



OPEN ACCESS

EDITED AND REVIEWED BY
Katherine Samaras,
St. Vincent's Hospital Sydney, Australia

*CORRESPONDENCE

Ozra Tabatabaei-Malazy
✉ tabatabaeim@sina.tums.ac.ir
Solaleh Emamgholipour
✉ semamgholipour@sina.tums.ac.ir

RECEIVED 14 May 2024
ACCEPTED 20 June 2024
PUBLISHED 03 July 2024

CITATION

Khosravi S, Tabatabaei-Malazy O,
Emamgholipour S, Sojoodi M and Shabani P
(2024) Editorial: Prebiotics in the
management of obesity and associated
metabolic disorders.
Front. Endocrinol. 15:1432530.
doi: 10.3389/fendo.2024.1432530

COPYRIGHT

© 2024 Khosravi, Tabatabaei-Malazy,
Emamgholipour, Sojoodi and Shabani. This is
an open-access article distributed under the
terms of the [Creative Commons Attribution
License \(CC BY\)](#). The use, distribution or
reproduction in other forums is permitted,
provided the original author(s) and the
copyright owner(s) are credited and that the
original publication in this journal is cited, in
accordance with accepted academic
practice. No use, distribution or reproduction
is permitted which does not comply with
these terms.

Editorial: Prebiotics in the management of obesity and associated metabolic disorders

Sepehr Khosravi¹, Ozra Tabatabaei-Malazy^{1,2*},
Solaleh Emamgholipour^{3,4*}, Mozhdeh Sojoodi⁵
and Parisa Shabani⁶

¹Non-Communicable Diseases Research Center, Endocrinology and Metabolism Population Sciences Institute, Tehran University of Medical Sciences, Tehran, Iran, ²Endocrinology and Metabolism Research Center, Endocrinology and Metabolism Clinical Sciences Institute, Tehran University of Medical Sciences, Tehran, Iran, ³Department of Clinical Biochemistry, School of Medicine, Tehran University of Medical Sciences, Tehran, Iran, ⁴Metabolic Disorders Research Center, Endocrinology and Metabolism Molecular-Cellular Sciences Institute, Tehran University of Medical Sciences, Tehran, Iran, ⁵Division of Surgical Oncology, Massachusetts General Hospital, Harvard Medical School, Boston, MA, United States, ⁶Frankel Institute For Heart and Brain Health, University of Michigan, Medial Center, Ann Arbor, MI, United States

KEYWORDS

weight management, overweight, obesity, microbiota, weight regain, prebiotics

Editorial on the Research Topic

Prebiotics in the management of obesity and associated metabolic disorders

The prevalence of obesity and metabolic comorbidities has considerably increased worldwide over the past decades (1). Extensive efforts have been made to address obesity, but current therapeutic approaches have not yielded satisfactory results, indicating a need for alternative solutions. Imbalanced gut microbiota or dysbiosis is thought to be a principal factor in the development of obesity and metabolic consequences. The approach of modulating the gut microbiota using oral supplementation with prebiotics has received the lion's share of attention for its beneficial effects on managing obesity and associated metabolic disorders. Preclinical and clinical studies have reported the benefits of prebiotics in combatting obesity and metabolic abnormalities through various mechanisms of action. However, the results are inconsistent, necessitating further studies to validate the conclusions. Further detailed studies are needed to understand how changes in microbial signatures affect the management of obesity and its related conditions, such as diabetes and fatty liver. This editorial on the research topic "Prebiotics in the Management of Obesity and Associated Metabolic Disorders" explores the therapeutic potential of modifying gut microbiota through various means, including diet, prebiotics, probiotics, and synbiotics. It highlights innovative strategies that could lead to significant advances in the management and treatment of metabolic disorders.

The burgeoning field of gut microbiota research offers novel insights into how our internal microbial communities affect our health, particularly about metabolic diseases such as obesity and nonalcoholic steatohepatitis (NASH). With obesity reaching epidemic proportions globally and NASH becoming more recognized as a major cause of liver disease, understanding the role of gut microbiota in these conditions is becoming crucial (1). Xiang et al. study delves into how the approximately 40 trillion microbes in our gut can

influence NASH progression. It highlights the critical role of microbial dysbiosis in altering hepatic lipid metabolism and inflammatory responses, pointing out that microbial metabolites, especially short-chain fatty acids (SCFAs) derived from dietary fibers, are vital. SCFAs like butyrate provide substantial energy to intestinal cells and have profound effects on intestinal and liver health (2, 3). It is well known that gut microbiota plays a key role in the metabolism of bile acids in liver homeostasis, facilitated by receptors such as the farnesoid X receptor (FXR) and Takeda G protein-coupled receptor 5 (TGR5) (4). Insufficient FXR and TGR5 activities under NASH conditions highlight the need to develop microbiota-centered therapies that could transform the treatment landscape for liver diseases by leveraging microbial interactions. Xiang et al. emphasize that the relationship between gut microbiota and NASH remains complex and requires further studies to establish a clear roadmap for the clinical application of microbiota.

Dietary treatment is recognized for its anti-obesity effects by altering gut microbiota composition (5, 6). Hence, it highlights a novel therapeutic strategy that could be pivotal in managing obesity and its associated metabolic complexities. Song et al. study explores the anti-obesity effects of Platycodon Grandiflorum Polysaccharide (PGNP) by gut microbiota modulation. This intervention has shown promising results in rebalancing gut microbial communities, notably increasing Bacteroidetes and reducing Firmicutes. These changes are associated with enhanced production of metabolites that positively influence lipid metabolism and inflammation—key players in obesity and metabolic syndrome.

