

CORRESPONDENCE

Open Access



How to define parenteral nutrition

Annika Reintam Blaser^{1,2*}, Antonella Cotoia³, Mette M. Berger⁴, Martin Padar^{1,5}, Yaseen M. Arabi⁶, Michael P. Casaer⁷, Jan Gunst⁷, Imre W. K. Kouw^{8,9}, Manu L. N. G. Malbrain¹⁰, Stefan J. Schaller^{11,12}, Joel Starkopf^{1,5}, Martin Sundström Rehal^{13,14}, Arthur R. H. van Zanten^{8,9}, Kaspar F. Bachmann^{1,15} and the GUTPHOS study sites and investigators

In the context of an international, multicentre observational study—the GUTPHOS study [1]—the investigators documented the use of parenteral nutrition (PN), including daily energy intake via this route. A secondary outcome, “days free of PN,” was planned to validate a gastrointestinal dysfunction score. During data quality check, we observed that the definition of PN varied among the participating sites. This observation was further confirmed in a subsequent survey, in which the sites were asked to report the approach they used to document PN. All 28 GUTPHOS study sites (21 from Europe, four from Asia, one from North America, one from South America and one from Oceania) responded and reported their approach (Fig. 1A). In addition to some expectable differences in practice (e.g., using multi-chamber bags vs. separate components), relevant conceptual variability was observed, leading the steering committee to develop a consensus definition for PN in GUTPHOS: “The administration of intravenous amino acids or lipids alone, or any combination of at least two macronutrient components is considered as being PN, whereas administration of only glucose in any concentration is not.” Sites adjusted their data accordingly.

In this correspondence, we aim to draw attention to the unclear definition of PN and call for action.

Despite numerous guidelines and studies addressing PN in the literature, clear guidance on which components and amounts are necessary to define PN for

clinical studies still needs to be provided. The European Society for Clinical Nutrition and Metabolism (ESPEN) defines PN as a nutrition therapy provided through intravenous (central or peripheral) administration of amino acids, glucose, lipids, electrolytes, vitamins and trace elements [2]. The American Society for Parenteral and Enteral Nutrition (ASPEN) defines PN as nutrition provided when patients cannot use the gastrointestinal (GI) tract or nutrition needs cannot be met through the GI tract alone [3]. ESPEN guidelines on ICU nutrition mention that parenteral and enteral feeding preparations differ because commercially available PN solutions contain amino acids and glucose with or without lipids (multi-chamber bags) but no micronutrients for stability reasons [4]. While it is clear that providing all macronutrients and micronutrients intravenously is considered PN, the minimum requirements for defining PN in clinical studies need to be clarified (Fig. 1B).

When PN was first introduced in the 1960s, it was initially called “total parenteral nutrition (TPN)”, designed to cover all nutrition needs, i.e. to provide fluid, protein, carbohydrate, fat, vitamins, trace elements and minerals. The term was simplified to PN after the start of the millennium, and this abbreviation has led to different interpretations. Our current observation suggests the need to define PN more precisely for clinical studies. In the ongoing GUTPHOS study, the question of the PN definition was never raised during the study protocol development nor by sites during data collection, although the Case Report Form manual listed dextrose-based maintenance solutions as non-nutritional calories, yet without further specification and guidance. Our observation suggests a high confidence level in personal or local interpretations,

*Correspondence:

Annika Reintam Blaser
annika.reintam.blaser@ut.ee

Full list of author information is available at the end of the article



© The Author(s) 2024. **Open Access** This article is licensed under a Creative Commons Attribution-NonCommercial-NoDerivatives 4.0 International License, which permits any non-commercial use, sharing, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if you modified the licensed material. You do not have permission under this licence to share adapted material derived from this article or parts of it. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <http://creativecommons.org/licenses/by-nc-nd/4.0/>.

