

Short Communication

The New American Addiction: How Weight-Loss Drugs Could Create the Next Epidemic

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Abstract

GLP-1 receptor agonists such as semaglutide (Ozempic, Wegovy) have rapidly gained popularity for obesity treatment, with unprecedented public enthusiasm and off-label use among otherwise healthy populations. While these agents demonstrate substantial weight-loss efficacy, emerging concerns highlight potential pharmaceutical dependency, ethical dilemmas, and long-term safety uncertainties. This article critically examines the mechanisms, benefits, and risks of GLP-1 agonists, emphasizing the need for regulatory oversight, psychological safeguards, and integration with lifestyle interventions. We propose a clinical framework in which GLP-1 agonists serve as a temporary adjunct, while structured exercise, dietary intervention, and behavioral therapy remain the standard of care for sustainable health.

More Information

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Submitted: September 04, 2025

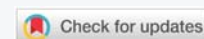
Approved: September 11, 2025

Published: September 12, 2025

How to cite this article: Peters F. The New American Addiction: How Weight-Loss Drugs Could Create the Next Epidemic. J Sports Med Ther. 2025; 10(3): 046-047. Available from: <https://dx.doi.org/10.29328/journal.jsmt.1001096>

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Keywords: GLP-1 receptor agonists; Obesity; Weight loss; Ethics; Policy; Exercise; Pharmacotherapy



Introduction

GLP-1 receptor agonists have shifted from niche pharmacotherapy for type 2 diabetes to mainstream weight-loss interventions, propelled by social media, celebrity endorsements, and aggressive direct-to-consumer marketing [1,2]. Obesity remains a leading public health challenge, contributing to cardiometabolic disease, disability, and healthcare costs [3]. Weight-loss outcomes with GLP-1 agonists are often dramatic; yet concerns regarding over-reliance on medication, rebound weight gain after discontinuation, and proliferation of unregulated compounded formulations demand critical evaluation [4].

Mechanisms and clinical benefits

GLP-1 receptor agonists mimic the action of endogenous incretin hormones, stimulating insulin secretion, suppressing glucagon release, delaying gastric emptying, and reducing appetite. Clinical trials consistently report significant weight loss and improved cardiometabolic profiles among individuals with obesity and type 2 diabetes [4-6]. These findings have accelerated off-label prescribing for non-diabetic populations, despite limited long-term safety data [7].

Emerging concerns: Dependency and discontinuation

A growing pattern of weight regain following

discontinuation raises concerns about psychological and behavioral dependency. Patients often describe a fear of weight regain and a loss of control, reflecting a cycle of pharmacologic reliance: appetite suppression, weight loss, cessation, rapid rebound and re-initiation [5,6]. Unlike medications such as anticoagulants, which prevent immediate, life-threatening events, GLP-1 agonists primarily influence behavioral drivers of weight gain, raising ethical questions about indefinite use for a condition that is largely modifiable through lifestyle.

Economic and regulatory challenges

The high cost of GLP-1 therapy, often exceeding \$1,000 monthly, combined with limited insurance coverage has fueled demand for compounded formulations marketed through online clinics and wellness spas. These compounded products frequently lack standardized dosing, quality control, and regulatory oversight, amplifying risks beyond those of FDA-approved agents [8-10]. Recent reports underscore the urgency of implementing stronger safeguards to protect patients from substandard or counterfeit medications while also addressing inequities in access to evidence-based obesity care [1].

Lifestyle interventions remain foundational

While pharmacotherapy offers short-term efficacy, empirical evidence consistently demonstrates that structured



exercise and dietary interventions remain essential for sustainable weight management, cardiometabolic health, and preservation of lean body mass [11,12]. Rapid weight loss without concomitant resistance training increases the risk of sarcopenia and frailty, particularly in middle-aged and older adults [13]. Healthcare practitioners should emphasize resistance training, protein intake, and long-term behavioral strategies to safeguard lean mass.

Ethical and policy implications

The normalization of GLP-1 agonists as cosmetic ‘quick fixes’ risks marginalizing behavioral medicine and perpetuating a pharmaceutical-first paradigm. Expanding insurance access, regulating compounded alternatives, and mandating integration of exercise and nutrition counseling into obesity treatment protocols represent essential steps toward responsible clinical practice [4,8,9]. Recent discussions have emphasized the need for equitable access, long-term safety monitoring, and safeguards against misuse in otherwise healthy populations [1].

Conclusion

GLP-1 receptor agonists represent a powerful but double-edged tool in obesity management. Their efficacy is clear; however, the risks of psychological dependence, economic burden, and erosion of lifestyle-based care demand a more balanced approach. Pharmacotherapy should complement, not replace, evidence-based exercise and nutrition interventions. Sustainable metabolic health requires empowering individuals through behavioral change rather than fostering long-term pharmaceutical reliance.

References

1. Brown T. Public perceptions and ethical considerations of GLP-1 agonists in obesity care. *Obes Rev.* 2023;24(3):e13550.
2. Patel R, Jones M. The rise of GLP-1 receptor agonists: Trends and implications. *J Obes Res.* 2022;30(6):455–63.
3. Hruby A, Hu FB. The epidemiology of obesity: A big picture. *Pharmacoeconomics.* 2022;40(2):101–15.
4. Rubino D. Long-term effects of GLP-1 receptor agonists in obesity treatment: What we do and don’t know. *Obes Rev.* 2023;24(2):e13528.
5. Wilding JPH. Weight regain after withdrawal of semaglutide: A randomized clinical trial. *Lancet Diabetes Endocrinol.* 2022;10(11):840–9.
6. Jastreboff AM, Aronne LJ, Ahmad NN, Wharton S, Connery L, Alves B, et al. Tirzepatide Once Weekly for the Treatment of Obesity. *N Engl J Med.* 2022;387(3):205–16. Available from: <https://doi.org/10.1056/NEJMoa2206038>
7. Garvey WT. Safety of GLP-1 receptor agonists: Current evidence and future directions. *Diabetes Obes Metab.* 2023;25(1):10–22.
8. Wharton S. Clinical practice guidelines for the treatment of obesity in adults. *CMAJ.* 2021;193(31):E1120–E1136.
9. Greenfield J. Ethical challenges in prescribing GLP-1 receptor agonists for obesity. *Lancet Diabetes Endocrinol.* 2022;10(12):890–9.
10. U.S. Food and Drug Administration. Consumer warning: The dangers of unregulated compounded semaglutide. FDA Public Health Advisory. 2024.
11. Jakicic JM. Role of physical activity and exercise in obesity treatment. *J Clin Endocrinol Metab.* 2022;107(4):1036–50.
12. Swift DL. The effects of exercise and physical activity on weight loss and maintenance. *Prog Cardiovasc Dis.* 2021;67(3):143–52.
13. Folli F. Sarcopenia, frailty, and rapid weight loss: Clinical implications. *J Cachexia Sarcopenia Muscle.* 2022;13(4):1800–12.