Furthermore, polyamine metabolites, essential for cellular health and immune function, can be synthesized by gut microbiota from amino acids and regulated by host cells, gut microbiota, and diet (7). The potential of polyamine synthesis enhancement in treating obesity-related type 2 diabetes and metabolic syndrome is gaining attention in the study of Bui et al. Increasing polyamine levels through diet and probiotic supplementation may mitigate the immune deficiencies and metabolic disruptions accompanying these conditions. This approach underscores the importance of gut microbiota in metabolic health and opens up new avenues for creating nutraceutical interventions that capitalize on these mechanisms.

Child obesity is another area of research focus, as this group requires further attention due to the long-term effects of obesity. Prebiotics may help fight childhood obesity by altering gut bacteria, which is a promising research area. Despite the lack of extensive clinical trials, preclinical evidence suggests that prebiotics can significantly influence host metabolism. Wang et al. highlight the importance of further research on how prebiotics work and developing focused treatments to help reduce rising childhood obesity rates. There are several key mechanisms, including the modulation of enteroendocrine function and the systemic effects of prebiotics and their fermentation products on lipid and glucose homeostasis (8).

Lastly, Rasaei et al. study shows the overall effectiveness of biotics—prebiotics, probiotics, and synbiotics—across various demographics. They found an inverse association between biotics consumption and overweight/obesity risk in adults. Moreover, the

consumption of prebiotic: 8–66 g/day, probiotic: 10^4 – 1.35×10^{15} colony-forming unit (CFU)/day, and synbiotic: 10^6 – 1.5×10^{11} CFU/day and 0.5–300 g/day for 2 to 104 weeks showed a favorable effect on the overweight/obesity indicators in adults. However, while generally effective in reducing obesity-related measures such as body weight and BMI in adults, their impact is less significant in pregnant women and infants. Some of the challenges in determining the effectiveness of currently available probiotic preparations for weight control are the differences in the strains and formulations of probiotics used, individual responses to probiotics, dietary habits, genetic predispositions, lifestyle factors, personal gut microbiota composition, intervention duration, specific target population, and the quality of study designs (9). These findings highlight the urgent need for well-designed clinical trials to explore how biotics can influence obesity and metabolic health comprehensively and effectively across diverse demographic groups.

As we unravel the complex interactions between our gut microbiota and metabolic health, targeted interventions focusing on this interplay may offer viable solutions for managing and potentially reversing the effects of metabolic diseases. The studies reviewed here provide a hopeful glimpse into the future of metabolic disorder treatments, suggesting that with further research and clinical trials, we can better tailor these interventions to benefit individuals at various stages of life, from infancy through adulthood. Emphasizing the need for continued research, these insights herald a new era in medical science where gut microbiota modulation becomes a cornerstone of innovative therapeutic strategies.

Author contributions

SK: Validation, Visualization, Writing – original draft, Writing – review & editing. OT-M: Conceptualization, Supervision, Validation, Visualization, Writing – review & editing, Writing – original draft. SE: Conceptualization, Supervision, Validation, Writing – review & editing. MS: Validation, Visualization, Writing – review & editing. PS: Validation, Visualization, Writing – review & editing.

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Publisher's note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

References

1. NCD Risk Factor Collaboration (NCD-RisC), Phelps NH, Singleton RK, Zhou B, Heap RA, Mishra A, et al. Worldwide trends in underweight and obesity from 1990 to 2022: a pooled analysis of 3663 population-representative studies with 222 million children, adolescents, and adults. *Lancet*. (2024) 403:1027–50. doi: 10.1016/S0140-6736(23)02750-2
2. Ramakrishna BS. Role of the gut microbiota in human nutrition and metabolism. *J Gastroenterol Hepatol*. (2013) 28 Suppl 4:9–17. doi: 10.1111/jgh.12294
3. Albillos A, de Gottardi A, Rescigno M. The gut-liver axis in liver disease: Pathophysiological basis for therapy. *J Hepatol*. (2020) 72:558–77. doi: 10.1016/j.jhep.2019.10.003
4. Li T, Chiang JYL. Bile acid-based therapies for non-alcoholic steatohepatitis and alcoholic liver disease. *Hepatobiliary Surg Nutr*. (2020) 9:152–69. doi: 10.21037/hbsn.2019.09.03
5. Chen J, Liu J, Yan C, Zhang C, Pan W, Zhang W, et al. Sarcodon aspratus polysaccharides ameliorated obesity-induced metabolic disorders and modulated gut microbiota dysbiosis in mice fed a high-fat diet. *Food Funct*. (2020) 11:2588–602. doi: 10.1039/C9FO00963A
6. Do Mh, Lee H-B, Oh M-J, Jhun H, Choi SY, Park H-Y. Polysaccharide fraction from greens of raphanus sativus alleviates high fat diet-induced obesity. *Food Chem*. (2020) 343:128395. doi: 10.1016/j.foodchem.2020.128395
7. Sagar NA, Tarafdar S, Agarwal S, Tarafdar A, Sharma S. Polyamines: functions, metabolism, and role in human disease management. *Med Sci*. (2021) 9:44. doi: 10.3390/medsci9020044
8. Rodriguez J, Delzenne NM. Modulation of the gut microbiota-adipose tissue-muscle interactions by prebiotics. *J Endocrinol*. (2021) 249:R1–r23. doi: 10.1530/JOE-20-0499
9. Vallianou N, Stratigou T, Christodoulatos GS, Dalamaga M. Understanding the role of the gut microbiome and microbial metabolites in obesity and obesity-associated metabolic disorders: current evidence and perspectives. *Curr Obes Rep*. (2019) 8:317–32. doi: 10.1007/s13679-019-00352-2