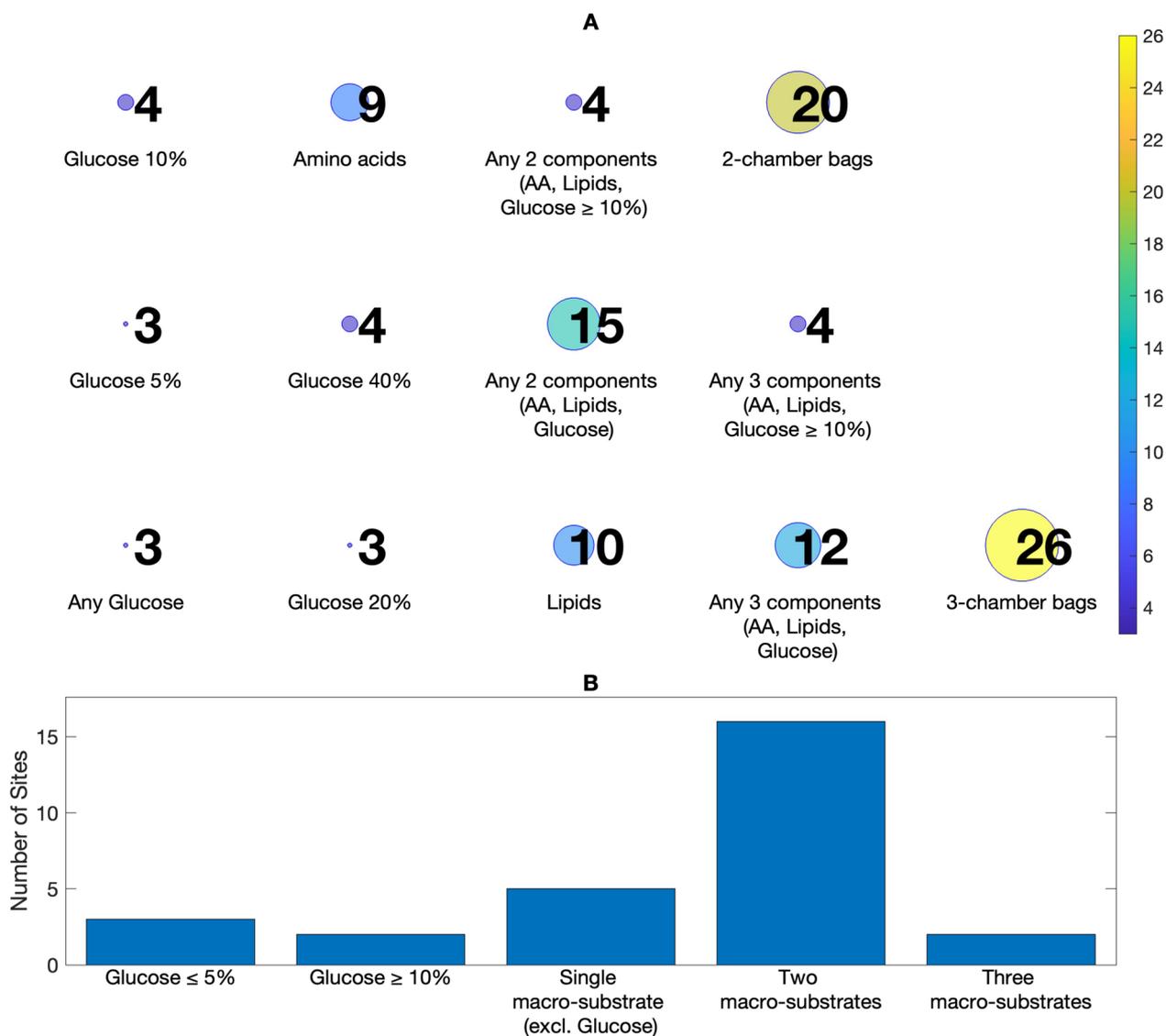


Fig. 1 Results of the survey among 28 sites participating in the GUTPHOS study. **A** Frequency of Positive Responses for PN Options. Bubble chart representing the frequency of responses for different Parenteral Nutrition (PN) options in the survey with 28 sites. Each bubble represents a PN option. Each bubble’s size and colour intensity correspond to the number of positive responses for that option, with larger and brighter bubbles indicating more frequent selection. The number inside each bubble indicates the exact count of positive responses. PN options are labelled beneath each bubble. AA amino acids. **B** Summary of minimum requirements to label a solution as PN. Bar chart illustrating the diversity in defining Parenteral Nutrition (PN) among survey respondents. The x-axis shows five categories of increasing complexity in PN definition, from "Any single component" to "Only commercial preparations." The y-axis and bar heights represent the number of responses in each category. The sites were categorised according to the minimal definition (left-to-right on the x-axis)

whereas, in reality, these interpretations are widely different. In randomised controlled trials (RCT) investigating specific PN products, the variation in definition may be less important than in observational studies. The most recent large NUTRIREA-3 RCT defined PN as “ternary admixture bags containing three groups of macronutrients used according to standard practice” [5]. However, in the NUTRIREA-2 and CALORIES RCTs that

compared enteral nutrition (EN) vs. PN, PN was defined as nutrition provided through a central venous catheter [6, 7], leaving considerable space for different interpretations. Accordingly, RCTs would benefit from a detailed definition of PN.

