

4.2a Composition of Enteral Nutrition: (Carbohydrate/fat): High fat/low CHO

Question: Does a high fat/low CHO enteral formula affect outcomes in the critically ill adult patient?

Summary of evidence: There were eight level 2 studies and one level 1 study that compared a high fat, low CHO formula to a standard formula. Two studies compared Pulmocare (55% fat, 28 % CHO); one compared Novasource Diabetic Plus (40% fat, 40 % CHO), one compared Diben (45% fat, 37% CHO), one compared Glucerna 1.5 (46% fat, 33% CHO) to standard formula (up to 35% fat and 53% CHO) and one compared Glucerna select (50 % fat, 30% CHO, 20 % protein 1 Kcal/mL) to Nutrison Protein Plus (35% fat, 45 % CHO, 20 % protein, 1.25 Kcal/mL). One study compared two hospital made formulas (45% fat, 35% CHO vs. 30% fat, 50% CHO). Two studies compared two different high fat formulas to a standard formula: Mesejo 2015's experimental EN formulas were Diaba HP (40% fat, 33% CHO) and Glucerna Select (49% fat, 30% CHO) and Nourohommadi 2017's experimental formulas contained 45% fat (50:50 olive and sunflower oil), 35% CHO and 45% fat (100% sunflower oil), 35% CHO. The data for the two intervention arms in Mesejo 2015 and Nourohommadi 2017 have been combined in the meta-analyses.

Mortality: Eight studies reported on mortality and there were no differences between the groups for overall mortality when the data were aggregated (RR 1.12, 95% CI 0.82, 1.55, $p=0.45$, test for heterogeneity $I^2=0\%$; Figure 1) and for ICU mortality (RR 1.12, 95% CI 0.78, 1.62, $p=0.52$, test for heterogeneity $I^2=0\%$; Figure 2).

Infections: Three studies (Mesejo 2003, Mesejo 2015 and Vahabzadeh 2019) reported infectious complications and found no differences between the two groups (RR 0.94, 95% CI 0.67, 1.33, $p=0.74$, test for heterogeneity $I^2=0\%$; Figure 3).

LOS: Two studies (Mesejo 2003, Nourohommadi 2017) reported on ICU length of stay as means and standard deviations and no differences were seen between the two groups when the data were aggregated (WMD -2.07, 95% CI -6.98, 2.84, $p=0.41$; figure 4). Data from four studies were not included in the analyses due to carrying reporting outcomes (three reported median and ranges, one reported ICU free days).

Ventilator days: Duration of mechanical ventilation was significantly lower in the high fat group in one study (Al Saady 1994 $p<0.001$), no difference found in the van de Berg 1994 study or the Mesejo 2003 study. For the two studies that reported ventilation duration in mean and standard deviation, a significant reduction in duration was seen in the high fat group (WMD -2.87, 95% CI -3.59, -1.14, $p=0.0002$; Figure 5).

Other complications: Six studies reported on glycemic control or glucose levels, three reported significantly lower blood sugars in the group receiving the higher fat, lower CHO formula (Mesejo 2003, Mesejo 2015 [Diaba HP group], Doola 2019). Wewalka 2018 and Vahabzadeh 2019 found no statistically significant differences in fasting blood glucose levels between groups. Van Steen 2018 showed a trend in a reduction of hyperglycemic events in the high fat group, but there was no difference between groups regarding hypoglycemic events. Insulin use was significantly lower in the high fat, low CHO group compared to the lower fat, higher CHO group in one study (Doola 2019) but not in the other study (Vahabzadeh 2019). Four studies reported on diarrhea and there was a trend towards a reduction in diarrhea in the high fat, low CHO formula fed groups (RR 0.81, 95% CI 0.64, 1.04, $p=0.10$, test for heterogeneity $I^2=0\%$; Figure 6).

