

Therapeutic Potential of Mangiferin Derivatives: Enhancing Bioavailability and Diverse Biological Activities

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DESCRIPTION

Mangiferin, a bioactive compound abundantly found in plants such as mangoes and *Anemarrhena asphodeloides*, has long captivated the interest of researchers for its potential therapeutic properties. However, despite its innate benefits, the full realization of mangiferin's therapeutic potential has been hampered by challenges such as limited bioavailability and specificity [1]. To overcome biological activities, scientists have turned to the synthesis of mangiferin derivatives.

This introduction will delve into the multifaceted mangiferin derivatives, elucidating how these modified compounds not only overcome the bioavailability constraints of natural mangiferin but also disclose a spectrum of therapeutic effects. From enhancing antioxidant and anti-inflammatory activities to potentially combating cancer and diabetes, mangiferin derivatives offer an optimistic frontier in drug development and disease management [2]. In this article, we will explore how the synthesis of mangiferin derivatives aims to enhance their bioavailability, enabling improved delivery and distribution within the body.

Furthermore, we will delve into the diverse biological activities exhibited by these derivatives, ranging from their antioxidant potential to their ability to modulate inflammatory pathways, inhibit cancer proliferation, and manage diabetes [3,4]. By the therapeutic potential of mangiferin derivatives, this article seeks to underscore their significance in advancing human health and paving the way for innovative therapeutic interventions.

Enhancing bioavailability of mangiferin

One of the primary motivations behind synthesizing mangiferin derivatives is to improve its bioavailability. Natural mangiferin faces limitations in absorption and distribution within the body, hindering its therapeutic efficacy. By modifying its structure through synthesis, researchers aim to enhance its solubility, stability, and cellular uptake. These modifications pave the way

for improved pharmacokinetic profiles, ensuring that mangiferin derivatives reach their target tissues more effectively [5].

Diverse biological activities

The synthesis of mangiferin derivatives has unlocked a plethora of biological activities beyond its innate properties. These derivatives exhibit versatile pharmacological effects, making them promising candidates for various therapeutic applications [6].

Antioxidant potential: Mangiferin derivatives have demonstrated robust antioxidant activity, scavenging free radicals and protecting cells from oxidative stress-induced damage [7]. These properties make them valuable in combating age-related diseases, neurodegenerative disorders, and cardiovascular conditions.

Anti-inflammatory effects: Inflammation underlies many chronic diseases, including arthritis, inflammatory bowel disease, and asthma. Mangiferin derivatives have shown potent anti-inflammatory effects by modulating key inflammatory pathways [8]. They inhibit the production of pro-inflammatory mediators, offering relief from inflammation and associated symptoms.

Anticancer properties: Cancer remains a formidable health challenge worldwide, necessitating the exploration of novel therapeutic agents. Mangiferin derivatives have garnered attention for their potential anticancer effects [9]. Studies indicate their ability to induce apoptosis, inhibit tumor proliferation, and suppress metastasis in various cancer types. Furthermore, their low toxicity profile offers a bright avenue for developing adjunctive cancer therapies.

Anti-diabetic activity: Diabetes mellitus imposes a significant burden on global healthcare systems, underscoring the need for effective treatment strategies. Mangiferin derivatives have emerged as potential candidates for managing diabetes due to their insulin-sensitizing and glucose-lowering effects. They improve glucose uptake, enhance insulin signaling, and mitigate complications associated with diabetes, offering hope for better disease management [10].

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CONCLUSION

The synthesis of mangiferin derivatives represents a guarantee approach to harnessing the therapeutic potential of this natural compound. By overcoming bioavailability limitations and expanding its biological activities, these derivatives offer new avenues for drug development and disease management. From antioxidant and anti-inflammatory effects to anticancer and anti-diabetic properties, mangiferin derivatives hold immense assurance in addressing a wide range of health conditions. Continued research into their synthesis and pharmacological mechanisms will likely disclose further insights, ultimately translating into novel therapies for improving human health.

REFERENCES

1. Barakat S, Nasr M, Ahmed RF, Badawy S, Mortada N. Recent formulation advances of mangiferin. *Revista Brasileira de Farmacognosia*. 2022;32(6):871-882.
2. Bhatia HS, Candelario-Jalil E, de Oliveira AC, Olajide OA, Martínez-Sánchez G, Fiebich BL. Mangiferin inhibits cyclooxygenase-2 expression and prostaglandin E2 production in activated rat microglial cells. *Arch Biochem Biophys*. 2008;477(2): 253-258.
3. Acosta J, Sevilla I, Salomón S, Nuevas L, Romero A, Amaro D. Determination of mangiferin solubility in solvents used in the biopharmaceutical industry. *J. Pharm. Pharmacogn. Res*. 2016;4(2): 49-53.
4. Dar A, Faizi S, Naqvi S, Roome T, Zikr-ur-Rehman S, Ali M, et al. Analgesic and antioxidant activity of mangiferin and its derivatives: The structure activity relationship. *Biol Pharm Bull*. 2005;28(4): 596-600.
5. Matkowski A, Kus P, Goralska E, Wozniak D. Mangiferin—a bioactive xanthonoid, not only from mango and not just antioxidant. *Mini Rev Med Chem*. 2013;13(3):439-455.
6. Oyama KI, Kondo T. Total Synthesis of flavocoumestrol, a component of the blue supramolecular pigment from *Commelina communis*, on the basis of direct 6-C-Glycosylation of flavan. *J Org Chem*. 2004;69(16): 5240-5246.
7. Wu Z, Wei G, Lian G, Yu B. Synthesis of mangiferin, isomangiferin, and homomangiferin. *J Org Chem*. 2010;75(16):5725-5728.
8. Andreu GP, Delgado R, Velho JA, Curti C, Vercesi AE. Iron complexing activity of mangiferin, a naturally occurring glucosylxanthone, inhibits mitochondrial lipid peroxidation induced by Fe²⁺-citrate. *Eur J Pharmacol*. 2005;513(1-2):47-55.
9. Lee B, Trinh HT, Bae EA, Jung K, Kim DH. Mangiferin inhibits passive cutaneous anaphylaxis reaction and pruritus in mice. *Planta Med*. 2009;75(13):1415-1417.
10. Guo F, Huang C, Liao X, Wang Y, He Y, Feng R, et al. Beneficial effects of mangiferin on hyperlipidemia in high-fat-fed hamsters. *Molecular Nutrition Food Research*. 2011;55(12):1809-1818.