



Apports protéiques

Réanimation SRLF 2024

H.U.B

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des Enfants Reine Fabiola
Universitair Kinderziekenhuis
Koningin Fabiola

Professeur Jean-Charles Preiser
HUB Hôpital Erasme Bruxelles

Liens d'intérêt

- Baxter
- DIM 3
- Fresenius
- Nestlé
- Nutricia - Danone

Agenda

- Constats cliniques
- Physiologie
- Apports protéiques
 - **Précoces : oui/non**
 - **Tardifs : revalidation nutritionnelle**
- Recommandations de pratique clinique
- Futur?

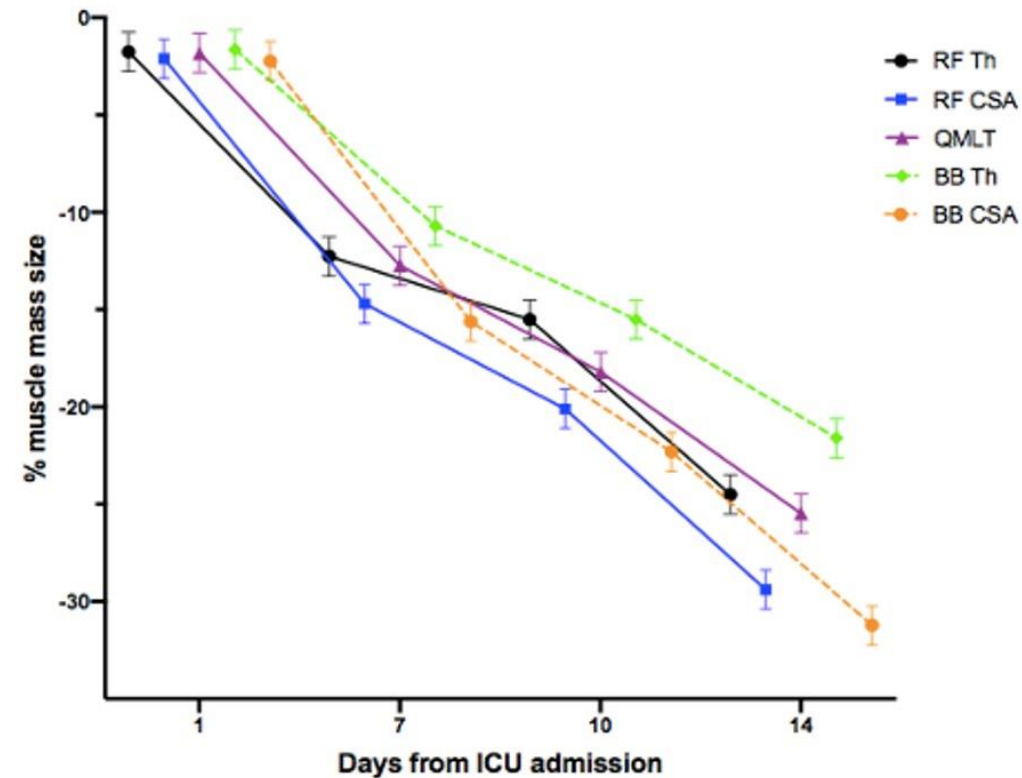
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The rate and assessment of muscle wasting during critical illness: a systematic review and meta-analysis

Brigitta Fazzini^{1*}, Tobias Märkl², Christos Costas³, Manfred Blobner^{4,5,6}, Stefan J. Schaller^{4,5}, John Prowle^{1,3}, Zudin Puthuchery^{1,3†} and Henning Wackerhage^{2†}



% RF th	-1.75	-12.25	-15.5	-24.5
% RF CSA	-2.10	-14.7	-20.1	-29.4
% QMLT	-1.82	-12.74	-18.2	-25.48
% BB Th	-1.64	-10.7	-15.5	-21.6
% BB CSA	-2.23	-15.61	-22.30	-31.22

Loss of muscle in the ICU : 2%/day

Covid-19

Evolution of the nutritional status of COVID-19 critically-ill patients: A prospective observational study from ICU admission to three months after ICU discharge

C. Rives-Lange ^{a, b, c, *}, A. Zimmer ^a, A. Merazka ^a, C. Carette ^{a, b, d}, A. Martins-Bexinga ^{b, c},
C. Hauw-Berlemont ^e, E. Guerot ^e, A.S. Jannot ^{b, f}, J.L. Diehl ^{b, e, g}, S. Czernichow ^{a, b, c},
B. Hermann ^{b, e, h}

Clin Nutr 2021;

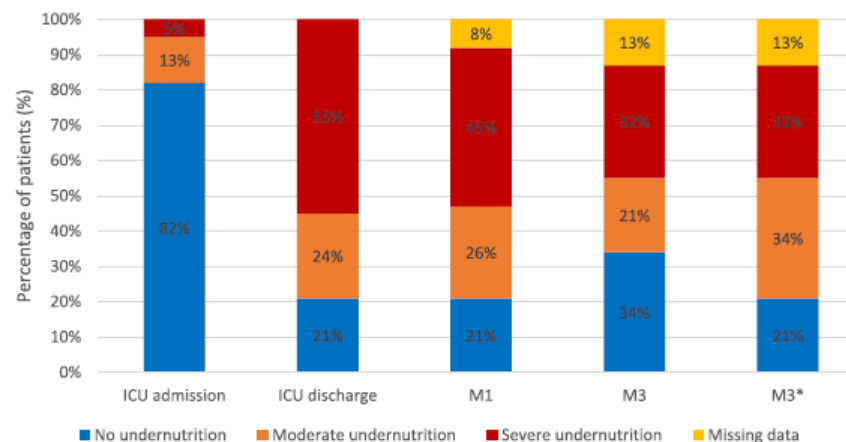


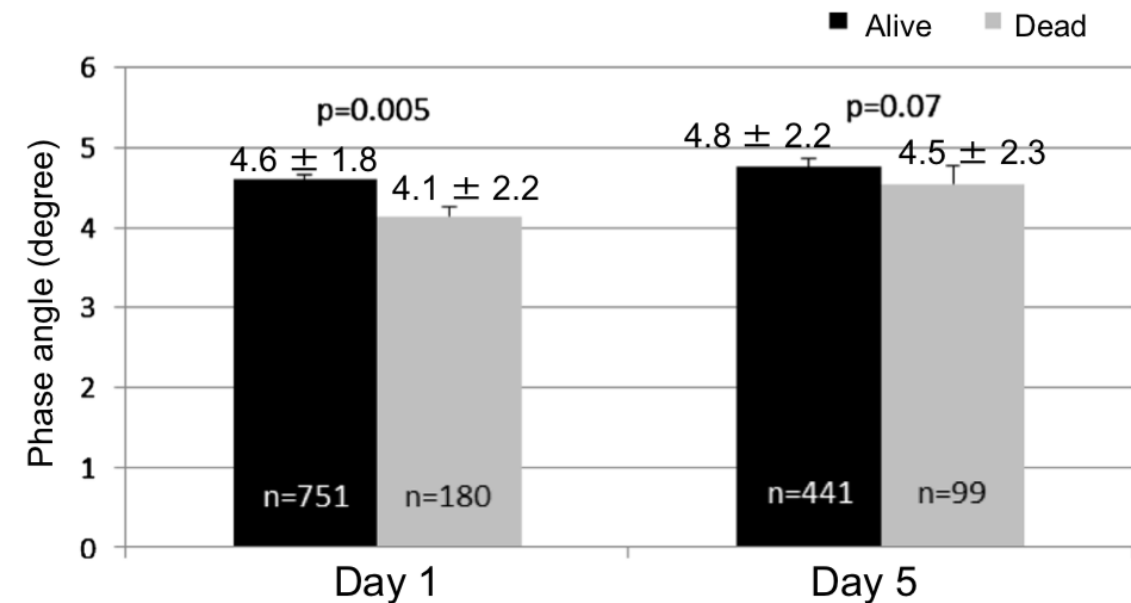
Fig. 3. Nutritional status of patients from ICU admission to 3 months post-ICU discharge. Nutritional status was at ICU admission, ICU discharge, one month (M1) and three months (M3) following ICU discharge was determined by body mass index and (kg/m^2) and weight loss (%) criteria. At M3, nutritional status was additionally determined by body composition and muscle function criteria for patients with on-site visit (M3*). Abbreviations: ICU: Intensive Care Unit.

931 Patients

Table 2 Logistic regression multivariable analysis of factors associated with 28-day mortality (n = 895)

Variables	aOR ^a	95 % CI	P value
Age	1.014	1.0016–1.0271	0.03
Day 1 phase angle	0.86	0.78–0.96	0.008
APACHE II	1.08	1.06–1.11	<0.001
Admission diagnosis: surgery vs. medicine	0.51	0.33–0.79	0.002
Other diagnosis	0.39	0.21–0.72	0.003

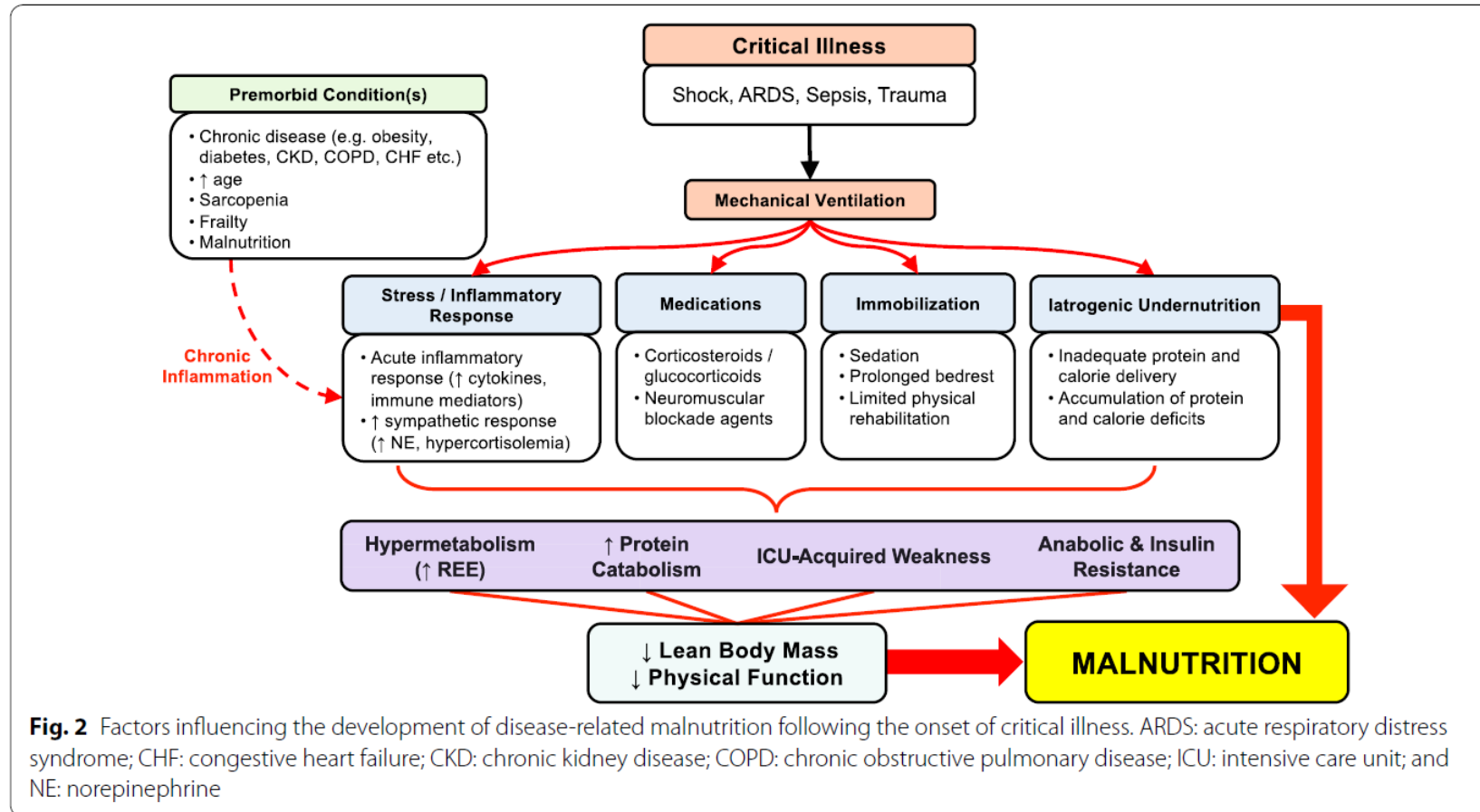
^a For continuous variables (age, day 1 phase angle and APACHE II), the adjusted odds ratio (aOR) is expressed for a 1-point increase



The role of nutrition rehabilitation in the recovery of survivors of critical illness: underrecognized and underappreciated