Conclusions:

- 1) A high fat, low CHO enteral formula may be associated with a reduction in ventilated days in medical ICU patients with respiratory failure and better glycemic control in critically ill patients with hyperglycemia.
- 2) A high fat, low CHO enteral formula has no effect on mortality, infections or LOS found between the critically ill patients receiving high fat/low CHO formula or standard.
- 3) A high fat, low CHO formula may be associated with less diarrhea 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

Table 1. Randomized Studies Evaluating High Fat/Low CHO Enteral Nutrition In Critically ill Patients

Study	Population	Methods (score)	Intervention	Mortality # (%)**		RR (CI) or p value	Infections # (%)		RR (CI) or p value
				High fat/low CHO NR	Standard NR		High fat/low CHO NR	Standard NR	
1. van den Berg 1994	Medical ICU patients with COPD Chronically ventilated N=32	C.Random: not sure ITT: yes Blinding: no (5)	55% fat, 28 % CHO (Pulmocare) vs 30 % fat, 53 % CHO (standard, Ensure Plus)	High fat/low CHO NR	Standard NR	NR	High fat/low CHO NR	Standard NR	NR
2. Al Saady 1989	Ventilated patients Acute respiratory failure N=40	C.Random: not sure ITT: no Blinding: double (9)	55% fat, 28 % CHO (Pulmocare) vs 30 % fat, 53 % CHO (standard, Ensure Plus)	3/9 (33)	3/11 (27)	1.22 (0.32-4.65)	NR	NR	NR
3. Mesejo 2003	Critically ill pts with Diabetes or hyperglycemia from 2 different centers N=50	C.Random: not sure ITT: yes Blinding: single (9)	40% fat, 40 % CHO (Novasource Diab Plus) vs. 29 % fat, 49 % CHO (Standard, Isosource Protein)	ICU 8/26 (31)	ICU 7/24 (29)	1.05 (0.45, 2.47)	10/26 (38.5)	8/24 (33)	1.15 (0.55, 2.43)
4) Mesejo 2015	Critically ill patients meeting ADA criteria for diabetes/hyperglycemia. Multi-centre. N=157	C.Random: yes ITT: no Blinding: single (11)	40% fat, 33% CHO (Diaba HP - experimental) vs 49% fat, 30% CHO (Glucerna Select – experimental) vs 34% fat, 44% CHO (Isosource Protein Fibra – control)	<u>Diaba HP</u> 28 day 11/52 (21.1) <u>6 Month</u> 16/52 (30.7) <u>Glucerna Select</u> 28 day 13/52 (25) <u>6 Month</u> 18/52 (34.6)	28 day 10/53 (18.9) 6 Month 20/53 (37.7)		<u>Diaba HP</u> 18/52 (34.6) <u>Glucerna Select</u> 23/52 (44.2)	23/53 (43.3)	
5) Nourohhamadi 2017	Mixed ICU patients. Single centre. N=42	C.Random: yes ITT: yes Blinding: double (10)	45% fat (half olive, half sunflower oil), 35% CHO vs 45% fat (all sunflower oil), 35% CHO vs 30% fat, 50% CHO.	<u>Olive/Sunflower</u> ICU 3/16 (18.7) <u>Sunflower</u> ICU 6/16 (37.5)	6/16 (37.5)		NR	NR	NR
6) Wewalka 2018	Medical ICU pts. Single centre. N=60	C.Random: no ITT: yes Blinding: no (9)	45% fat, 37% CHO (Diben) vs 30% fat, 55% CHO (Fresubin original fibre). Formulas contain 2.3 g fibre/100ml and 1.5 g fibre/100 ml, respectively.	ICU 13/30 (43)	ICU 9/30 (30)		NR	NR	

