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Rutin Nanoparticles: Pioneering New Frontier skincare**Deep Sharma¹, Anjana Devi², Vijay Sharma³, Vivek Sharma³, Kamal Jeet^{4*}**¹Research scholar, Department of Pharmacy, School of Pharmaceutical and Health Sciences, Career Point University, Hamirpur-176041, India.²Associate Professor, Department of Pharmacy, School of Pharmaceutical and Health Sciences, Career Point University, Hamirpur-176041, India.³Department of Pharmacy, School of Pharmaceutical and Health Sciences, Career Point University, Hamirpur-176041, India.³Department of Pharmacy, School of Pharmaceutical and Health Sciences, Career Point University, Hamirpur-176041, India.⁴Principal, SFC Institute of Pharmacy, Moga, Punjab, India.***Email:** kamaljeetisf@gmail.com**Article Information**

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Dermatological issues, spanning from common inflammatory conditions to skin aging and cancers, present significant challenges for effective treatment. The application of topical medications provides a promising method to specifically target skin problems while reducing systemic side effects. *Rutin*, a naturally occurring flavonoid recognized for its strong antioxidant, anti-inflammatory, and healing properties, presents considerable therapeutic potential for various dermatological applications. However, its limited ability to penetrate the skin and low bioavailability hinder its clinical effectiveness. Nanotechnology provides a practical solution to overcome these challenges by encapsulating *Rutin* in nanoparticles. This review examines the potential of *Rutin*-loaded nanoparticles as innovative surface modification techniques for skin-related uses. We explore the benefits of *Rutin* for skin health, the advantages of nanoparticle-delivery systems that are based on collaboration, alongside the joint effects of integration *Rutin* employing nanotechnology to improve local effectiveness. In addition, we investigate the different types of nanoparticles suitable for *Rutin* encapsulation, their techniques for enhancing drug delivery, and the promising preclinical and recent clinical information supporting the use of *Rutin*-loaded nanoparticles in tackling skin ailments. In the end, we explore the challenges and potential avenues for transforming this innovative approach into effective clinical topical therapies.

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1. INTRODUCTION

The skin, acknowledged as the largest organ in the human body, plays an essential role as a protective barrier against outside factors. It is vulnerable to various skin-related issues, including inflammatory conditions like eczema and psoriasis, infectious diseases, and skin cancers, pigmentary disorders, and the effects of aging. Topical therapies are crucial in skin care, providing a focused approach to apply medical agents directly to the affected areas, which reduces overall exposure and potential

side effects. However, the outer layer corneum, the epidermis, poses a significant barrier for drug absorption, diminishing the effectiveness of multiple topical therapies. *Rutin* (quercetin-3-O-Rutinoside), a common flavonoid found in plants such as buckwheat, citrus fruits, and apples, has garnered significant interest due to its various biological impacts, including strong antioxidant and anti-inflammatory characteristics, vasoprotective, and influences on wound recovery. These therapeutic properties position *Rutin* as a practical option for addressing different skin issues, such as indications of aging, hyperpigmentation, inflammatory skin disorders, and tissue restoration. However, *Rutin*'s natural physicochemical characteristics, including low water solubility and decreased permeability through the stratum corneum, and rapid metabolism, pose challenges for its effective topical use and absorption at the desired site within the skin.

Nanotechnology, which enables the alteration of substances at the nanoscale (1-100 nm), provides innovative solutions to address the limitations of traditional topical drug delivery. Various types of nanoparticles, such as liposomes, solid lipid nanoparticles (SLNs), nanostructured lipid transporters (NLCs), polymer-based nanoparticles, and metallic nanoparticles, can encapsulate therapeutic agents like *Rutin*. This encapsulation enhances skin penetration, supports controlled release, and allows for targeted delivery^{8, 9}. By incorporating *Rutin* into nanoparticles, we can boost its solubility, protect it from degradation, enhance its ability to infiltrate the layer corneum, and promote its accumulation at the specific region within the skin¹⁰. This analysis seeks to investigate the joint efficacy of *Rutin*-loaded nanoparticles as creative regional treatment methods for various skin-related applications, emphasizing the benefits, challenges, and potential advancements in this rapidly evolving field¹⁻³.

2. *Rutin* as Bioactive Compound:

Rutin, a glycosylated derivative of quercetin and part of the flavonoid class of polyphenolic compounds, have diverse pharmacological impacts that are significant for skin wellness.