Lesley L. Moisey^{1*}, Judith L. Merriweather² and John W. Drover³

Moisey et al. *Critical Care* (2022) 26:270



Metabolic aspects of muscle wasting during critical illness

van Gassel RJJ

Curr Opin Clin Nutr Metab Care 2020 Mar;23(2):96-101

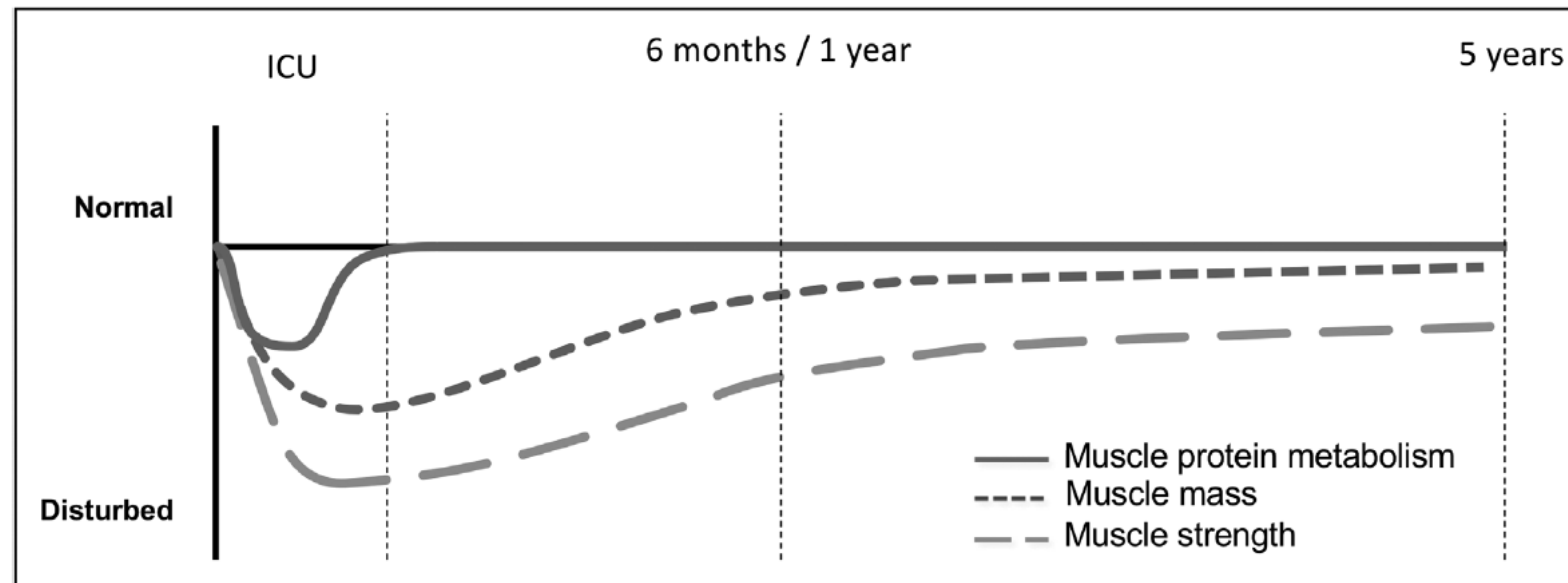
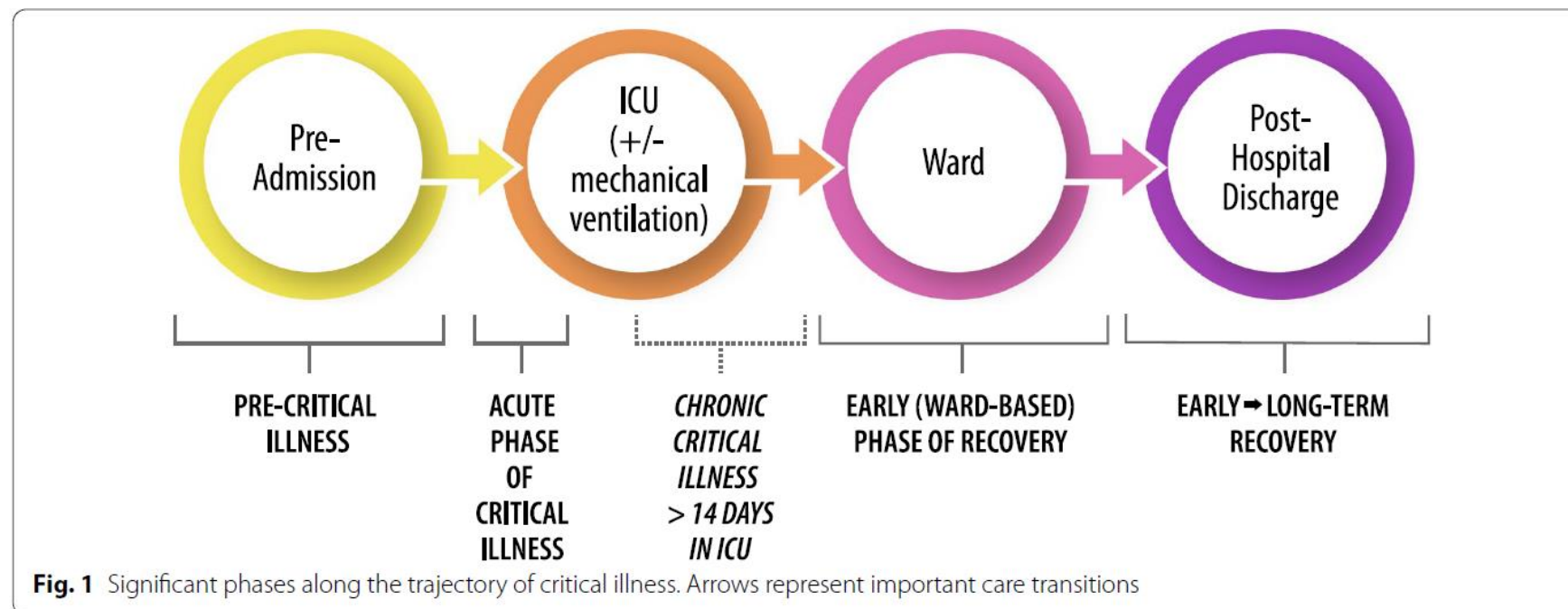


FIGURE 1. Protein metabolism in critical illness and impact on muscle mass and function. Changes over time in muscle protein metabolism, mass and function, distinguishes a short catabolic phase during ICU stay with subsequent persistent impact on muscle mass and function. Put together, this timeline further emphasizes the long-lasting impact of the relative short period of disturbed protein metabolism.

The role of nutrition rehabilitation in the recovery of survivors of critical illness: underrecognized and underappreciated

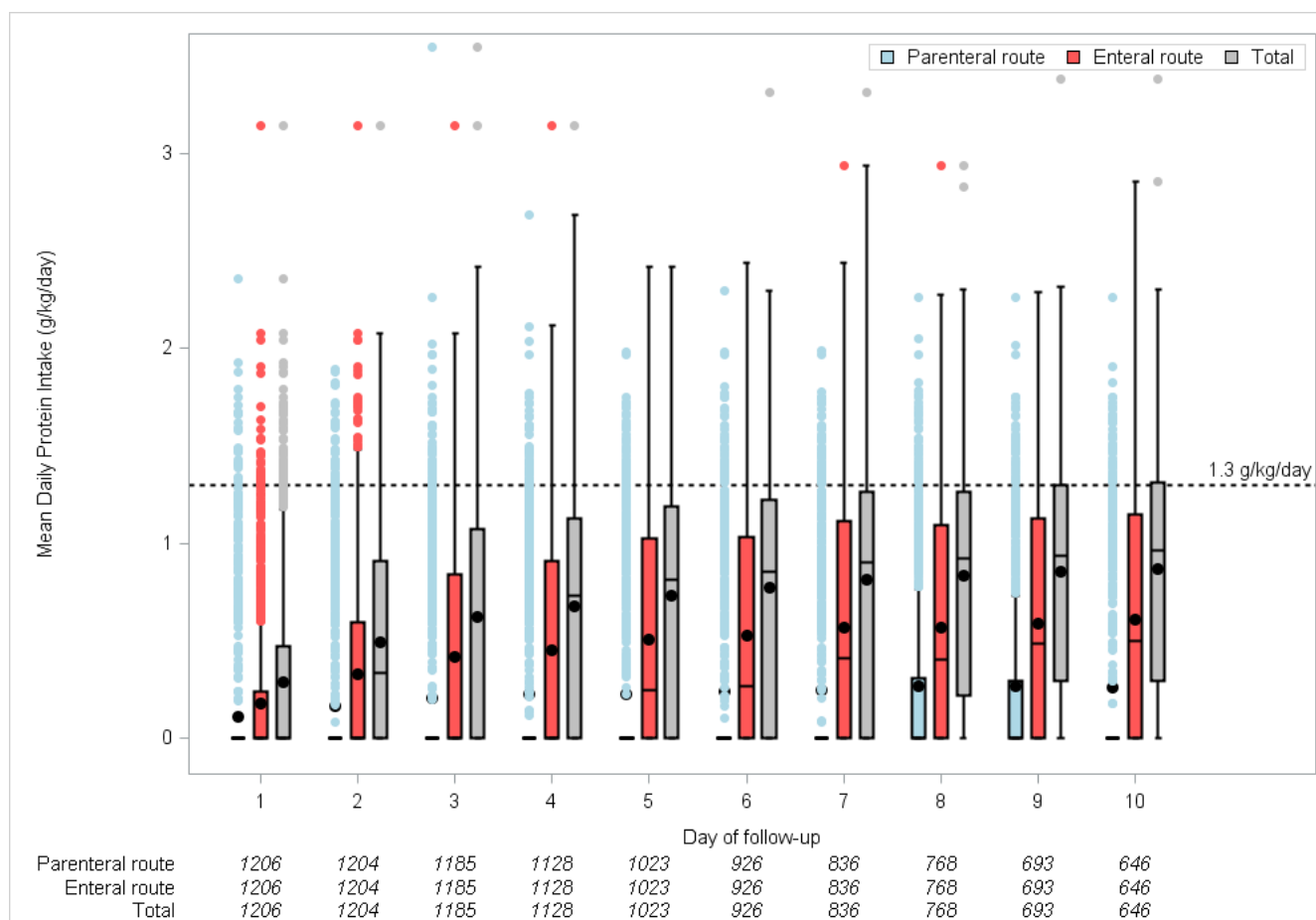
Lesley L. Moisey^{1*}, Judith L. Merriweather² and John W. Drover³

Moisey et al. *Critical Care* (2022) 26:270



Enquête binationale de pratique nutritionnelle lors d'un séjour prolongé en soins intensifs : l'étude FRANS (FRench-speAking intensive care unit Nutritional Survey).

Apports protéiques



Per os – première semaine suivant l'admission

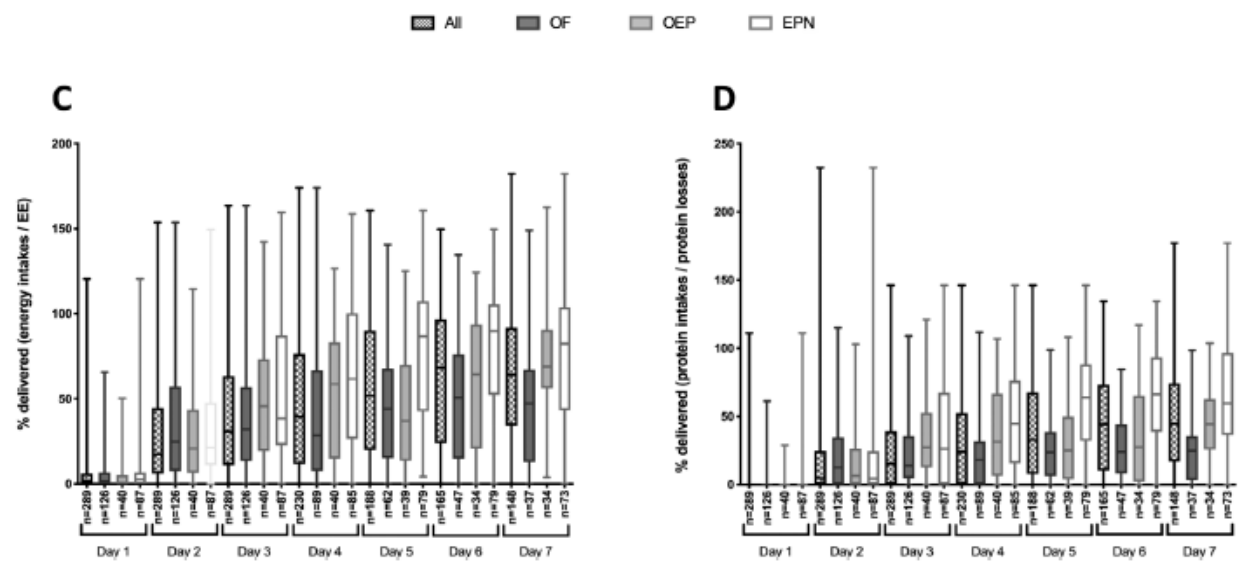
- Mixed ICU (LOS ≥ 3 j) N = 289 patients

Apports oraux étaient plus faibles comparés aux autres supports nutritionnels et déficit plus important pour les protéines

Nutrition During Critical Care: An Audit on Actual Energy and Protein Intakes

Loïc Rougier, MD¹; Jean-Charles Preiser, PhD² ; Marjorie Fateur, RD³; Anne-Marie Verbrugge, RD³; Nicolas Paquot, PhD^{3,4}; Didier Ledoux, PhD^{3,5}; Benoit Misset, MD⁵; and Anne-Françoise Rousseau, PhD^{3,5} 

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xxxx 2020 1–10
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DOI: 10.1002/jpen.1962
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Apports en Kcalorie/protéines moyens /jour

	Kcal/kg/j	g/kg/j
Per os seul	9,7	0,35
Per os+ NE/PN	15,1	0,57
NE et/ou PN seul	17,60	0,81

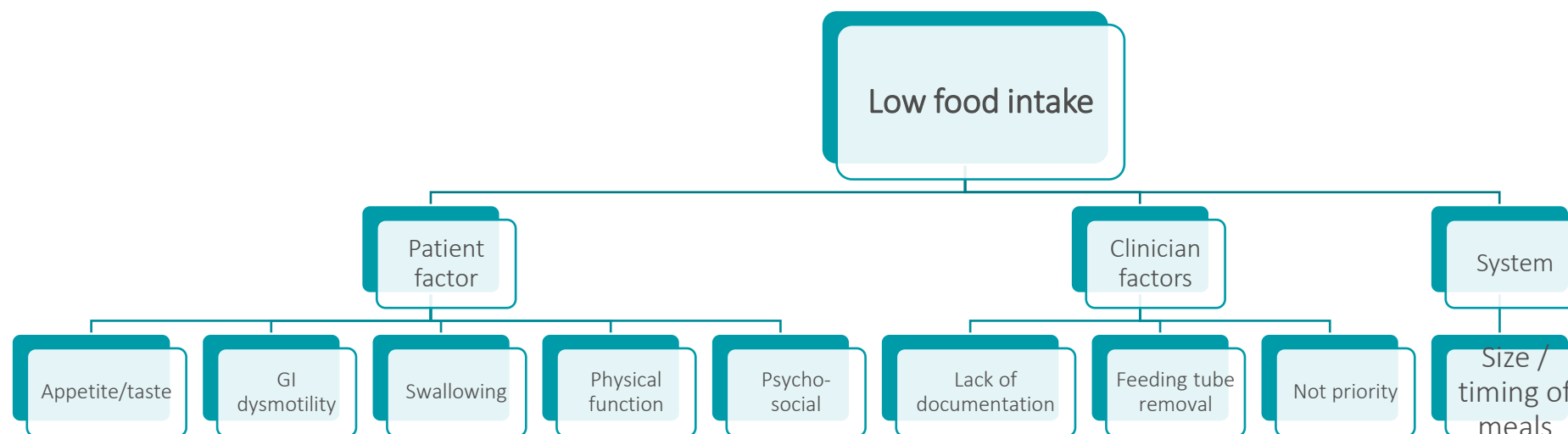
Figure 2. Daily energy (A) or protein (B) intakes and daily percentage of delivered energy (C) or protein (D) during the first 7 days, in the whole cohort and in the 3 groups. EPN, enteral/parenteral nutrition; OEP, oral + enteral/parenteral; OF, orally fed.



Nutrition intake in the post-ICU hospitalization period

Emma J. Ridley^{a,b}, Lee-anne S. Chapple^{c,d}, and Marianne J. Chapman^{c,d}

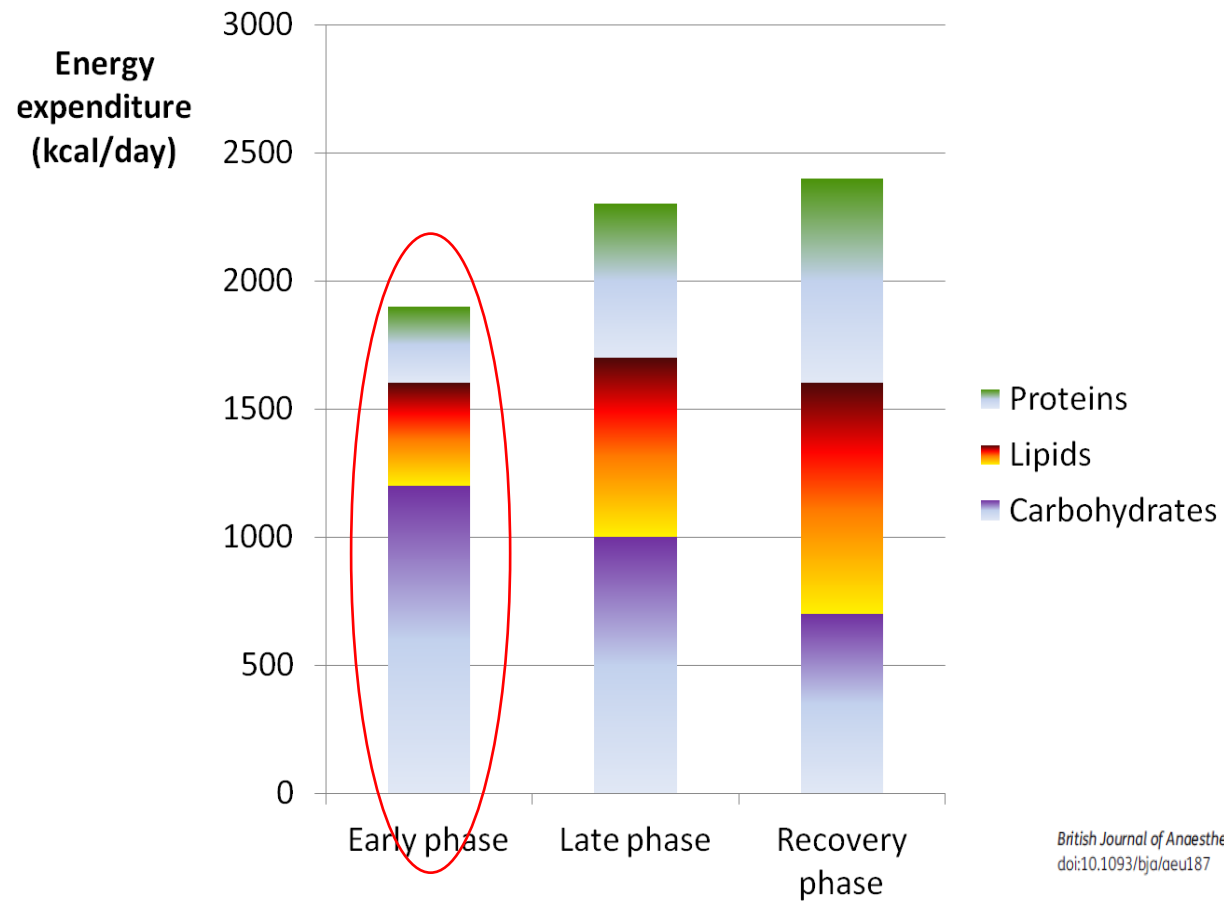
Curr Opin Clin Nutr Metab Care 2020, 23:111–115