<p>7) Van Steen 2018</p>	<p>Medical and surgical critically ill patients N=107</p>	<p>C.Random: yes ITT: no Blinding: no (8)</p>	<p>46% fat, 33% CHO, 21% protein (Glucerna 1.5) vs 35% fat, 50% CHO, 15% protein (Fresubin Energy Fibre + protein supplement (Resource Instant Protein) 3x qd to make relatively equal in protein to intervention group.</p>	<p>ICU 9/52 (17)</p>	<p>ICU 8/49 (16)</p>		<p>NR</p>	<p>NR</p>	
<p>8) Doola 2019</p>	<p>Critically ill patients requiring insulin for hyperglycemia while on EN N=41</p>	<p>C.Random: yes ITT: no Blinding: double (8)</p>	<p>50 % fat, 30% CHO, 20 % protein (Glucerna select 1 Kcal/mL) vs. 35% fat, 45 % CHO, 20 % protein (Nutrison Protein Plus (1.25 Kcal/mL) Target for both 25 kcal/kg; 1.2 g protein/kg for 2 days</p>	<p>28 day 1/21 (5%)</p>	<p>28 day 2/20 (10%)</p>	<p>0.60</p>	<p>NR</p>	<p>NR</p>	
<p>9) Vahabzadeh 2019</p>	<p>Critically ill patients with hyperglycemia while on EN N=88</p>	<p>C.Random: no ITT: no Blinding: double (5)</p>	<p>45% fat,35% CHO, 20% protein hospital made formula vs. 30% fat, 50% CHO, 20% protein hospital made formula. Target for both 25-30 Kcal/kg for up to 15 days</p>	<p>ICU 6/41 (14%)</p>	<p>ICU 4/39 (10%)</p>		<p>Sepsis 0/41</p>	<p>Sepsis 1/39 (2.6%)</p>	

Table 1. Randomized Studies Evaluating High Fat/Low CHO Enteral Nutrition In Critically ill Patients (continued)

Study	LOS days		Ventilator days		Other	
	High fat/low CHO	Standard	High fat/low CHO	Standard	High fat/low CHO	Standard
1. van den Berg 1994	NR	NR	4 (median)	6 (median)	High fat/low CHO Gastric retention 1/15 (7)	Standard 1/17 (6)
2. Al Saady 1994	NR	NR	3.6 ± 0.7	6.2 ± 1.5	Diarrhea 3/9 (33) 3/11 (27)	
3. Mesejo 2003	ICU 14.8 ± 9.4	ICU 14.8 ± 8.8	8.7 ± 6.2	9.4 ± 6.0	Plasma Glucose Levels (mmol/L) 9.8 ± 2.4 12.4 ± 2.6	
4) Mesejo 2015	<u>Diaba HP</u> ICU+ 13 (9-20) Hospital+ 27 (18-50) <u>Glucerna Select</u> ICU+ 11.5 (7.5-18) Hospital+ 30.5 (14 – 46.5)	ICU+ 12 (7-21) Hospital+ 25 (17-51)	<u>Diaba HP</u> + 7 (4-13) <u>Glucerna Select</u> + 6 (3-12)	6 (2-11)+	Plasma Glucose Levels (mg/dL) Diaba HP: 138.6 (39.1) Glucerna Select: 143.9 (45.9) Isocource: 146.1 (49.9)	
5) Nourohamadi 2017	<u>Olive/Sunflower</u> ICU* 16.6 ± 6.7 <u>Sunflower</u> ICU* 19.6 ± 8.3	ICU* 23.2 ± 12.5	NR	NR	Diarrhea Olive/sunflower: 2/16 (13.5) Sunflower: 3/16 (19.7) Control: 3/16 (19.7)	
6) Wewalka 2018	NR	NR	NR	NR	Fasting Plasma Glucose (mg/dL) 128 (110-170) 123 (98-153) Diarrhea 22/30 26/30	
7) Van Steen 2018	ICU 4.6 (2-12.6)+	ICU 4.2 (2.4-11.4)+	NR+	NR+	Patients with hypoglycemia 0/51 1/49 Patients with hyperglycemia 2/51 7/49	
8) Doola 2019	ICU 7 (4-11)* Hospital 18 (14-30)*	ICU 8 (6-11)*; p=0.80 Hospital 15 (11-20)*; p=0.10	141 [94-184]*	160 [106-219]*; p=0.70	Mean insulin use, units per hour 1.01 2.31 (p=0.017) Patients with Glycemic variability 12.6% 15.9%; p=0.01 Mean glucose control, mmol/L 8.7 10.1; p=0.002 Diarrhea 2/21 (9.5) 3/20 (15); p=0.70	