2.1 Antioxidant Properties:

Oxidative stress, caused by reactive oxygen species (ROS) due to exposure to UV rays, environmental pollution, and routine metabolic processes, is a key factor in the development of numerous skin problems, including aging, inflammation, and skin cancer⁴. *Rutin* exhibits strong antioxidant properties by counteracting free radicals, hindering lipid peroxidation, and improving the activity of

natural antioxidant enzymes such as superoxide dismutase (SOD) and catalase. This antioxidant capability of *Rutin* can shield skin cells from oxidative harm, reducing the negative impacts of environmental stressors and promoting skin health and anti-aging⁵.

2.2 Anti-inflammatory Activity:

Inflammation plays a vital role in many skin conditions, such as eczema, psoriasis, and acne. *Rutin* has shown considerable anti-inflammatory properties by influencing multiple inflammatory pathways. It can reduce the production of pro-inflammatory cytokines like TNF- α , IL-1 β , and IL-6, and impede the activity of inflammatory enzymes such as cyclooxygenase-2 (COX-2) and inducible nitric oxide synthase (iNOS). By reducing the concentrations of inflammatory substances, *Rutin* may help reduce skin irritation, redness, and itching, making it a potential treatment for inflammatory skin issues^{6,7}.

2.3 Wound Healing Promotion:

Wound healing is a multifaceted process that encompasses cell growth, movement, and the restructuring of the extracellular matrix. *Rutin* has been demonstrated to enhance the healing process of wounds by promoting collagen production, the creation of new blood vessels, and the renewal of the skin layer. It can support the growth and movement of fibroblasts, boost collagen accumulation, and foster the maturation of blood vessels at the injury location, ultimately leading to quicker and more effective wound closure. These attributes make *Rutin* a promising option for external use in managing wounds and promoting healing from burns⁸.

2.4 Skin Pigmentation Modulation:

Hyperpigmentation issues, such as melasma and post-inflammatory hyperpigmentation, are common issues related to the skin. *Rutin* has demonstrated potential in controlling skin color by inhibiting melanogenesis, the process responsible for melanin production. It can decrease the activity of tyrosinase, an essential enzyme in melanin production, and interfere with the transfer of melanosomes to keratinocytes. This ability to inhibit melanogenesis indicates that *Rutin* might be advantageous in tackling hyperpigmentation issues and promoting a more uniform skin tone⁹.

2.5 Photoprotective Effects:

Ultraviolet (UV) radiation significantly contributes to the aging process of the skin, hyperpigmentation, and the development of skin cancer. *Rutin* has shown photoprotective properties by absorbing UV radiation, counteracting reactive oxygen species (ROS) generated by UV exposure, and

safeguarding DNA from photo damage. Applying *Rutin* applying it to the skin could help reduce damage caused by UV rays, which encompasses erythema, photoaging, and the probability of skin cancers ¹⁰.

3. Nanoparticles as Distribution Mechanisms for Enhanced Surface Delivery:

Nanotechnology provides a flexible approach to enhance the surface utilization of bioactive compounds such as *Rutin*. Due to their unique physicochemical characteristics—such as tiny size, large surface area, and customizable surface features—nanoparticles can address the limitations of traditional topical preparations and improve the delivery of drugs to the skin ¹¹.

3.1 Enhanced Skin Penetration:

The tiny size of nanoparticles allows them to penetrate the layer corneum effectively, either through the intercellular route (between corneocytes) or the transcellular route (through corneocytes). Nanoparticles can navigate through the tight passages of the layer corneum, delivering *Rutin* deeper into the epidermis and dermis, thereby reaching target cells and tissues more efficiently than larger drug carriers or free drug molecules ¹².

3.2 Controlled and Sustained Release:

Delivery systems based on nanoparticles can be configured to provide controlled and extended distribution of *Rutin* directly at the target site. This method of administration can improve the therapeutic effects, decrease the frequency of doses, and lessen the potential side effects that arise from varying drug concentrations in the skin. Various nanoparticle materials and production methods can be utilized to customize the release kinetics of *Rutin*, ensuring that the efficacy of the therapy is maintained throughout time ^{13, 14}.

3.3 Targeted Distribution and Enhanced Bioavailability:

Nanoparticles can be adjusted with targeting ligands, such as antibodies, peptides, or aptamers, to particularly target unhealthy skin cells or tissues. This exact technique can improve the local concentration of *Rutin* at the action location, enhancing its therapeutic effectiveness while minimizing off-target impacts. Additionally, enclosing *Rutin* in nanoparticles can shield it from degradation caused by enzymes and metabolic activities in the skin, thereby improving its bioavailability and extending its therapeutic impacts ¹⁵.