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The 3 post-injury phases



British Journal of Anaesthesia Page 1 of 10
doi:10.1093/bja/aeu187

BJA

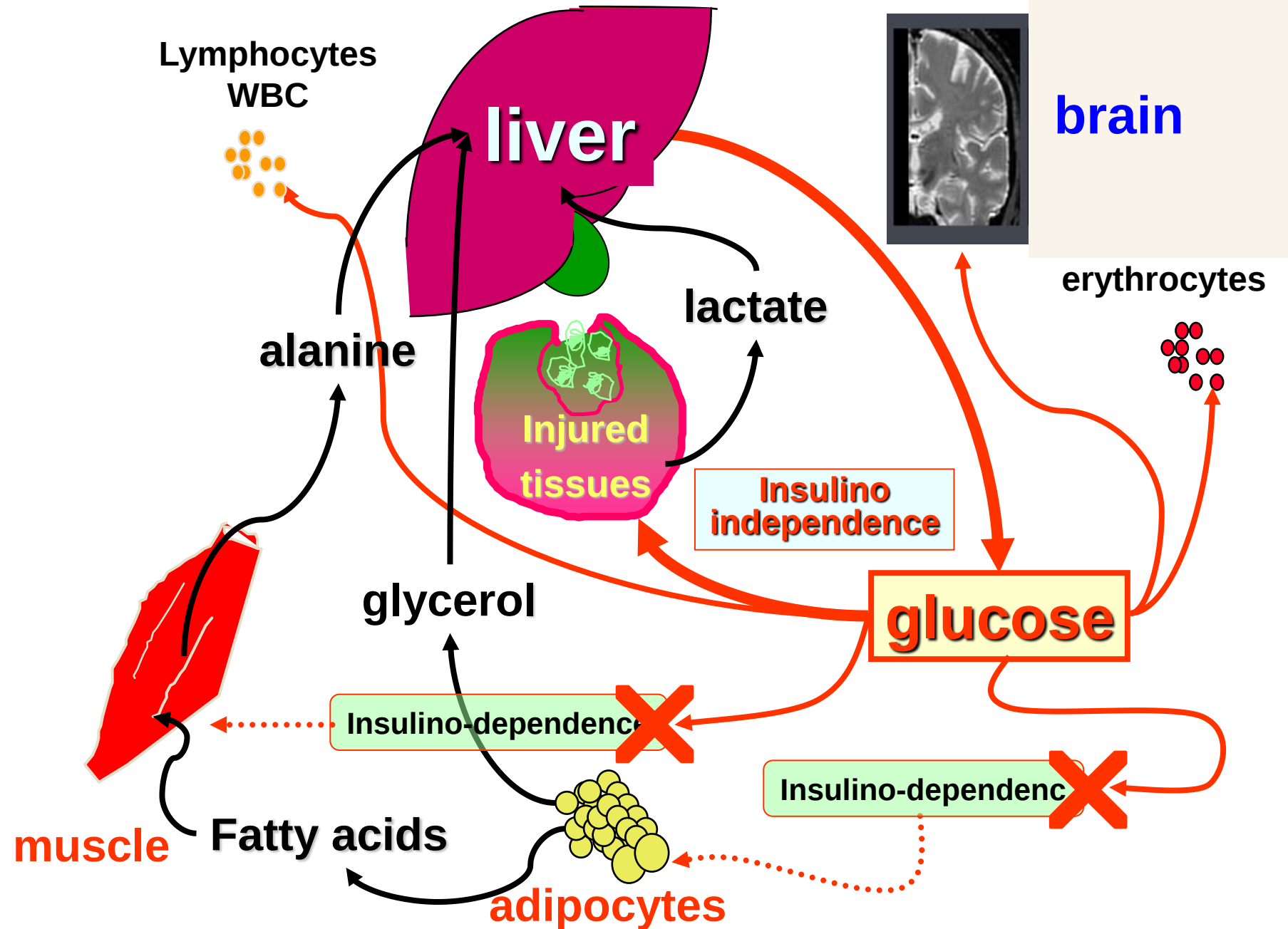
Metabolic response to the stress of critical illness

Preiser, Ichai, Orban, Groeneveld

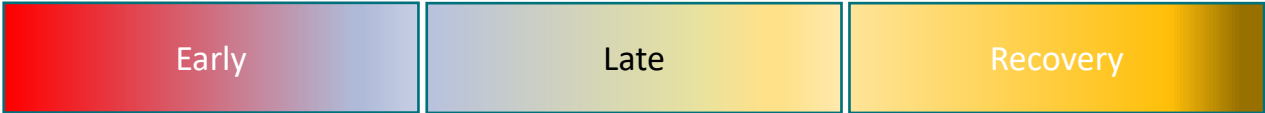
Phase aiguë / précoce Quelques heures à 3-7 jours..

- Anorexie
- Dépense énergétique limitée
- Utilisation préférentielle du glucose comme substrat énergétique

METABOLIC ADAPTATION TO STRESS



Institution	Localisation du patient			
Hôpital A	USI		USI	Salle
Hôpital B	USI	Middle care	Salle	Revali/Domicile



Critical Care Is a Concept, Not a Location*

KEYWORDS: COVID-19; high-flow nasal oxygen; hypoxemia; resource utilization; respiratory failure

Vikramjit Mukherjee, MD¹
Ryan C. Maves, MD²

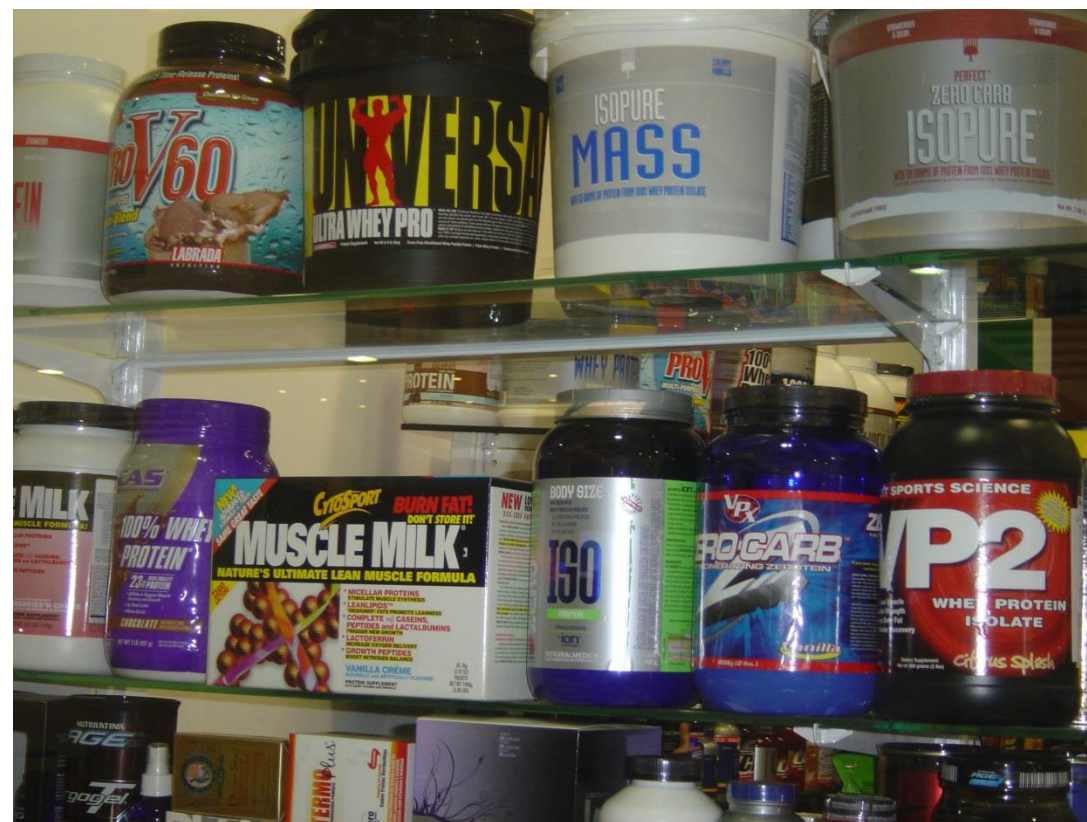
www.ccmjournal.org

January 2024 • Volume 52 • Number 1

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Combien de protéines ?



‘High protein intake during the early phase of critical illness: yes or no?’

JC Preiser, Crit Care 2018 .

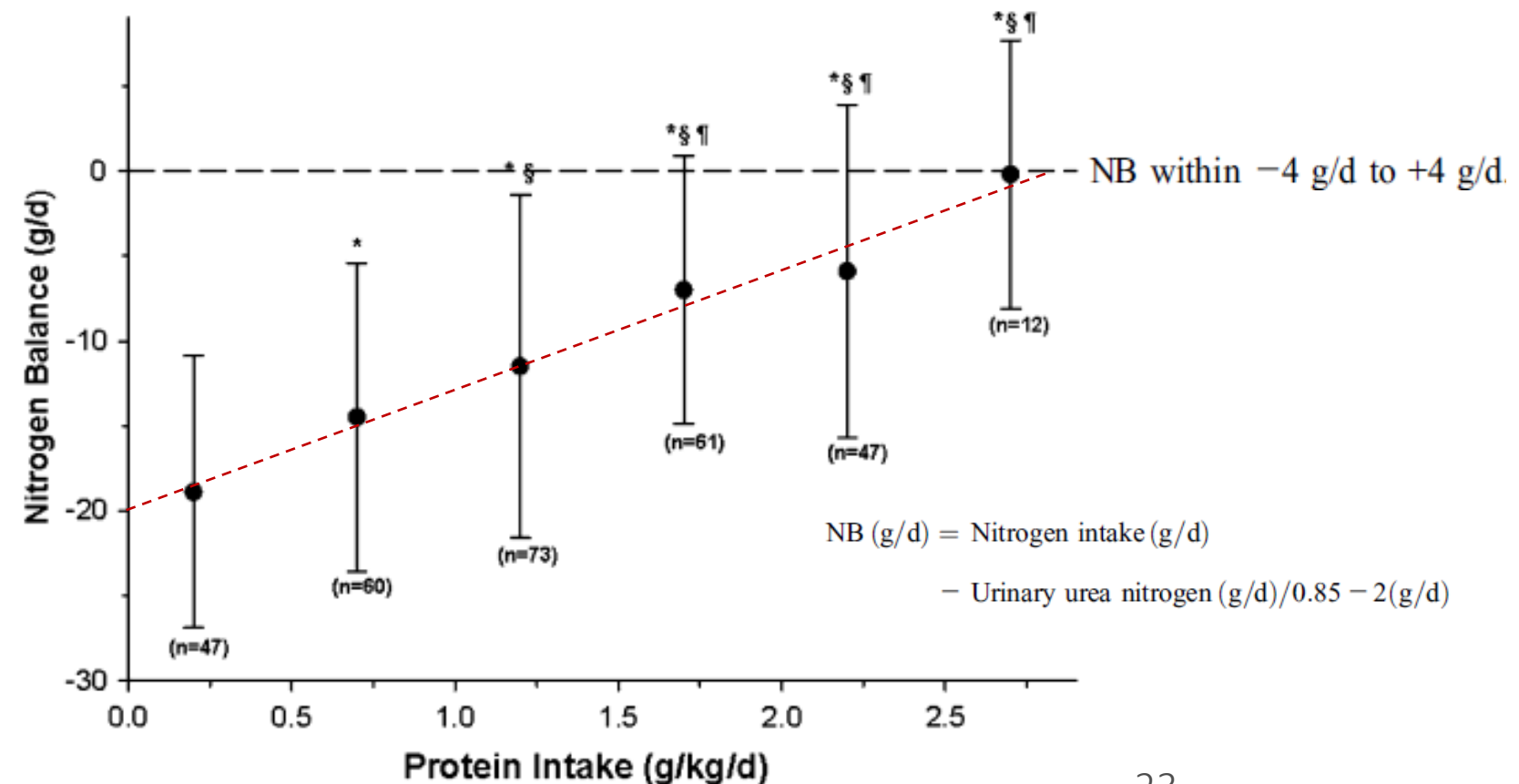
- Increase muscle protein synthesis
- Easily absorbed
- IV infusion safe

PRO

- Increases ureagenesis and oxidation of AA
- No effect on muscle protein breakdown
- Fuel auto-cannibalism
- Glucagon release

CON

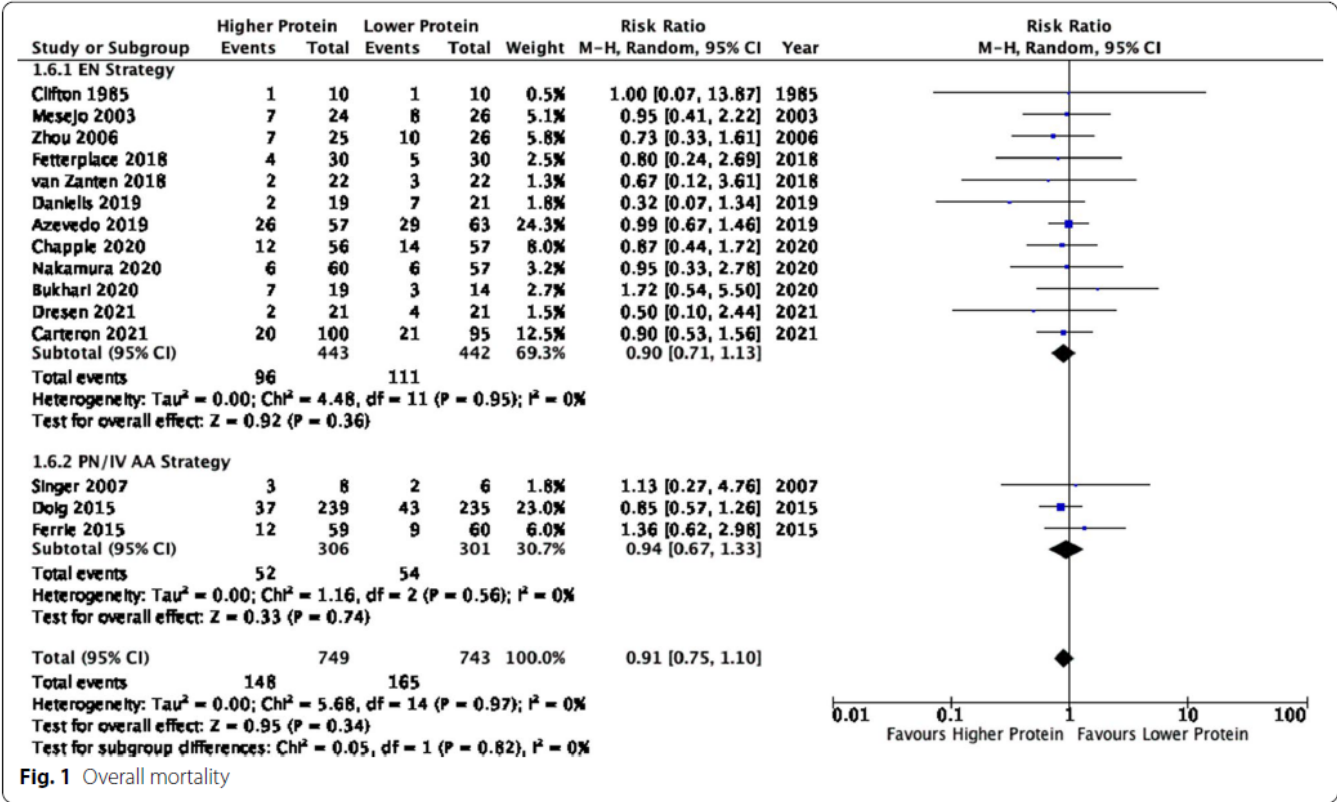
A reappraisal of nitrogen requirements for patients with critical illness and trauma