Developing a precise PN definition is challenging. On the one hand, any substrate (glucose, amino acid, lipid) delivered intravenously, independent of the amount

and indication, are parenteral nutrients that could be considered PN. On the other hand, from the perspective of sustaining life over an extended period of time, PN would be defined as covering the body's requirement for all macro- and micro-nutrients. Therefore, another relevant question arises: Should supplemental PN (SPN), used in conjunction with oral or enteral intake, be defined differently regarding its composition? With such differentiation, PN administered together with EN is not always considered SPN when capturing different feeding routes in studies or databases.

Taken together, despite a simple initial definition of PN that seemed intuitively clear to most clinicians, a precise, unified definition for data comparison in clinical studies is not available and needs to be developed. The above definition, retrospectively applied to the GUTPHOS study, should only be understood as a compromise aiming at unifying results from different sites in a particular study rather than a consensus definition proposal. We wish to draw attention to varying interpretations of "PN" as a pitfall hindering comparisons between sites or studies. We call critical care and nutrition societies for a joint effort to find a global consensus on a detailed definition of PN.

Abbreviations

ASPEN	American Society for Parenteral and Enteral Nutrition
EN	Enteral nutrition
ESPEN	European Society for Clinical Nutrition and Metabolism
ICU	Intensive care unit
PN	Parenteral nutrition
RCT	Randomized controlled trial

Acknowledgements

We thank all collaborators in all study sites.

the GUTPHOS study sites and investigators: Gelderse Vallei Hospital, Ede, The Netherlands: Yvonne Swaen-Dekkers; Tartu University Hospital, Tartu, Estonia: Anna-Liisa Voomets; Karolinska University Hospital Huddinge, Stockholm, Sweden: Rebecca Lindström; Karolinska University Hospital Solna, Solna, Sweden: Jonas Blixt; Luzerner Kantonsspital, Luzern, Switzerland: Benjamin Hess; CHUV, Centre hospitalier universitaire vaudois, Lausanne, Switzerland: Olivier Pantet; King Abdullah International Medical Research Center, Riyadh, Saudi Arabia: Yasir Alzoubi; UZ Leuven, Leuven, Belgium: Liese Mebis; Charité—Universitätsmedizin Berlin, Corporate Member of Freie Universität Berlin and Humboldt Universität zu Berlin, Department of Anaesthesiology and Intensive Care Medicine (CCM/CVK), Berlin, Germany: Linus O. Warner; Klinikum rechts der Isar Technische Universität München, Munich, Germany: Kristina Fuest; Azienda Ospedaliero Universitaria "Policlinico Riuniti", Foggia, Italy: Francesco Cardinale; Beijing Hospital, Beijing, China: Zhigang Chang; University Hospitals of North Midlands, North Midlands, UK: Ramprasad Matsa; H.I.G.A. San Martín de La Plata, Buenos Aires, Argentina: Cecilia Loudet; General Hospital of Atikki, KAT, Athens, Greece: Maria Theodorakopoulou; USL Romagna: Azienda USL della Romagna, Cesena, Italy: Giuliano Bolondi; Královské Vinohrady University Hospital, Prague, Czech Republic: František Duška; Hospital Clínic de Barcelona, Barcelona, Spain: Juan Carlos Lopez-Delgado; Auckland City Hospital, Auckland, New Zealand: Varsha Asrani; Univerity Clinical Center Niš, Niš, Serbia: Natalija Vukovic; North Estonia Medical Centre, Tallinn, Estonia: Oskar Appelberg; The Methodist Hospital Research Institute, Houston, USA: Raul Sanchez Leon; CHU de Besançon, Besançon, France: Guillaume Besch; Universität Leipzig, Leipzig, Germany: Sirak Petros; Södersjukhuset AB, Stockholm, Sweden: Rebecka Rubenson Wahlin; West China Hospital, Chengdu, China: Qin Wu; University Clinical Centre of Serbia, Belgrade, Serbia: Jovana Stanisavljevic; The First Hospital of Jilin University, Jilin, China: Zhang Dong

Author contributions

ARB, AC, MMB, MP and KFB conceptualised and drafted the manuscript, KFB visualised the results, and all the co-authors reviewed and revised the manuscript.