3.4 Improved Formulation Durability and Cosmetical Acceptability:

Nanoparticles can enhance the physicochemical

stability of *Rutin*, safeguarding it from degradation due to light, oxidation, or hydrolysis. This improved stability can prolong the shelf life of topical products. Furthermore, nanoparticle-based creations can be transformed into aesthetically pleasing objects that offer enhanced texture, spreadability, and tactile feeling, which could lead to better patient compliance and approval ^{16, 17}.

4. *Rutin*-Loaded Nanoparticles: Joint Opportunities for Skin Therapies:

The combination of *Rutin*'s therapeutic properties coupled with the benefits of nanoparticle-based drug delivery promotes a cooperative strategy for developing innovative topical treatments for an array of skin conditions.

4.1 *Rutin*-Loaded Nanoparticles for Inflammatory Skin Disorders:

Rutin-loaded nanoparticles have shown considerable promise in treating inflammatory skin conditions like eczema, psoriasis, and dermatitis. By encapsulating *Rutin* in nanoparticles, its ability to penetrate the skin is enhanced, allowing targeted delivery to inflamed areas within the epidermis and dermis. The extended release from these nanoparticles can extend the anti-inflammatory advantages of *Rutin*, helping to alleviate skin redness, itching, and overall inflammation. Research conducted on animal models of dermatitis and psoriasis has demonstrated that the topical application of *Rutin*-loaded nanoparticles significantly lowers skin irritation, enhances epidermal thickness, and the amounts of inflammatory markers compared to free *Rutin* or traditional formulations ¹⁸.

4.2 *Rutin*-Loaded Nanoparticles for Age-Defiance and Dermal Shielding:

The antioxidant and photoprotective qualities of *Rutin*, combined with enhanced skin absorption via nanoparticles, position *Rutin*-loaded nanoparticles as an effective approach for safeguarding skin and combating aging. These nanoparticles can efficiently transport *Rutin* into the more profound layers of the skin, safeguarding skin cells from oxidative harm initiated by UV rays, degradation of collagen, and signs of photoaging. *Rutin*'s ability to stimulate collagen production and its antioxidant properties can aid in minimizing wrinkles, enhancing skin elasticity, and contributing to a more youthful appearance. Items containing *Rutin*-loaded nanoparticles can be blended with sunscreens and anti-aging products to boost their effectiveness and offer prolonged protection against environmental influences ¹⁹.

4.3 Rutin-Loaded Nanoparticles for Wound Healing:

The wound-healing effects of *Rutin* can be greatly enhanced through the utilization of nanoparticle delivery systems. These *Rutin*-loaded nanoparticles boost the efficiency of medication uptake at the injury location, allowing for an ongoing release of *Rutin* that encourages fibroblast growth, collagen production, and the formation of new blood vessels. Research indicates that employing *Rutin*-loaded nanoparticles. Administering therapies directly to wound models leads to faster wound recovery, improved collagen formation, and enhanced re-epithelialization in relation to treatments employing free *Rutin*. This points to the capacity of *Rutin*-loaded nanoparticles as an effective method for speeding up the healing process in chronic injuries, burns, and surgical cuts²⁰.

4.4 Rutin-Loaded Nanoparticles for Hyperpigmentation Disorders:

Rutin's ability to inhibit melanogenesis, combined with nanoparticle delivery, provides a focused strategy for addressing hyperpigmentation disorders. These nanoparticles can transport *Rutin* directly to the melanocytes in the outer skin layer, effectively preventing tyrosinase activity and diminished melanin production. The controlled release from the nanoparticles allows an extended inhibition of melanogenesis, resulting in a progressive and effective enhancement of skin luminosity. Additional research is required to assess the effectiveness of *Rutin*-loaded nanoparticles in treating specific hyperpigmentation issues such as melasma and post-inflammatory hyperpigmentation in clinical environments²¹.

5. Types of Nanoparticles for Rutin Delivery and Formulation Strategies:

A range of nanoparticles has been investigated for the encapsulation and exterior application of *Rutin*, each offering distinct characteristics and benefits:

5.1 Liposomes:

These are structures encased in membranes made up of lipid bilayers, acknowledged for their synergy with biological frameworks and their capacity for natural decomposition. They can encapsulate both hydrophilic and hydrophobic pharmaceuticals, encompassing *Rutin*. While liposomes can improve skin hydration and absorption, their stability could pose challenges.

5.2 Solid Lipid Nanoparticles (SLNs) and Nanostructured Lipid Carriers (NLCs):

Composed of solid lipids (SLNs) or a combination of solid and liquid lipids (NLCs), these

nanoparticles provide enhanced stability, increased drug loading capacity, and controlled release when compared to liposomes. They are similarly very compatible with biological systems and cost-effective for large-scale manufacturing^{22, 23}.