The effect of higher versus lower protein delivery in critically ill patients: a systematic review and meta-analysis of randomized controlled trials

Lee et al. *Crit Care* (2021) 25:260
<https://doi.org/10.1186/s13054-021-03693-4>

Zheng-Yii Lee¹, Cindy Sing Ling Yap¹, M. Shahnaz Hasan¹, Julia Patrick Engkasan², Mohd Yusof Barakatun-Nisak^{3,6}, Andrew G. Day⁴, Jayshil J. Patel⁵ and Daren K. Heyland^{4*}

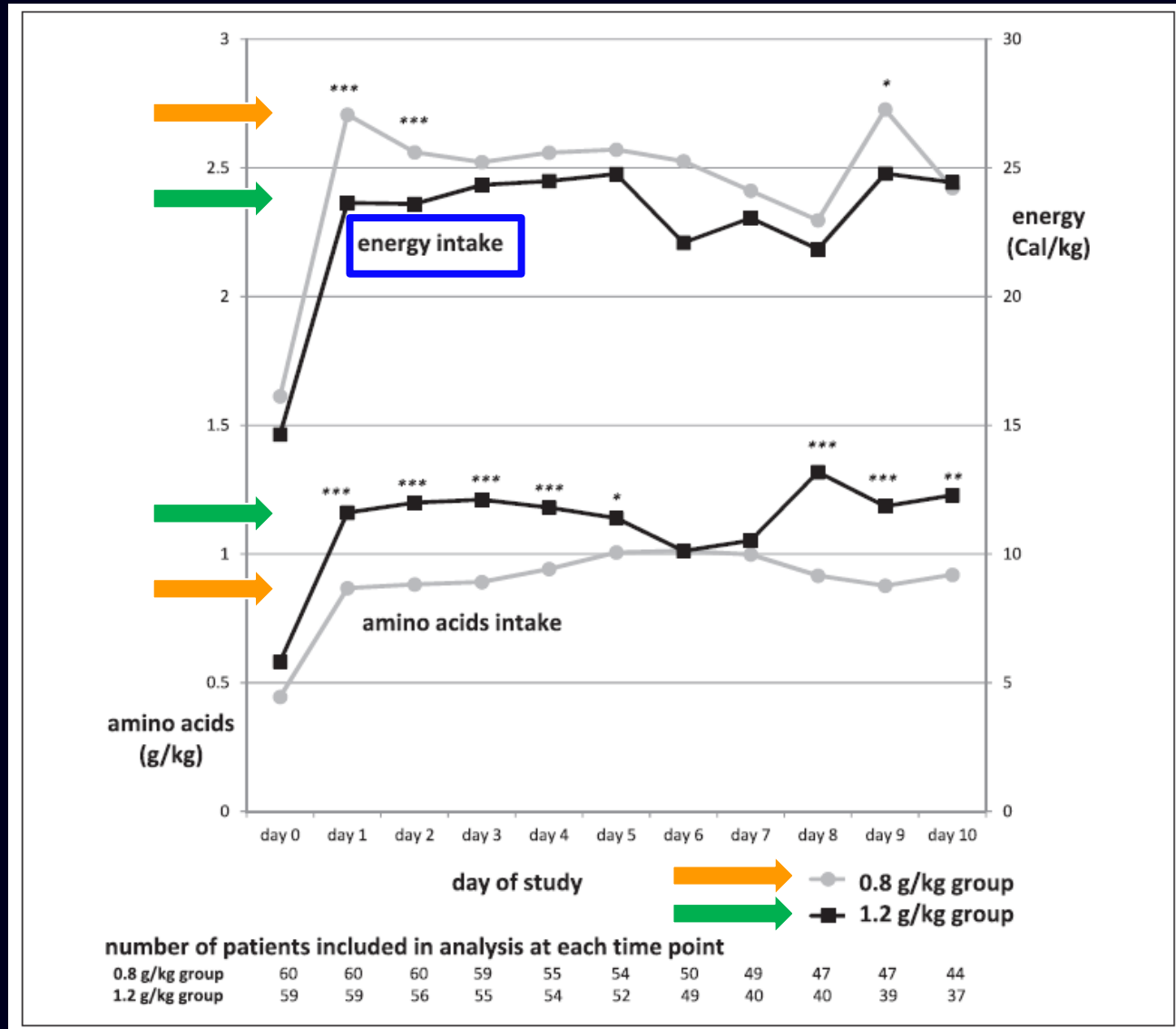


RCT in 119 critically ill patients Requiring PN (ICU day 1 – 2, mechanically ventilated, Apache 24,...)

Amino Acid Rich PN
(1,2 g/kg)
(energy poor)

Versus

Amino Acid Poor PN
(0,8 g/kg)



© Bol.com

RCT in 119 critically ill patients Requiring PN (ICU day 1 – 2, mechanically ventilated, Apache 24,...)

Table 4. Intention-to-Treat Analysis Comparing Outcomes (0.8 g/kg vs 1.2 g/kg Amino Acids).

Outcome Measures	0.8 g/kg Amino Acids (n = 60)	1.2 g/kg Amino Acids (n = 59)	P Value Between Groups
Handgrip strength on discharge from ICU, mean (SD), kg	15.8 (10.3)	18.5 (10.4)	← .054
% Expected value	45	51	
Handgrip strength at study day 7, mean (SD), kg	18.5 (11.8)	22.1 (10.1)	← .025*
% Expected value	52	62	
Nitrogen balance at study day 3, mean (SD), g/d	-5.6 (1.8)	-0.5 (3.1)	<.0001***
Nitrogen balance at study day 7, mean (SD), g/d	-4.7 (2.4)	-4.9 (7.9)	.92
ICU length of stay, median (Q1–Q3), d	6.0 (3.8–10.0)	5.0 (3.0–8.0)	.16
Hospital length of stay, median (Q1–Q3), d	27.5 (18.8–55.8)	25.0 (16.8–41.3)	.41
Dialysis days, median (Q1–Q3)	6.0 (5.3–7.0)	7.0 (2.0–8.8)	.87
Ventilator days, median (Q1–Q3)	2.0 (1.0–5.0)	2.0 (1.0–3.0)	.22
ICU mortality, No. (%)	→ 6 (10)	8 (14)	.55
Hospital mortality, No. (%)	→ 9 (15)	12 (20)	.47
6-month mortality, No. (%)	9 (15)	15 (25)	.17

ICU, intensive care unit; Q1, quartile 1; Q3, quartile 3; SD, standard deviation.

* $P < .05$. ** $P < .01$. *** $P < .001$.



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Early goal-directed nutrition versus standard of care in adult intensive care patients: the single-centre, randomised, outcome assessor-blinded EAT-ICU trial

Matilde Jo Allingstrup¹, Jens Kondrup², Jørgen Wiis¹, Casper Claudius¹, Ulf Gøttrup Pedersen¹,
Rikke Hein-Rasmussen¹, Mads Rye Bjerregaard¹, Morten Steensen¹, Tom Hartvig Jensen¹, Theis Lange^{3,4},
Martin Bruun Madsen¹, Morten Hylander Møller¹ and Anders Perner^{1*} 

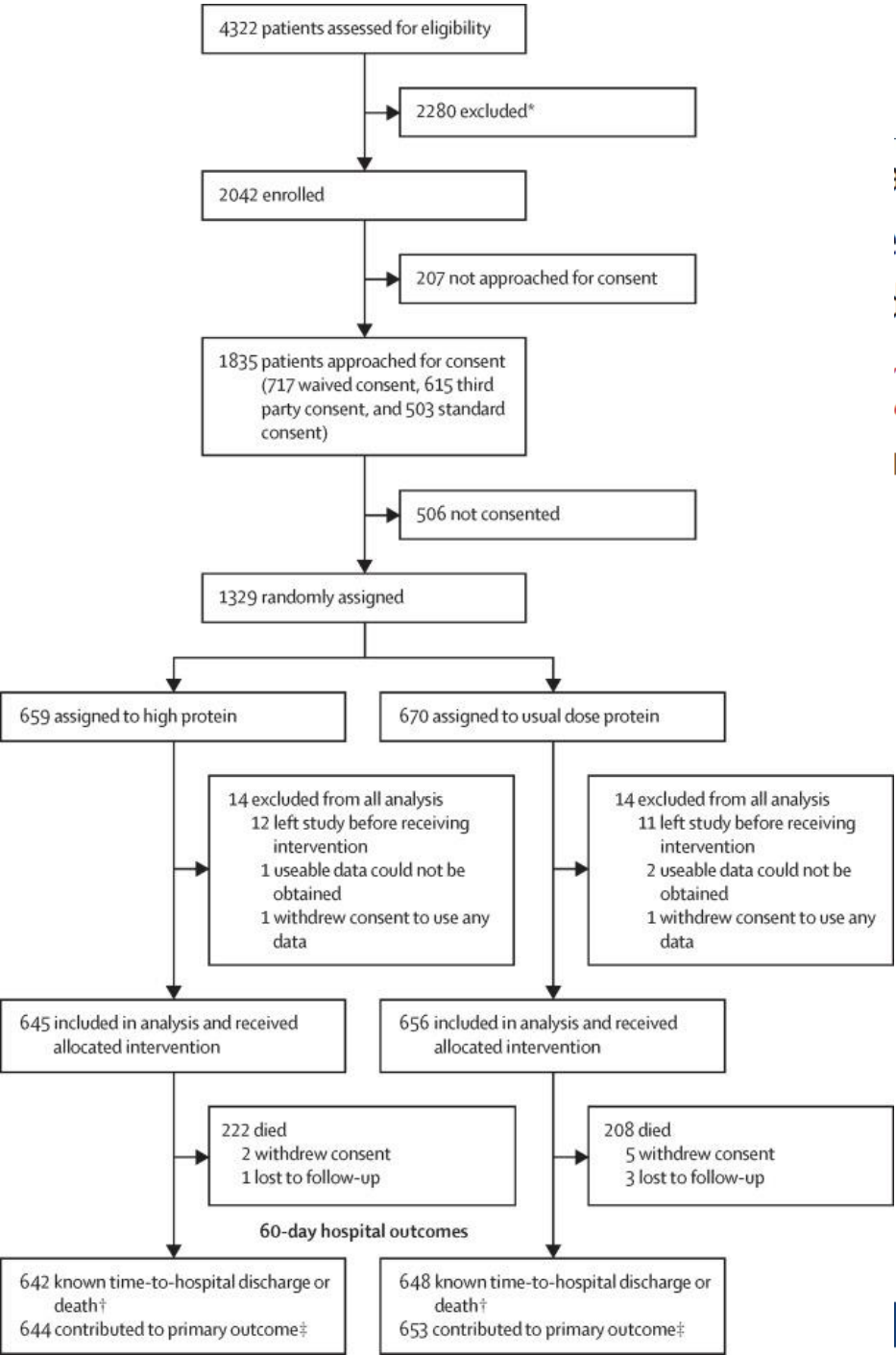
Intensive Care Med (2017) 43:1637–1647

Table 2. Nutrition characteristics in ICU after randomisation ^a

Variable	Early Goal-directed Nutrition (N=100)	Standard of Care (N=99)
Measured ^b energy requirement, kcal/day	2069 (1816 - 2380)	1887 (1674 - 2244)
Calculated ^c energy requirement, kcal/day	1950 (1750 - 2125)	1875 (1650 - 2100)
Energy intake, kcal/day	1877 (1567 - 2254)	1061 (745 - 1470)
Energy balance ^d , kcal/day	-66 (-157 - -6)	-787 (-1223 - -333)
Measured ^e protein requirement, g/kg/day	1.63 (1.36 - 2.05)	1.16 (0.89 - 1.62)
Protein intake, g/kg/day	1.47 (1.13 - 1.69)	0.50 (0.29 - 0.69)
Protein balance ^d , g/kg/day	-0.28 (-0.76 - 0.11)	-0.69 (-1.02 - -0.38)
P-urea, mmol/l	13.5 (8.7 – 21.9)	9.0 (5.6 – 14.4)
24-hour urinary urea, mmol/day	516 (368 – 760)	320 (175 – 482)

The effect of high protein with high multicenter

Daren K Heyland, Ja
Courtney Wedemire
EFFORT Protein Trial



Patients International, trial

'Reilly, Racquel Regala,
e

hed online January 25, 2023

The effect of higher protein dosing in critically ill patients with high nutritional risk (EFFORT Protein): an international, multicentre, pragmatic, registry-based randomised trial

Daren K Heyland, Jayshil Patel, Charlene Compher, Todd W Rice, Danielle E Bear, Zheng-Yii Lee, Victoria C González, Kevin O'Reilly, Racquel Regala, Courtney Wedemire, Miguel Ibarra-Estrada, Christian Stoppe, Luis Ortiz-Reyes, Xuran Jiang, Andrew G Day, on behalf of the EFFORT Protein Trial team

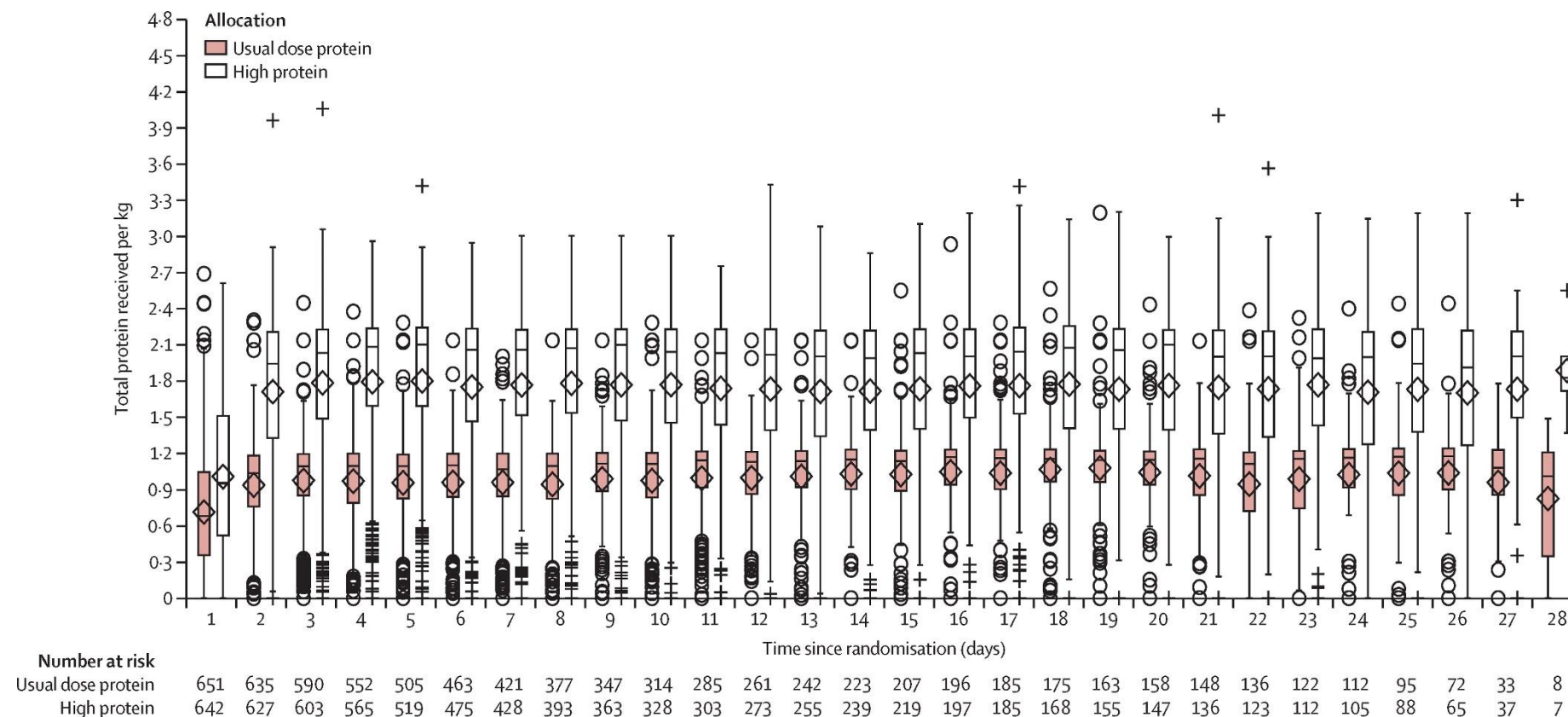
www.thelancet.com Published online January 25, 2023