Funding

The GUTPHOS study was funded through an ESICM Fresenius Kabi Clinical Nutrition Award 2023 and the Estonian Research Council (PRG1255).

Availability of data and materials

Data will be made available on reasonable request.

Declarations

Ethics approval and consent to participate

The GUTPHOS study was primarily approved by the University of Tartu Ethics Committee on May 29th, 2023 (approval number 377/T-15), and each participating site obtained local ethics committee approval according to local site country and institutional regulations.

Consent for publication

Not applicable.

Competing interests

ARB received a consultancy fee from VIPUN Medical and is holding a grant from the Estonian Research Council (PRG1255). MMB received lecture honoraria from Baxter, Fresenius Kabi and Nestlé Health Int. SJS received grants and non-financial support from Reactive Robotics GmbH (Munich, Germany), ASP GmbH (Attendorn, Germany), STIMIT AG (Biel, Switzerland), ESICM (Geneva, Switzerland), grants, personal fees, and non-financial support from Fresenius Kabi Deutschland GmbH (Bad Homburg, Germany), grants from the Innovationsfond of The Federal Joint Committee (G-BA), personal fees from Springer Verlag GmbH (Vienna, Austria) for educational purposes and Advanz Pharma GmbH (Bielefeld, Germany), non-financial support from national and international societies (and their congress organisers) in the field of anaesthesiology and intensive care medicine, outside the submitted work. SJS holds stocks in small amounts from Alphabet Inc., Bayer AG, and Siemens AG; these holdings have not affected any decisions regarding his research or this study. MLNGM is member of the medical advisory Board of Pulsion Medical Systems (now fully part of Getinge group), Sentinel Medical and Baxter. He consults for BBraun, Becton Dickinson, Spiegelberg, Medtronic, MedCaptain, and Holtech Medical, and received speaker's fees from PeerVoice. He holds stock options for Sentinel, Serenno and Potrero. AvZ reported receiving honoraria for advisory board meetings, lectures, research, and travel expenses from AOP Pharma, Abbott, Baxter, Cardinal Health, Danone-Nutricia, DIM3, Dutch Medical Food, Fresenius Kabi, GE Healthcare, InBody, Mermaid, Rousselot, and Lyric. MPC and JG receive a senior clinical investigator fellowship funded by Research Foundation-Flanders. AC, YA, JS, MSR, MP and IWKK report no conflicts of interest.

Author details

¹Department of Anaesthesiology and Intensive Care, Institute of Clinical Medicine, University of Tartu, Tartu, Estonia. ²Lucerne Cantonal Hospital, Lucerne, Switzerland. ³Department of Anesthesia and Intensive Care, University Hospital of Foggia, Foggia, Italy. ⁴Faculty of Medicine and Biology, Lausanne University, Lausanne, Switzerland. ⁵Tartu University Hospital, Tartu, Estonia. ⁶College of Medicine, King Saud Bin Abdulaziz University for Health Sciences, King Abdullah International Medical Research Center, Intensive Care Department, King Abdulaziz Medical City, Ministry of National Guard-Health Affairs, Riyadh, Saudi Arabia. ⁷Clinical Division and Laboratory of Intensive Care Medicine, Department of Cellular and Molecular Medicine, KU Leuven, Leuven, Belgium. ⁸Gelderse Vallei Hospital, Ede, The Netherlands. ⁹Division of Human Nutrition and Health, Nutritional Biology, Wageningen University and Research, Wageningen, The Netherlands. ¹⁰First Department of Anaesthesiology and Intensive Therapy, Medical University of Lublin, Lublin, Poland. ¹¹Department of Anaesthesia, Intensive Care Medicine and Pain Medicine, Division of General Anaesthesia and Intensive Care Medicine, Medical University of Vienna, Vienna, Austria. ¹²Department of Anaesthesiology and Intensive Care Medicine (CCM/CVK), Charité - Universitätsmedizin Berlin, Corporate Member of Freie Universität Berlin and Humboldt Universität Zu Berlin, Berlin, Germany. ¹³Perioperative Medicine and Intensive Care, Karolinska University Hospital, Stockholm,