<p>9) Vahabzadeh 2019</p>	<p>ICU free days 0 (0-0)</p>	<p>ICU free days 0 (0-1); p=0.11</p>	<p>Ventilator dependency 35/41 (85.4%)</p>	<p>Ventilator dependency 34/39 (87.2%); p=0.81</p>	<p>Reduction in blood glucose by end of study, mg/dL 66.75 51.74; p=0.35 Insulin use, IU/day by end of study 0 (0-6) 0 (0-8); p=0.18</p>
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C.Random: concealed randomization
 ITT: intent to treat
 NR: Not reported

± : Mean ± Standard deviation
 RR= relative risk, CI= Confidence intervals
 *not able to analyze as not reported as mean and SD

*data obtained from correspondence with author
 **presumed to be ICU mortality unless otherwise stated

Figure 1. Overall Mortality

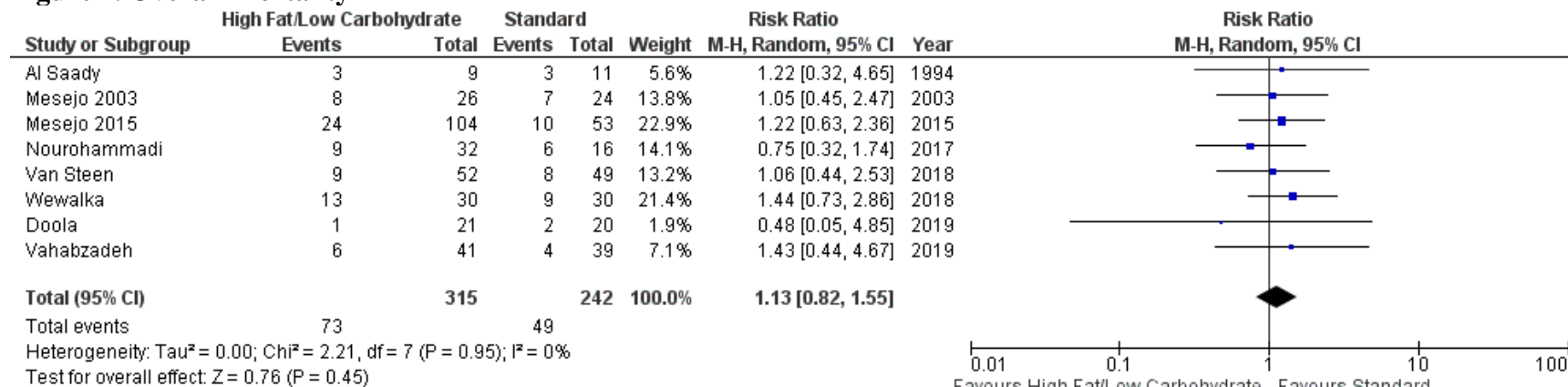


Figure 2. ICU Mortality

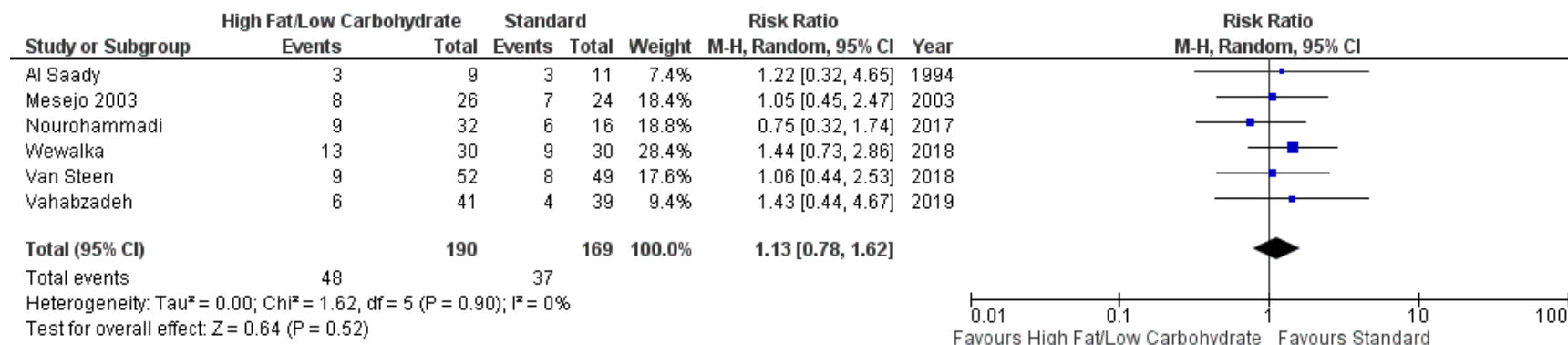


Figure 3. Infections

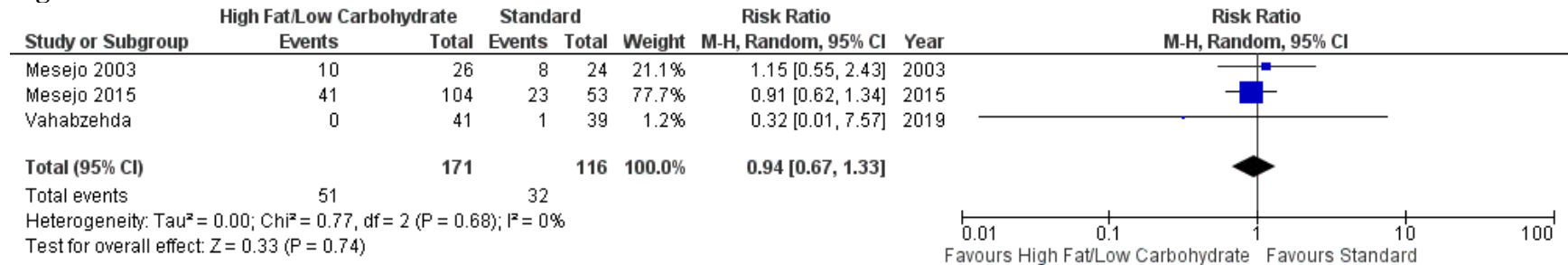


Figure 4. ICU LOS

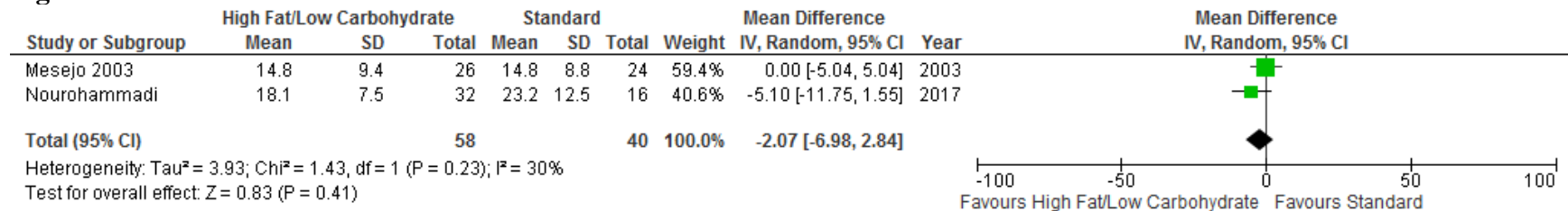


Figure 5. Mechanical Ventilation

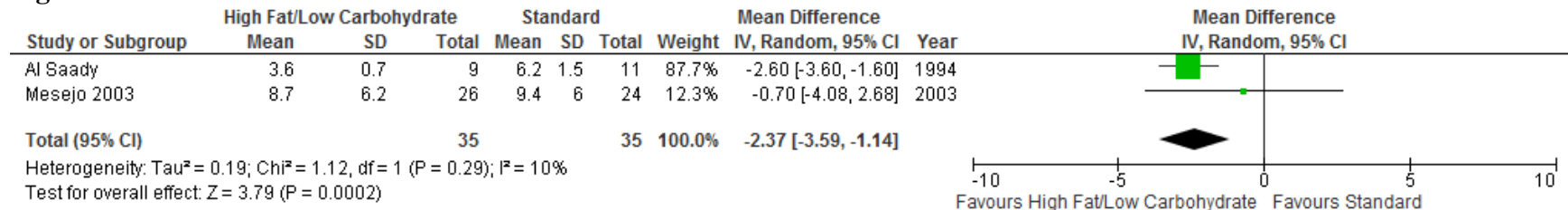
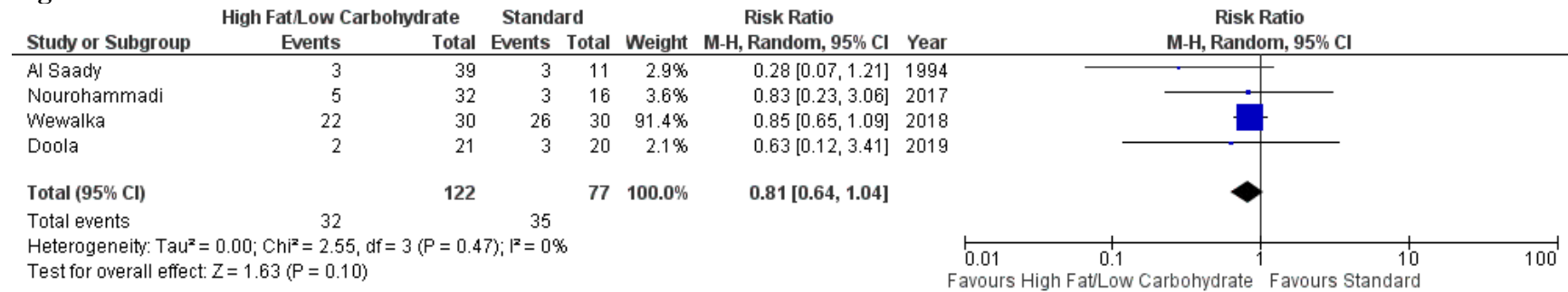


Figure 6. Diarrhea



References

Included Studies

1. Van den Berg B, Bogaard JM, Hop WC. High fat, low carbohydrate, enteral feeding in patients weaning from the ventilator. *Intensive Care Med.* 1994 Aug; 20(7): 470-5.