	High protein (n=645)	Usual dose protein (n=656)
Age, years	57 (17 [18–95])	57 (17 [18–93])
Sex		
Male	395 (61%)	388 (59%)
Female	250 (39%)	267 (41%)
Admission category		
Medical	548 (85%)	540 (82%)
Surgical elective	24 (4%)	19 (3%)
Surgical emergency	73 (11%)	97 (15%)

The effect of higher protein dosing in critically ill patients with high nutritional risk (EFFORT Protein): an international, multicentre, pragmatic, registry-based randomised trial

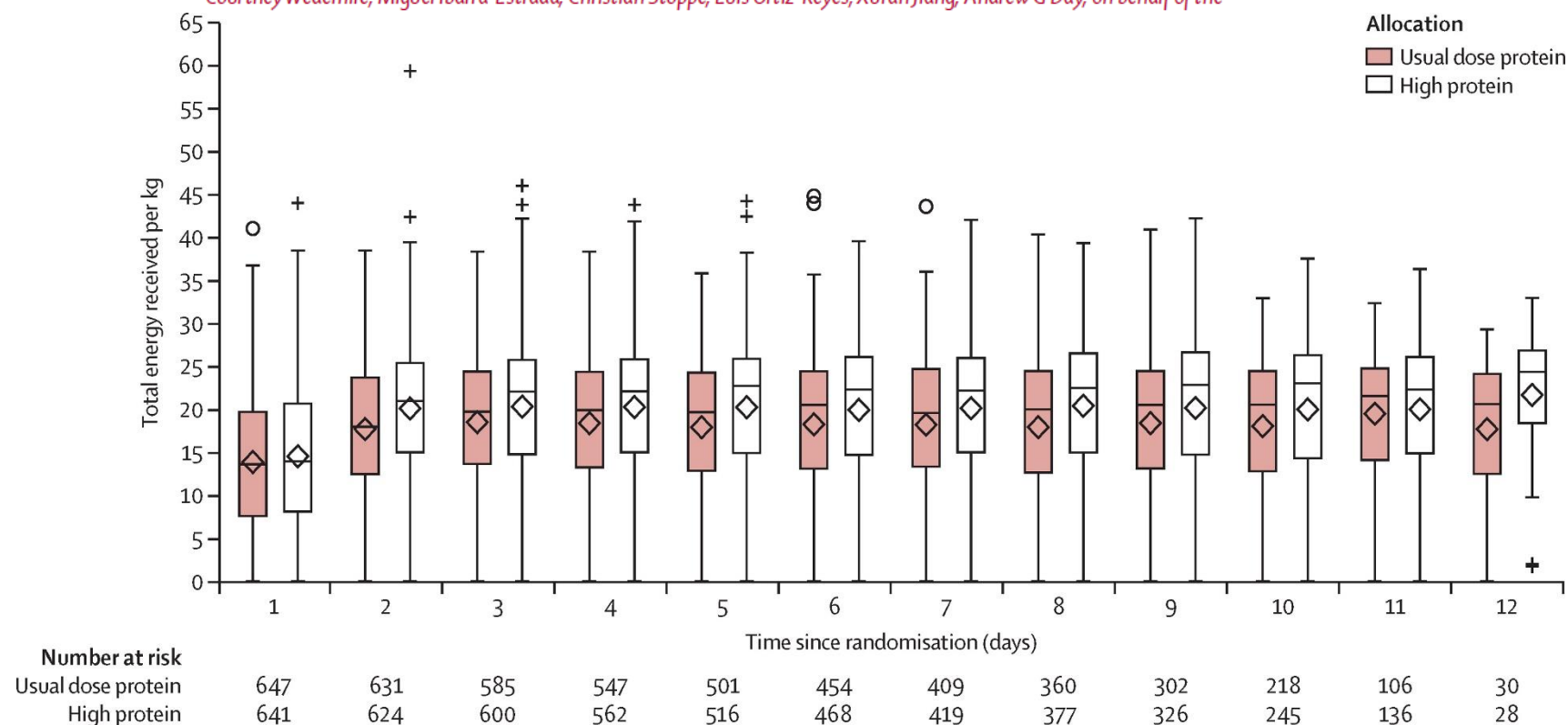
Daren K Heyland, Jayshil Patel, Charlene Compher, Todd W Rice, Danielle E Bear, Zheng-Yii Lee, Victoria C González, Kevin O'Reilly, Racquel Regala, Courtney Wedemire, Miguel Ibarra-Estrada, Christian Stoppe, Luis Ortiz-Reyes, Xuran Jiang, Andrew G Day, on behalf of the EFFORT Protein Trial team

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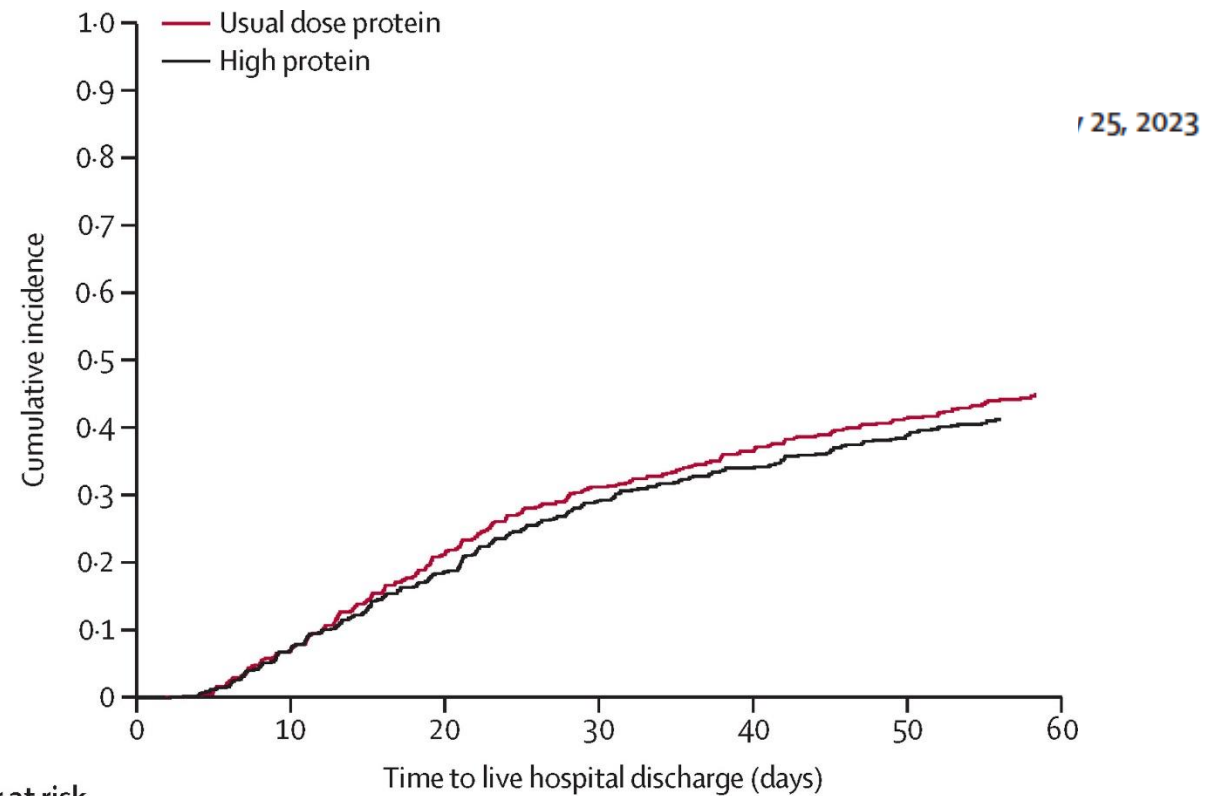


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The effect of higher protein dosing in critically ill patients with high nutritional risk (EFFORT Protein): an international, multicentre, pragmatic, registry-based randomised trial



Number at risk		Time to live hospital discharge (days)					
		0	10	20	30	40	50
Deaths (high dose protein)	..	95	162	184	207	218	222
Deaths (usual protein)	..	91	142	175	185	198	208
Hospitalisation (high protein)	644	499	353	261	204	165	144
Hospitalisation (usual dose protein)	653	513	364	261	216	175	145

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Daren K Heyland, Jayshil Patel, Charlene Compher, Todd W Rice, Danielle E Bear, Zheng-Yii Lee, Victoria C González, Kevin O'Reilly, Racquel Regala, Courtney Wedemire, Miguel Ibarra-Estrada, Christian Stoppe, Luis Ortiz-Reyes, Xuran Jiang, Andrew G Day, on behalf of the EFFORT Protein Trial team

www.thelancet.com Published online January 25, 2023

Table S7 – Metabolic tolerance

	High Protein (n=645)	Usual Protein (n=656)	p-value*
Insulin (total units/day)	(N=364) 44.9±48.2 (1.0-379.0)	(N=365) 40.4±41.1 (1.0-265.0)	0.39
Creatinine (μmol/L)	(N=641) 125.8±121.2 (17.8-1589.2)	(N=654) 117.5±115.5 (17.7-1442.0)	0.42
Highest Glucose (mmol/L)	(N=598) 10.3±3.0 (5.5-24.4)	(N=597) 10.4±2.9 (3.7-28.7)	0.39
Lowest Glucose (mmol/L)	(N=598) 7.2±1.5 (3.9-13.7)	(N=597) 7.1±1.3 (3.7-12.8)	0.07
Highest Urea (mmol/L)	(N=634) 14.0±8.5 (2.4-50.7)	(N=647) 11.9±7.2 (1.8-47.7)	<0.0001
Urea ≥30 mmol/L	43 (6.7%)	20 (3.0%)	0.002
Highest Triglycerides (mmol/L)	(N=160) 2.2±1.4 (0.3-7.9)	(N=175) 2.5±2.0 (0.4-16.8)	0.28

Legend: Values reflect the average ± standard deviation (range) of the within patient averages over the study period.

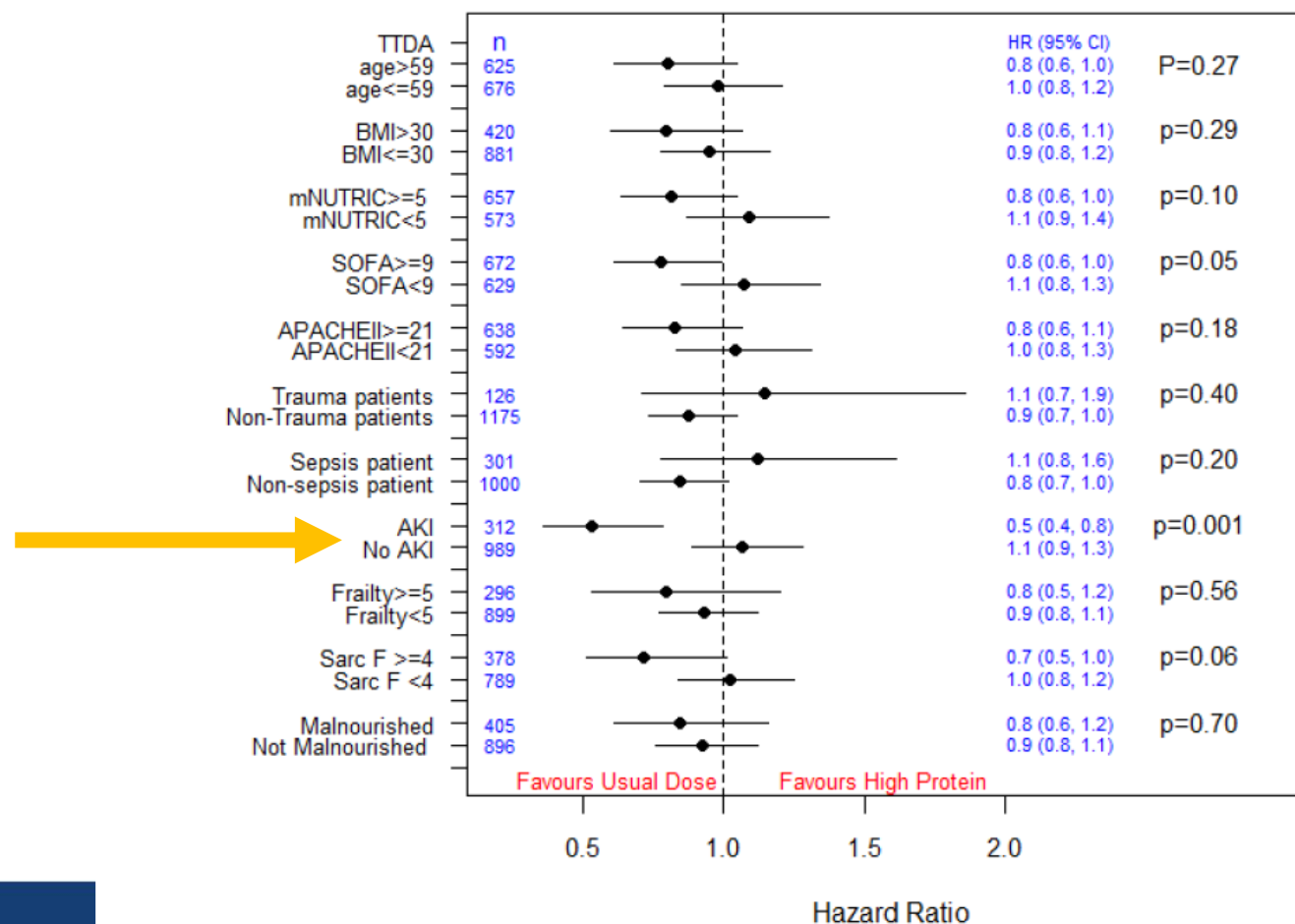
*All p-values unadjusted and by the Wilcoxon rank-sum test except Urea ≥30 mmol/L by the chi-squared test.