Sweden. ¹⁴Division of Anaesthesia and Intensive Care, Department of Clinical Science, Intervention and Technology (CLINTEC), Karolinska Institutet, Stockholm, Sweden. ¹⁵Department of Intensive Care Medicine, Inselspital, Bern University Hospital, University of Bern, Bern, Switzerland. ¹⁶Karolinska University Hospital Huddinge, Stockholm, Sweden. ¹⁷Karolinska University Hospital Solna, Solna, Sweden. ¹⁸CHUV, Centre Hospitalier Universitaire Vaudois, Lausanne, Switzerland. ¹⁹King Abdullah International Medical Research Center, Riyadh, Saudi Arabia. ²⁰UZ Leuven, Leuven, Belgium. ²¹Klinikum Rechts der Isar Technische Universität München, Munich, Germany. ²²Azienda Ospedaliero Universitaria "Policlinico Riuniti", Foggia, Italy. ²³Beijing Hospital, Beijing, China. ²⁴University Hospitals of North Midlands, North Midlands, UK. ²⁵H.I.G.A. San Martín de la Plata, Buenos Aires, Argentina. ²⁶General Hospital of Atikki, KAT, Athens, Greece. ²⁷AUSL Romagna: Azienda USL Della Romagna, Cesena, Italy. ²⁸Královské Vinohrady University Hospital, Prague, Czech Republic. ²⁹Hospital Clínic de Barcelona, Barcelona, Spain. ³⁰Auckland City Hospital, Auckland, New Zealand. ³¹University Clinical Center Niš, Niš, Serbia. ³²North Estonia Medical Centre, Tallinn, Estonia. ³³The Methodist Hospital Research Institute, Houston, USA. ³⁴CHU de Besançon, Besançon, France. ³⁵Universität Leipzig, Leipzig, Germany. ³⁶Södersjukhuset AB, Stockholm, Sweden. ³⁷West China Hospital, Chengdu, China. ³⁸University Clinical Centre of Serbia, Belgrade, Serbia. ³⁹The First Hospital of Jilin University, Jilin, China.

Received: 24 October 2024 Accepted: 29 October 2024

Published online: 19 November 2024

References

1. Kouw IWK, Melchers M, Mändul M, Arabi YM, Casaer MP, Cotoia A, et al. Prospective multicenter study to validate the gastrointestinal dysfunction score (GIDS) in intensive care patients: study protocol for Part A of the international GUTPHOS study. *Clin Nutr ESPEN*. 2024;63:702–8.
2. Cederholm T, Barazzoni R, Austin P, Ballmer P, Biolo G, Bischoff SC, et al. ESPEN guidelines on definitions and terminology of clinical nutrition. *Clin Nutr*. 2017;36(1):49–64.
3. American Society for Parenteral and Enteral Nutrition website: [https://www.nutritioncare.org/About_Clinical_Nutrition/What_Is_Parenteral_Nutrition/#:~:text=Parenteral%20nutrition%20\(Parenteral%20%5Bpah%20REN,through%20the%20GI%20tract%20alone](https://www.nutritioncare.org/About_Clinical_Nutrition/What_Is_Parenteral_Nutrition/#:~:text=Parenteral%20nutrition%20(Parenteral%20%5Bpah%20REN,through%20the%20GI%20tract%20alone).
4. Singer P, Reintam Blaser A, Berger MM, Alhazzani W, Calder PC, Casaer MP, et al. ESPEN guideline on clinical nutrition in the intensive care unit. *Clin Nutr*. 2019;38(1):48–79.
5. Reignier J, Plantefeve G, Mira JP, Argaud L, Asfar P, Aissaoui N, et al. Low versus standard calorie and protein feeding in ventilated adults with shock: a randomised, controlled, multicentre, open-label, parallel-group trial (NUTRIREA-3). *Lancet Respir Med*. 2023;11(7):602–12.
6. Reignier J, Boisramé-Helms J, Brisard L, Lascarrou JB, Ait Hssain A, Anguel N, et al. Enteral versus parenteral early nutrition in ventilated adults with shock: a randomised, controlled, multicentre, open-label, parallel-group study (NUTRIREA-2). *Lancet*. 2018;391(10116):133–43.
7. Harvey SE, Parrott F, Harrison DA, Bear DE, Segaran E, Beale R, et al. Trial of the route of early nutritional support in critically ill adults. *N Engl J Med*. 2014;371(18):1673–84.

Publisher's Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.