2. Al Saady NM, Blackmore CM, Bennett ED. High fat, low carbohydrate, enteral feeding lowers PaCO₂ and reduces the period of ventilation in artificially ventilated patients. *Intensive Care Med* 1989;15:290-5.B
3. Mesejo A, Acosta JA, Ortega C, Vila J, Fernandez M, Ferreres J, Sanchis JC, Lopez F. Comparison of a high-protein disease-specific enteral formula with a high-protein enteral formula in hyperglycemic critically ill patients. *Clin Nutr.* 2003 Jun; 22(3): 295-305.
4. Mesejo A, Montejo-González JC, Vaquerizo-Alonso C, Lobo-Tamer G, Zabarte-Martinez M, Herrero-Meseguer JI, Acosta-Escribano J, Blesa-Malpica A, Martinez-Lozano F. Diabetes-specific enteral nutrition formula in hyperglycemic, mechanically ventilated, critically ill patients: a prospective, open-label, blind-randomized, multicenter study. *Crit Care.* 2015 Nov 9;19:390.
5. Nourmohammadi M, Moghadam OM, Lahiji MN, Hatamian S, Shariatpanahi ZV. Effect of Fat-based versus Carbohydrate-based Enteral Feeding on Glycemic Control in Critically Ill Patients: A Randomized Clinical Trial. *Indian J Crit Care Med.* 2017 Aug;21(8):500-505.