The effect of higher protein dosing in critically ill patients with high nutritional risk (EFFORT Protein): an international, multicentre, pragmatic, registry-based randomised trial

Figure S2 – Subgroup Analysis Based on Effect on Time-to-Discharge-Alive from Hospital

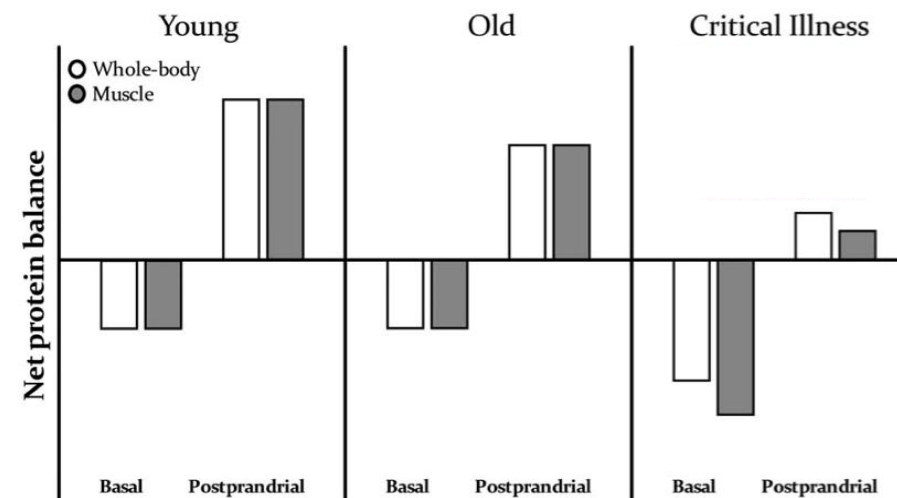
Regala,

January 25, 2023



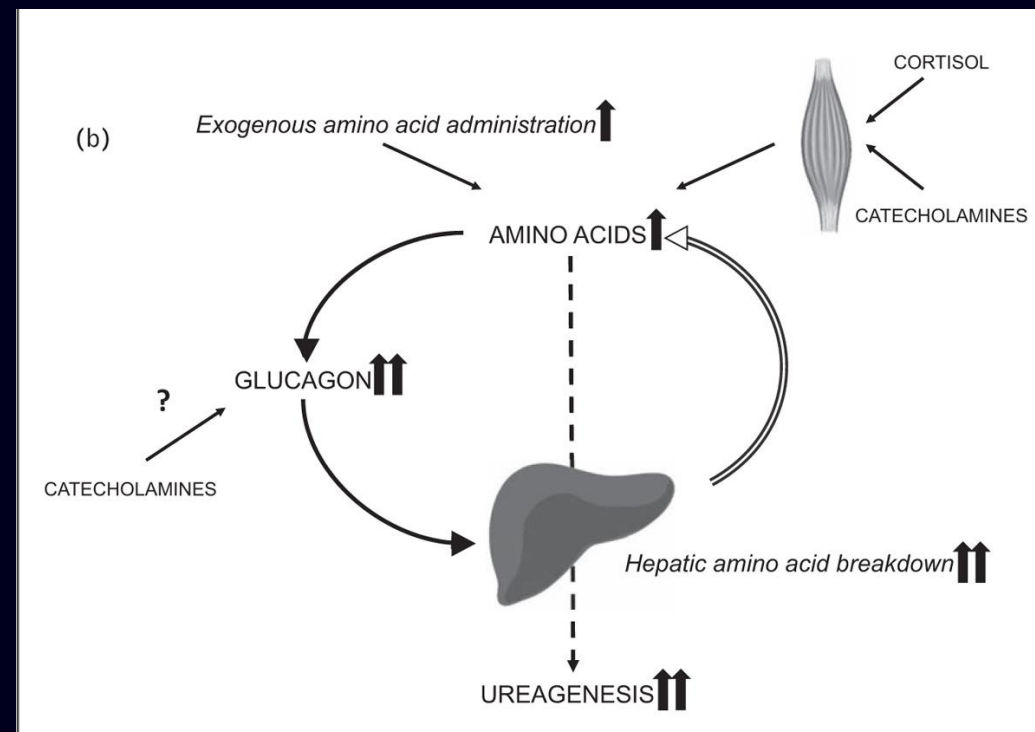
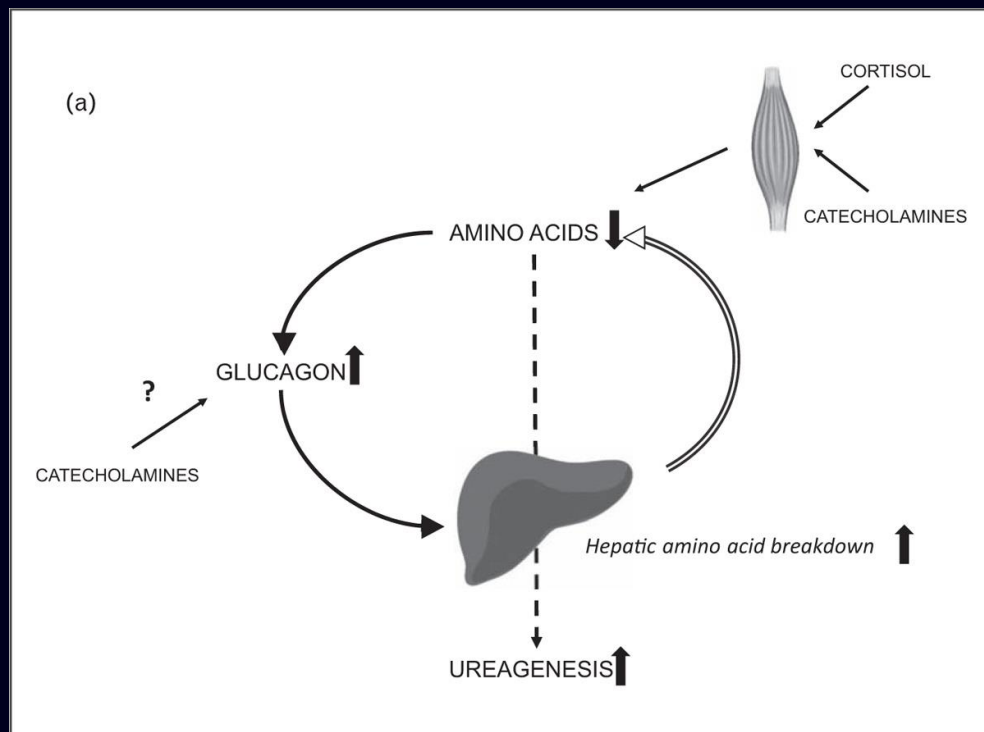
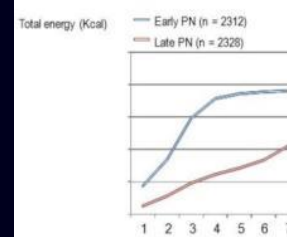
Critical Illness is Associated With Anabolic Resistance

Morton Current Opinion in Critical Care: April 2018 - Volume
24 - Issue 2 - p 124-130





What drives protein breakdown?

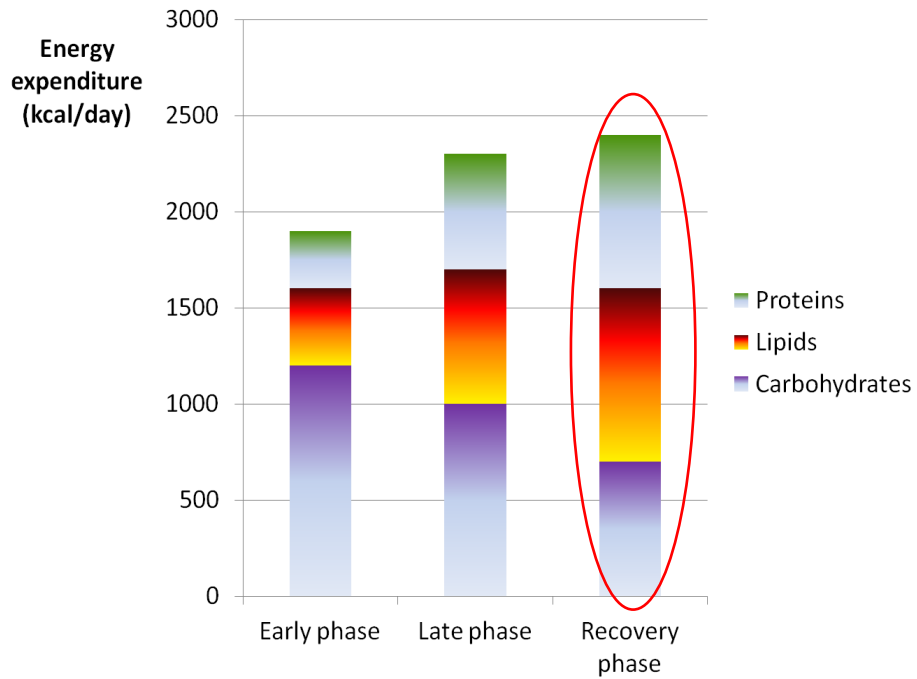


+/- Amino Acid Infusion

+/- Glucagon Immuno-neutralisation

Thiessen S. et al AJRCM 2017
Thiessen S. & Gunst J. Current Opinion in Critical Care 2018

Three metabolic phases after stress



British Journal of Anaesthesia Page 1 of 10
doi:10.1093/bja/oeu187

BJA

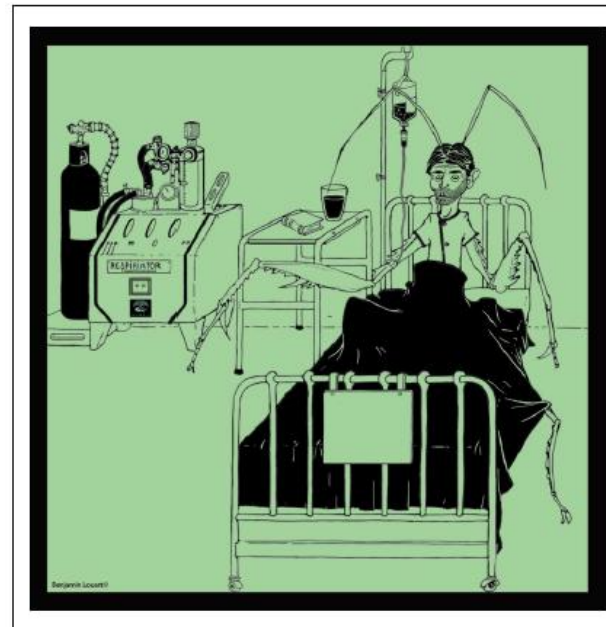
Metabolic response to the stress of critical illness

Preiser, Ichai, Orban, Groeneveld

Tutorial

Nutrition Rehabilitation in the Intensive Care Unit

Pablo Lucas Massanet, MD¹; Laurent Petit, MD²; Benjamin Louart, MD¹;
Philippe Corne, MD, PhD³; Celine Richard, RD⁴; and Jean Charles Preiser, MD, PhD⁵



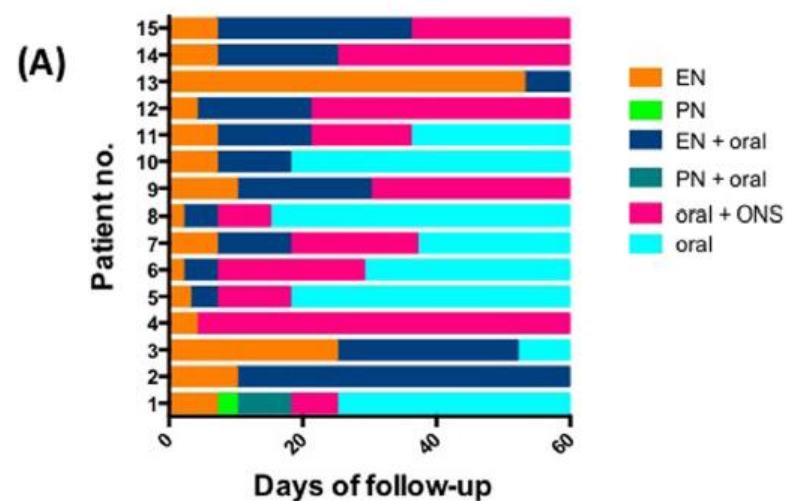
Journal of Parenteral and Enteral
Nutrition
Volume 39 Number 4
May 2015 391–400

Figure 1. Kafka's nightmare illustrates how deep the physical changes are for a patient after a critical illness and an intensive care unit stay.

Nutrition evaluation and management of critically ill patients with COVID-19 during post-intensive care rehabilitation

Alice Hoyois¹ | Asuncion Ballarin² | Justine Thomas² | Olivier Lheureux³ |
Jean-Charles Preiser³ | Emmanuel Coppens⁴ | Silvia Perez-Bogerd⁵ |
Olivier Taton⁵ | Sylvie Farine⁶ | Pauline Van Ouytsel⁶ | Marianna Arvanitakis¹

JPEN J Parenter Enteral Nutr. 2021;45:1153–1163.



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JPEN J Parenter Enteral Nutr. 2021;45:1153–1163.

(B)

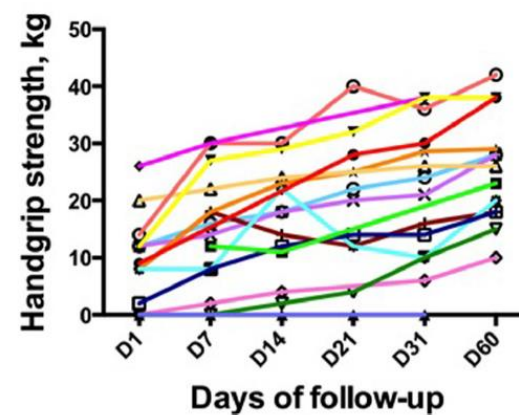


Figure 4 (A) Correlation between body mass index (BMI) and mid-arm circumference. (B) Individual dominant handgrip strength at the time of intensive care unit discharge and at day (D) 7, 14, 30, and 60 for 13 patients. Each patient is represented with a shade of gray

The role of nutrition rehabilitation in the recovery of survivors of critical illness: underrecognized and underappreciated

