher JA, Musngi MD, Frisch DA, Nielson J, Barber A, MacIntyre AD, Coates JE, Fildes JJ. Beta-hydroxy-beta-methylbutyrate supplementation in critically ill trauma patients. J Trauma. 2007 Jan;62(1):125-31; discussion 131-2. PubMed PMID: 17215743.
- 2. Nakamura K, Kihata A, Naraba H, Kanda N, Takahashi Y, Sonoo T, Hashimoto H, Morimura N. β-Hydroxy-β-methylbutyrate, Arginine, and Glutamine Complex on Muscle Volume Loss in Critically III Patients: A Randomized Control Trial. JPEN J Parenter Enteral Nutr. 2020 Feb;44(2):205-212. doi: 10.1002/jpen.1607. Epub 2019 May 27. PMID: 31134640.

E	cluded Articles	Reason
1.	Hsieh LC, Chien SL, Huang MS, Tseng HF, Chang CK. Anti-inflammatory and anticatabolic effects of short-term	No clinical outcomes
	beta-hydroxy-beta-methylbutyrate supplementation on chronic obstructive pulmonary disease patients in	
	intensive care unit. Asia Pac J Clin Nutr. 2006;15(4):544-550.	
2.	Bear DE, Langan A, Dimidi E, et al. β-Hydroxy-β-methylbutyrate and its impact on skeletal muscle mass and	Systematic review
	physical function in clinical practice: a systematic review and meta-analysis. Am J Clin Nutr. 2019;109(4):1119-	
	1132. doi:10.1093/ajcn/nqy373	