6. Wewalka M, Drolz A, Seeland B, Schneeweiss M, Schmid M, Schneeweiss B, Zauner C. Different enteral nutrition formulas have no effect on glucose homeostasis but on diet-induced thermogenesis in critically ill medical patients: a randomized controlled trial. *Eur J Clin Nutr.* 2018 Apr;72(4):496-503.
7. van Steen SC, Rijkenberg S, Sechterberger MK, DeVries JH, van der Voort PHJ. Glycemic Effects of a Low-Carbohydrate Enteral Formula Compared With an Enteral Formula of Standard Composition in Critically Ill Patients: An Open-Label Randomized Controlled Clinical Trial. *JPEN.* 2018;42(6):1035-45.
8. Vahabzadeh, D., Valizadeh Hasanloei, M.A. & Vahdat Shariatpanahi, Z. Effect of high-fat, low-carbohydrate enteral formula versus standard enteral formula in hyperglycemic critically ill patients: a randomized clinical trial. *Int J Diabetes Dev Ctries* 39, 173–180 (2019). <https://doi.org/10.1007/s13410-018-0660-z>
9. Doola R, Deane AM, Tolcher DM, Presneill JJ, Barrett HL, Forbes JM, Todd AS, Okano S, Sturgess DJ. The effect of a low carbohydrate formula on glycaemia in critically ill enterally-fed adult patients with hyperglycaemia: A blinded randomised feasibility trial. *Clin Nutr ESPEN.* 2019 Jun;31:80-87. doi: 10.1016/j.clnesp.2019.02.013. Epub 2019 Mar 11. PMID: 31060838.