Lesley L. Moisey^{1*}, Judith L. Merriweather² and John W. Drover³

Moisey et al. *Critical Care* (2022) 26:270

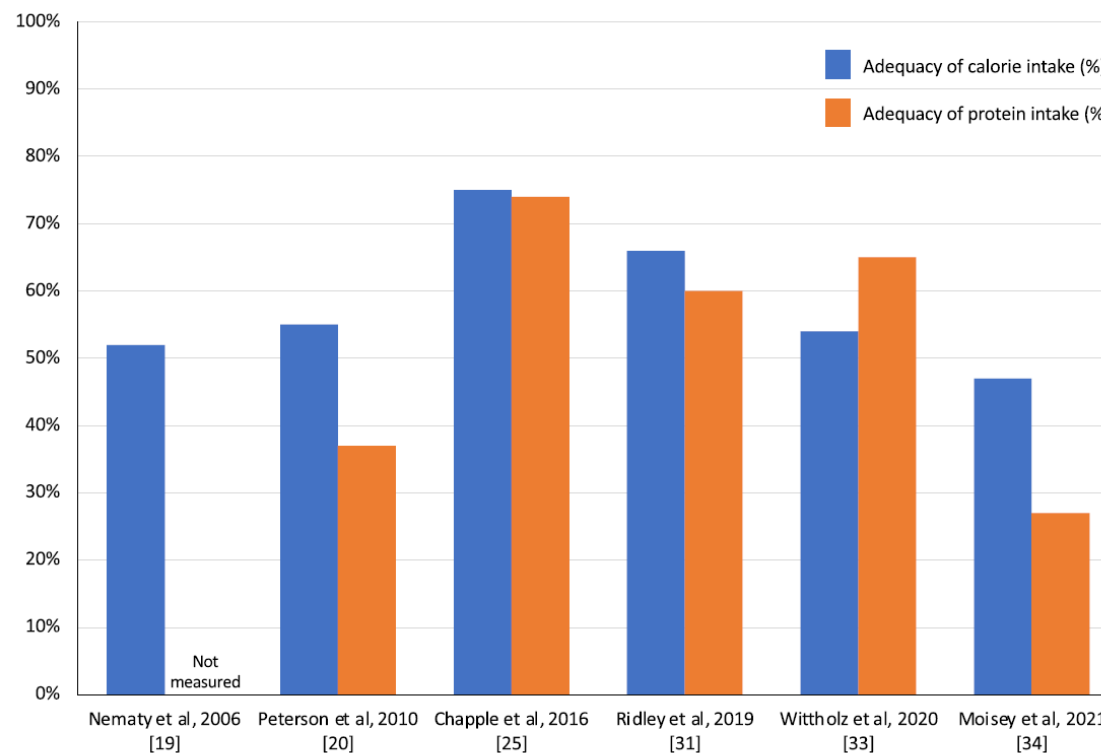
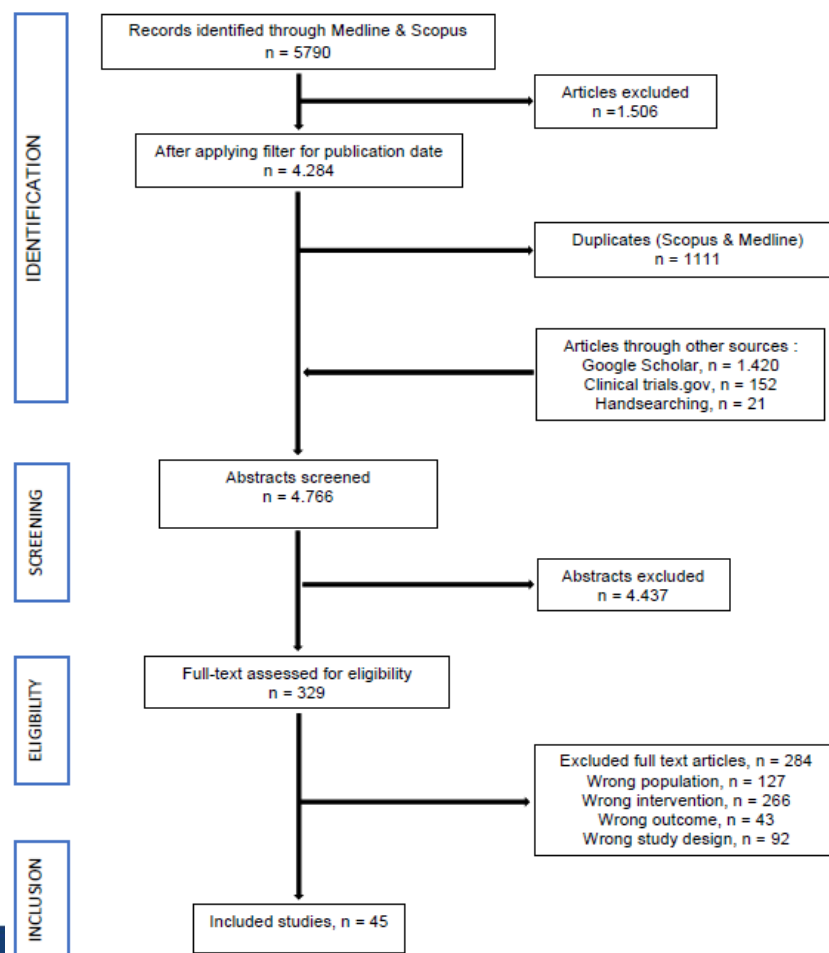


Fig. 3 Adequacy of calorie and protein intake in relation to estimated or prescribed amounts in patients solely prescribed oral diets in hospital following ICU discharge

Nutritional rehabilitation after acute illness among older patients: A systematic review and meta-analysis

Sylwia Szklarzewska^{a,*}, Raphael Mottale^b, Edgard Engelman^c, Sandra De Breucker^a,
Jean-Charles Preiser^d

Clinical Nutrition 42 (2023) 309–336



Non-overlapping data

Parameters	Hedge's g Mean [95% confidence interval]	Percent of non- overlapping data ** Mean [95% confidence interval]	Odds Ratio *** [95% confidence interval]	p-value
Function status *	0.27 [0.08 ; 0.46]	19.4 [6.2 ; 30.8]	1.63 [1.15 ; 2.3]	p=0.003
Muscle mass *	0.53 [0.11 ; 0.94]	34.6 [8.4 ; 53.1]	2.61 [1.22 ; 5.5]	p=0.01
Quality of life *	0.63 [-0.17 ; 1.44]	39.6 [-14.5 ; 69.2]	3.13 [0.73 ; 13.6]	p=0.12

* A positive value of Hedge's g indicates a better effect of food supplements compared with control

** For all parameters: the higher the percentage the better the effect of food supplements compared with control. A negative value indicates of better effect of control

Agenda

- Constats cliniques
- Physiologie
- Apports protéiques
 - **Précoces : oui/non**
 - **Tardifs : revalidation nutritionnelle**
- **Recommandations de pratique clinique**
- Futur?

How much proteins ?

REVIEW

Open Access

A guide to enteral nutrition in intensive care units: 10 expert tips for the daily practice



Jean-Charles Preiser^{1*}, Yaseen M. Arabi², Mette M. Berger³, Michael Casaer⁴, Stephen McClave⁵, Juan C. Montejó-González⁶, Sandra Peake^{7,8}, Annika Reintam Blaser^{9,10}, Greet Van den Berghe⁴, Arthur van Zanten¹¹, Jan Wernerman¹² and Paul Wischmeyer¹³

Critical Care (2021) 25:424

Suggested answer	ASPEN/SCCM guidelines	ESPEN guidelines
Low dose (e.g. 0.8 g/kg/day) during the early phase - to be increased to >1.2 g/kg/d later	Suggestion: Administer sufficient (high-dose) protein in the range of 1.2–2.0 g/kg actual body weight per day and may likely be even higher in burn or multitrauma patients [Quality of Evidence: Very Low]	During critical illness, 1.3 g/kg protein equivalents per day can be delivered progressively Grade of recommendation: 0: strong consensus

ESPEN ICU guidelines revised (2023)

During critical illness, 1.3 g/kg protein equivalents per day can be delivered progressively.

(R22, updated, Grade 0, strong consensus, 92%)

Commentary

Numerous observational studies showed associations between high-dose protein delivery and improved outcomes (morbidity or mortality).

However, RCTs did not confirm the expected benefit of higher protein intakes alone, or combined with physical activity or energy

In accordance with the Lee et al. meta-analysis, there is no support to recommend higher doses than 1.3 g/kg/d, except for increased muscle mass in individual patients.

The timing of administration has been mainly studied retrospectively or in post hoc analysis [62]. Recent stable isotope studies [63,64] demonstrate that amino acid conversion is improved with time, allowing an increase in whole body protein production only after the early period of the acute phase, even increasing further in the post-acute phase.

This supports the progressive increase of protein intakes.

There is a need for well-conducted RCTs to answer the question of the best timing and dose of protein administration in the ICU.

ESPEN ICU guidelines revised (2023)

Statement 3: Physical activity may improve the beneficial effects of nutritional therapy.

(S 3, consensus, 86%)

Commentary

Exercise has been suggested in several studies to be effective in preventing anabolic resistance, reducing morbidity and improving the level of activity.

Administration of increased protein intake together with increased physical activity during the late phase of critical illness should be further explored.

Agenda

- Constats cliniques
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- Apports protéiques
 - **Précoces : oui/non**
 - **Tardifs : revalidation nutritionnelle**
- Recommandations de pratique clinique
- Futur?

Individualised nutritional support in medical inpatients at nutritional risk: a randomised clinical trial

Philipp Schuetz, Rebecca Fehr, Valerie Baechli, Martina Geiser, Manuela Deiss, Filomena Gomes, Alexander Kutz, Pascal Tribolet, Thomas Bregenzer, Nina Braun, Claus Hoess, Vojtech Pavlicek, Sarah Schmid, Stefan Bilz, Sarah Sigrist, Michael Brändle, Carmen Benz, Christoph Henzen, Silvia Mattmann, Robert Thomann, Claudia Brand, Jonas Rutishauser, Drahomir Aujesky, Nicolas Rodondi, Jacques Donzé, Zeno Stanga, Beat Mueller**

Lancet 2019; 393: 2312-21

Nutrition risk screening (NRS 2002) within 48 h of hospital admission in all patients

If increased risk for malnutrition → individual assessment of the patient → if risk for malnutrition is present and nutritional therapy is not contraindicated → establish a strategy to achieve individual nutritional targets

Individual nutrition targets

Caloric requirements
Harris-Benedict equation
with adjusted bodyweight
or indirect calorimetry

Protein requirements
1.2–1.5 g/kg bodyweight
per day (0.8 g/kg of
bodyweight per day in
patients with renal failure
with no dialysis)

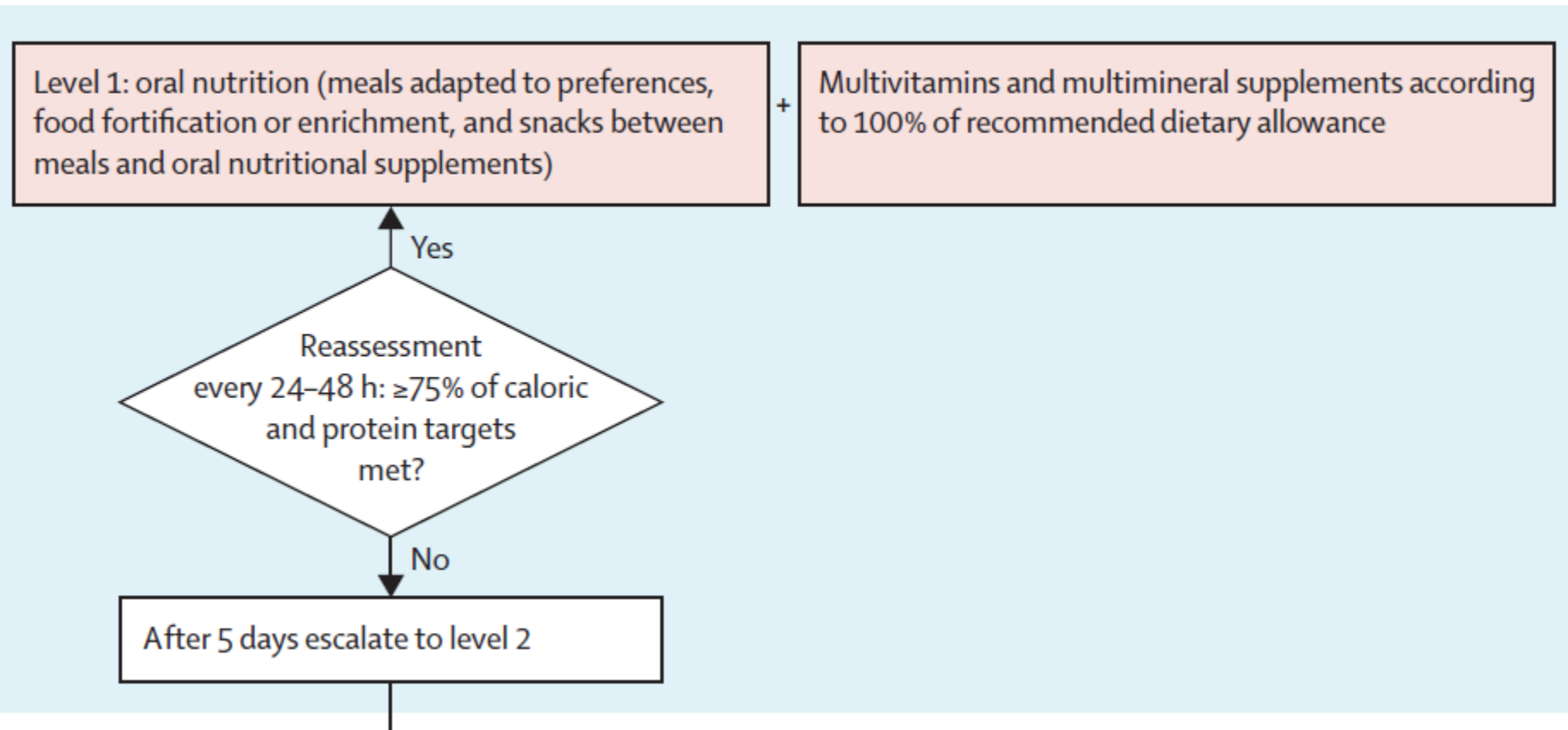
**Micronutrient
requirements**
Multivitamin use; other
micronutrients
according to specific
laboratory results

Specific targets
Disease-specific
adaptations
(eg, medium-chain
triglycerides, low
potassium in patients
with renal failure)

Individualised nutritional support in medical inpatients at nutritional risk: a randomised clinical trial

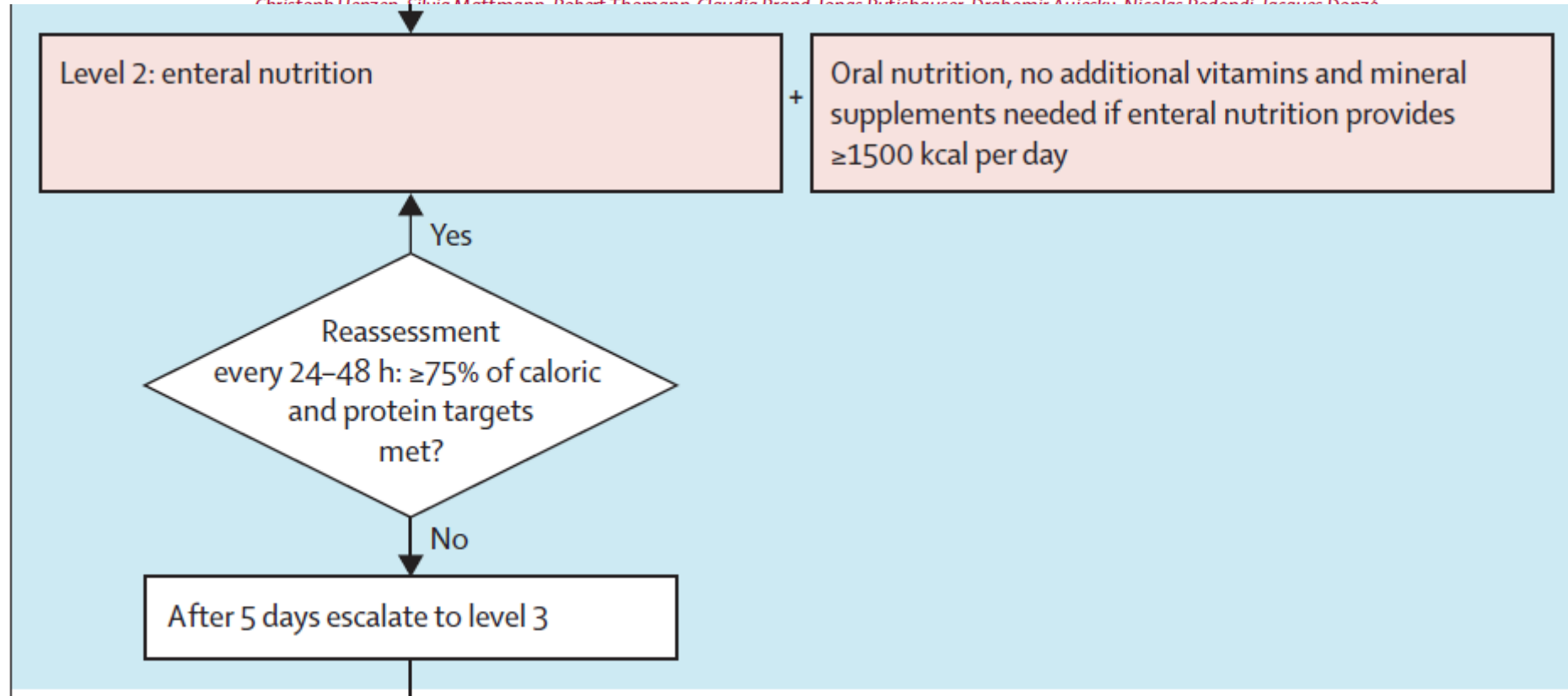
Philippe Schryvers, Barbara Fek, Valérie Desbailly, Martine Colles, Monique Deles, Florence Gosses, Alexander Kutz, Pascal Tikhelst