Excluded Studies	Reasons for exclusion
1. Schneeweiss B, Graninger W, Ferenci P, Druml W, Ratheiser K, Steger G, Grimm G, Schurz B, Laggner AN, Siostrzonek, et al. Short-term energy balance in patients with infections: carbohydrate-based versus fat-based diets. <i>Metabolism</i> . 1992 Feb; 41(2): 125-30.	No clinical outcomes
2. Diboune M, Ferard G, Ingenbleek Y, Tulasne PA, Calon B, Hasselmann M, Sauder P, Spielmann D, Metais P. Composition of phospholipid fatty acids in red blood cell membranes of patients in intensive care units: effects of different intakes of soybean oil, medium-chain triglycerides, and black-currant seed oil. <i>JPEN J Parenter Enteral Nutr</i> 1992 Mar-Apr; 16(2): 136-41.	No clinical outcomes
3. Adams S, Yeh YY, Jensen GL. Changes in plasma and erythrocyte fatty acids in patients fed enteral formulas containing different fats. <i>JPEN J Parenter Enteral Nutr</i> . 1993 Jan-Feb; 17(1): 30-	No clinical outcomes
4. Tappy L, Berger M, Schwarz JM, et al. Hepatic and peripheral glucose metabolism in intensive care patients receiving continuous high- or low-carbohydrate enteral nutrition. <i>JPEN J Parenter Enteral Nutr</i> . 1999;23(5):260-268. doi:10.1177/0148607199023005260	No clinical outcomes
5. Pohl M, Mayr P, Merti-Roetzer et al. Glycaemic control in type II diabetic tube-fed patients with a new enteral formula low in carbohydrates and high in monounsaturated fatty acids: a randomised controlled trial. <i>Eur J Clin Nutr</i> 2005;59:1221-1232.	Not ICU pts
6. Zhang G, Zou J. [Clinical application of enteral immune nutrition for chronic obstructive pulmonary disease patients]. <i>Zhonghua Yi Xue Za Zhi</i> . 2015;95(19):1501-4. Chinese	No clinical outcomes
7. Faramawy MAES, Allah AA, Batrawy SE, Amer H. Impact of high fat low carbohydrate enteral feeding on weaning from mechanical ventilation. <i>Egyptian Journal of Chest Diseases and Tuberculosis</i> . 2014;63(4):931-938.	Irreproducible findings (blenderized feeds) and possible erroneous stats (SE not SD reported?)