Strategy to reach the nutrition targets



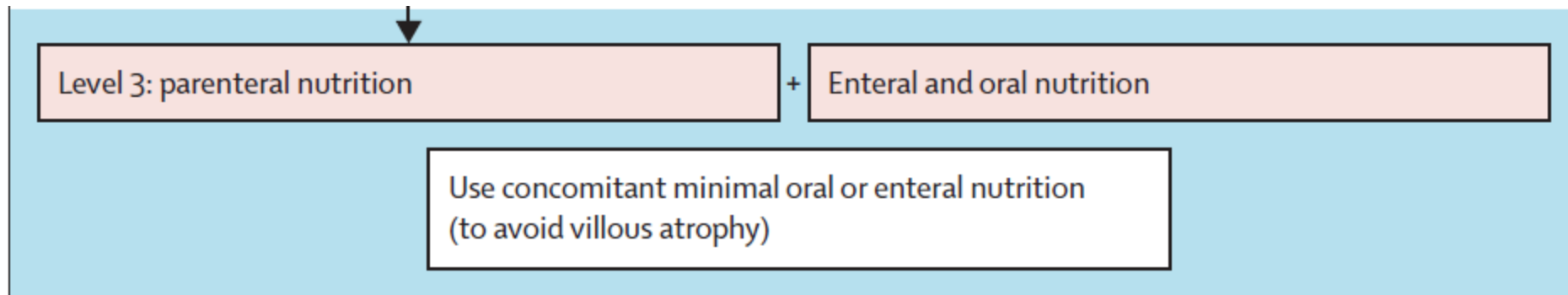
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Philipp Schuetz, Rebecca Fehr, Valerie Baechli, Martina Geiser, Manuela Deiss, Filomena Gomes, Alexander Kutz, Pascal Tribolet, Thomas Bregenzer, Nina Braun, Claus Hoess, Vojtech Pavlicek, Sarah Schmid, Stefan Bilz, Sarah Sigrist, Michael Brändle, Carmen Benz, Christof Henzen, Silvia Mettmann, Robert Thomann, Claudia Brand, Jeroen Butcher, Deshaun Avieson, Nicolas Bodondi, Jacques Denz

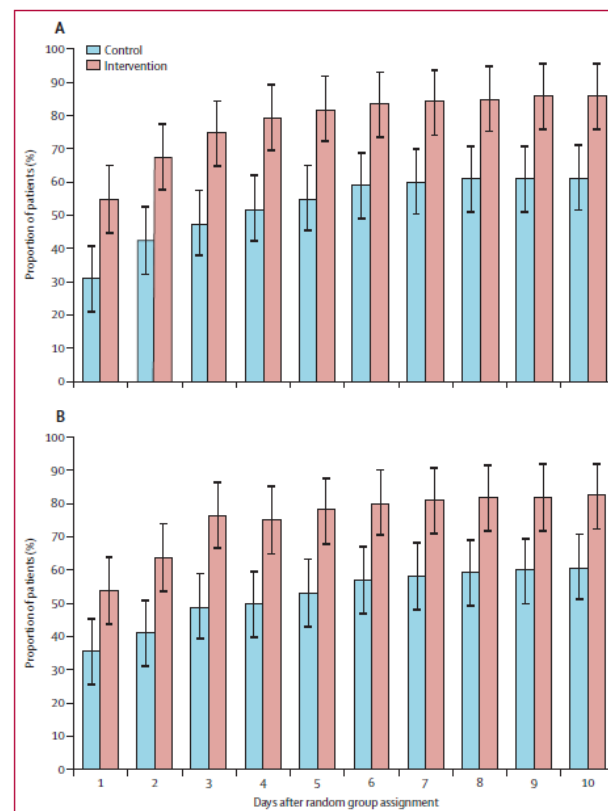


Individualised nutritional support in medical inpatients at nutritional risk: a randomised clinical trial

Philipp Schuetz, Rebecca Fehr, Valerie Baechli, Martina Geiser, Manuela Deiss, Filomena Gomes, Alexander Kutz, Pascal Tribolet, Thomas Bregenzer, Nina Braun, Claus Hoess, Vojtech Pavlicek, Sarah Schmid, Stefan Bilz, Sarah Sigrist, Michael Brändle, Carmen Benz, Christoph Henzen, Silvia Mattmann, Robert Thomann, Claudia Brand, Jonas Rutishauser, Drahomir Aujesky, Nicolas Rodondi, Jacques Donzé, Zeno Stanga, Beat Mueller**



Individualised nutritional support in medical inpatients at nutritional risk: a randomised clinical trial



a Gomes, Alexander Kutz, Pascal Tribolet,
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ser, Drahomir Aujesky, Nicolas Rodondi, Jacques Donzé,

Lancet 2019; 393: 2312–21

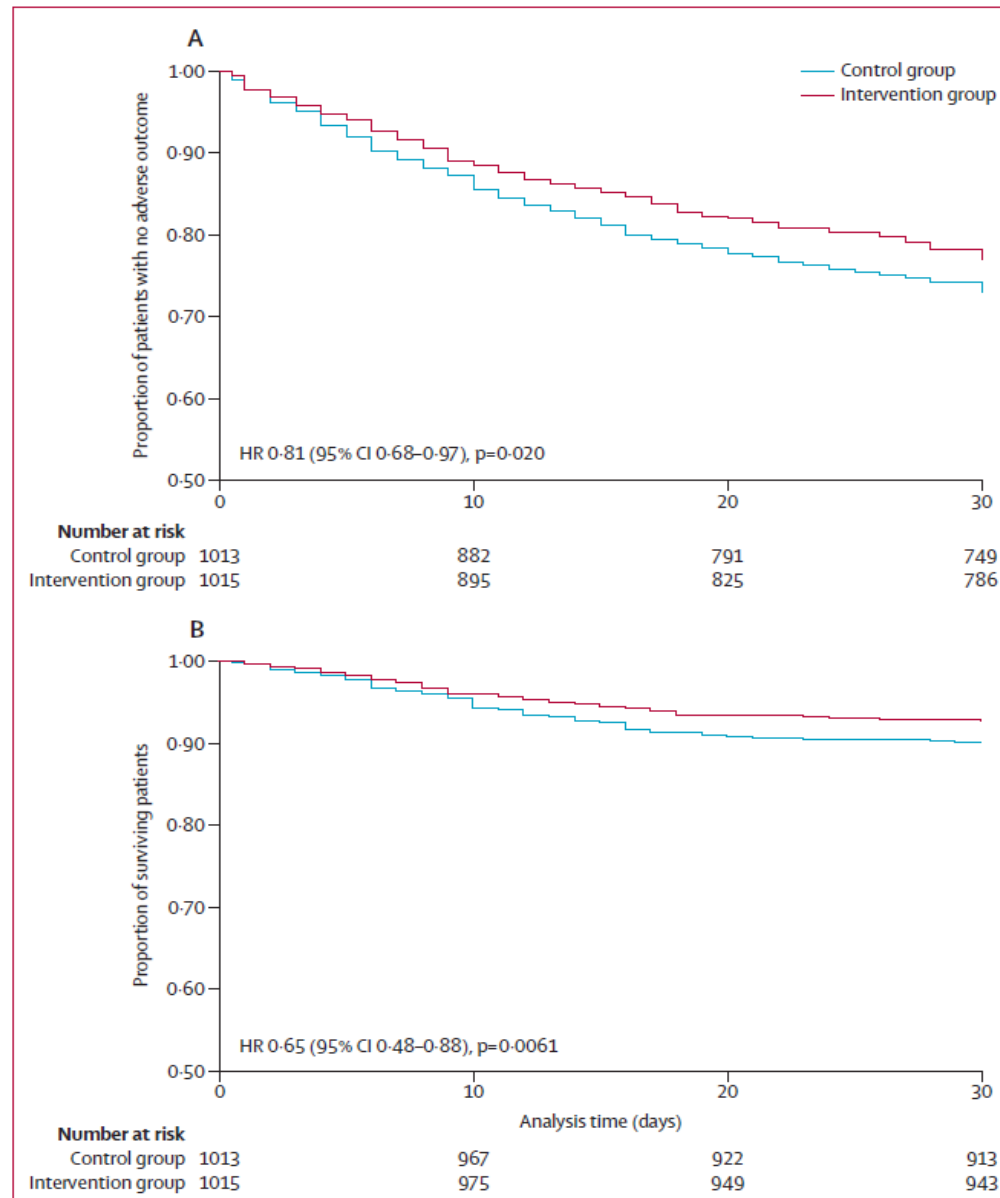
Figure 3: Proportion of patients reaching caloric (A) and protein (B) requirements during the first 10 days after random group assignment

Individualised nutritional support in medical inpatients at nutritional risk: a randomised clinical trial

Philipp Schuetz, Rebecca Feh
Thomas Bregenzer, Nina Bra
Christoph Henzen, Silvia Mai
Zeno Stanga*, Beat Mueller*

tz, Pascal Tribolet,
xel Brändle, Carmen Benz,
Nicolas Rodondi, Jacques Donzé,

Lancet 2019; 393: 2312-21



Trial Design in Critical Care Nutrition: The Past, Present and Future

Lee-anne S. Chapple ^{1,2,3,*} , Emma J. Ridley ^{4,5}  and Marianne J. Chapman ^{1,2,3} 

Nutrients **2020**, *12*, 3694;

Table 1. Summary of critical care nutrition study design in the past, present, and future.

	Past	Present	Future
Study methodology	Small physiological, observational, and single-centre comparative trials; poorly defined and underdeveloped processes	Large phase III trials (usually in the 1000s); a priori defined protocols, outcomes, and interim analyses	Sophisticated statistical techniques; larger sample sizes (tens of thousands); adaptive trial designs
Population	Small, specific populations often from specialist centres	General, heterogenic populations	Homogenous; selected based on anticipated response to nutrition intervention (e.g., malnourished); includes non-invasively ventilated cohorts
Intervention	Calories; early vs. late initiation; route of delivery e.g., gastric vs. jejunal, EN vs. PN, protein- vs. glucose-based PN, medium- vs. long-chain triglycerides and branched chain amino acids; immunonutrition	General interventions for general questions focused only on the period in ICU	Synergistic; patient-specific; based on mechanisms; extension of interventions beyond ICU discharge considering illness trajectory
Outcomes	Calorie delivery; nitrogen balance; incidence of infection; mortality (but underpowered to show an effect in the latter)	Robust clinical outcomes such as mortality	Patient recovery; functional outcomes; valid surrogate markers

Abbreviations: EN = enteral nutrition; ICU = intensive care unit; PN = parenteral nutrition

Poor physical recovery after critical illness: incidence, features, risk factors, pathophysiology, and evidence-based therapies

www.co-criticalcare.com

Yente Florine Niké Boelens^{a,b}, Max Melchers^{a,b},
and Arthur Raymond Hubert van Zanten^{a,b}

Volume 28 • Number 4 • August 2022

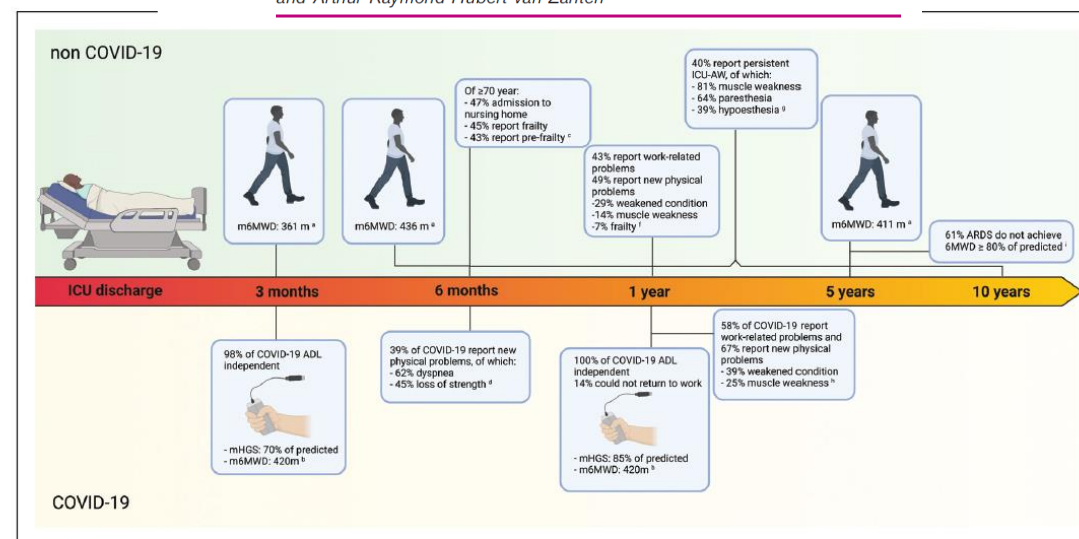


FIGURE 1. Visualized summary of study results regarding prevalence and incidence of impaired physical function following ICU-discharge in COVID-19 and non-COVID-19 patients. Legend: ^a Systematic review and meta-analysis including 16 studies and 1755 ICU survivors [12^{***}]; ^b Prospective single-center study including 114 COVID-19 ICU survivors, of which 98 analyzed at 3 months, and 51 analyzed at one year [17^{*}]; ^c Prospective cohort study including 266 ≥ 70-year-old ICU-survivors [13]; ^d Prospective multicenter cohort study including 212 COVID-19 ICU-survivors [15^{*}]; ^e Population-based cohort study including 546 ≥ 70-year-old ICU-survivors [33]; ^f Prospective multicenter cohort study including 2345 adult ICU-survivors [4^{***}]; ^g Prospective multicenter cohort study including 301 COVID-19 ICU survivors, of which 246 analyzed at one year [14^{***}]; ^h Single-center cohort study including 149 ICU-survivors [10^{*}]; ⁱ Single-center cohort study including 109 ARDS ICU-survivors, of which 94 were analyzed at 5 years [11]. 6MWD, Six Minute Walking Distance; ADL, Activities of Daily Living; COVID-19, Coronavirus Disease 2019; ICU, intensive care unit; ICU-AW, ICU-acquired weakness; m6MWD, Mean Six Minute Walking Distance; mHGS, mean Hand Grip Strength. Created with BioRender.com.



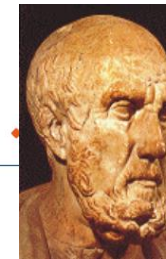
Core Outcome Measures for Clinical Effectiveness Trials of Nutritional and Metabolic Interventions in Critical Illness: An International Modified Delphi Consensus Study Evaluation (CONCISE)

Thomas William Davies; Rob van Gassel; Marcel van de Poll; Jan Gunst; Michael P Casaer; Kenneth B Christopher; Jean-Charles Preiser; Aileen Hill; Kursat Gundogan; Annika Reintam Blaser; Anne-Françoise Rousseau; Carol Hodgson; Dale M Needham; Melina Castro; Stefan Schaller; Thomas McClelland; James J Pilkington; Carla M Sevin; Paul E Wischmeyer; Zheng Yii Lee; Deepak Govil; Andrew Li; Lee-anne Chapple; Linda Denehy; Juan Carlos Montejo-González; Beth Taylor; Danielle E Bear; Rupert Pearce; John Prowle; Angela McNelly; Zudin A Puthuchearu

Crit Care, 2022

“Standardized methods to assess and report these outcomes should be a priority to enable robust detection of interventions on these outcomes”

Compher, C, 2021; JPEN



Hippocrate (470-377 av. JC)

« Que ta nourriture soit ton médicament! »

*Primum non nocere... avec une nutrition inadéquate:
apports excessifs à la phase aiguë, insuffisants en phase
tardive*