5.3 Polymeric Nanoparticles:

Biodegradable and biocompatible polymers like chitosan, PLGA, and hyaluronic acid can be utilized to create polymeric nanoparticles for *Rutin* encapsulation. These nanoparticles can be customized for specific drug release profiles, surface modifications for targeting, and improved stability.

5.4 Metallic Nanoparticles:

Gold and silver nanoparticles, primarily studied for their antimicrobial properties or employed as diagnostic tools, can also serve as carriers for *Rutin*. However, potential toxicity and compatibility issues must be carefully evaluated when employing metallic nanoparticles²⁴.

5.5 Nanoemulsions and Microemulsions:

These are uniform blends of oil and water that remain stable either thermodynamically or kinetically, and are maintained by surfactants. They can solubilize poorly water-soluble drugs like *Rutin* and improve skin penetration. Choosing the right type of nanoparticle and the formulation strategy depends on various factors, including the desired drug release pattern, targeted area within the skin, stability needs, biocompatibility, and manufacturing feasibility. Optimizing the nanoparticle formulation comprises elements like particle dimensions and surface characteristics, which are crucial for achieving maximum treatment efficacy and skin suitability^{25, 26}.

6. Advantages, Challenges, and Future Directions:

6.1 Advantages of Rutin-Loaded Nanoparticles:

- **Enhanced Topical Delivery:** Nanoparticles significantly improve the skin's ability to absorb *Rutin*, ensuring it reaches the intended site effectively.
- **Controlled and Sustained Delivery:** These nanoparticles can offer extended therapeutic advantages, resulting in less frequent dosing.
- **Targeted Delivery Ability:** They can be tailored for precise delivery to specific skin cells or tissues.
- **Improved Stability and Cosmetical Acceptability:** Nanoparticles enhance the uniformity of formulations and contribute to the general aesthetic appeal of topical products.
- **Reduced Systemic Exposure:** Employing topical application greatly diminishes the

chances of systemic side effects when contrasted with oral or injectable methods.

6.2. Challenges and Future Directions:

- **Scalability and Production Costs:** Developing nanoparticles attaining this on a large scale at a reasonable cost poses a significant challenge for economic viability.
- **Long-Term Consistency and Duration:** It is crucial to guarantee that nanoparticle-based formulations maintain their stability and durability to be marketable.
- **Skin Irritation and Safety Concerns:** A thorough evaluation of potential skin irritation, toxicity, and long-term safety of nanoparticles is necessary.
- **Clinical Translation:** Transitioning from preclinical studies to clinical trials and ultimately to market-ready products necessitates comprehensive clinical evaluation and regulatory approvals.
- **Personalized Dermatology:** Future research could concentrate on customizing *Rutin*-loaded nanoparticle formulations created to address distinct patient needs and resolve specific skin concerns for personalized dermatological treatments.
- **Combination Treatments:** Investigating the joint effects of *Rutin*-loaded nanoparticles, the addition of extra active elements or treatment strategies could improve the efficacy of the therapy.

Future studies should aim to address these challenges through advanced nanomaterial design, enhanced formulation methods, comprehensive biocompatibility and safety evaluations, and meticulously structured clinical trials. Additionally, exploring innovative nanomaterials and methods for altering surfaces will be crucial.

CONCLUSION:

Rutin-loaded nanoparticles offer a creative method to improve the localized management and therapeutic effectiveness of *Rutin* in skincare. By addressing the constraints of traditional topical products, these nanoparticles can efficiently transport *Rutin* to specific areas of the skin, enhancing its antioxidant, anti-inhibitory, tissue-healing, and pigment-modulating characteristics. Initial studies have shown considerable potential of *Rutin*-loaded nanoparticles in tackling various skin issues, such as inflammatory conditions, signs of aging, wounds, and hyperpigmentation. Despite current challenges pertaining to scalability, stability, and real-world implementation in medical environments, ongoing research and technological improvements are facilitating the creation of effective and clinically applicable topical

treatments using *Rutin*-loaded nanoparticles. This innovative approach has the potential to transform dermatological care and improve outcomes for those dealing with various skin conditions.

REFERENCE:

1. Kessler, M., Ubeaud, G., & Jung, L. (2003). Anti- and pro-oxidant activity of *Rutin* and quercetin derivatives. *Journal of Pharmacy and Pharmacology*, 55(1), 131–142. <https://doi.org/10.1211/002235702559>.
2. Arima, H., Ashida, H., & Danno, G. (2002). *Rutin*-enhanced Antibacterial Activities of Flavonoids against *Bacillus cereus* and *Salmonella enteritidis*. *Bioscience Biotechnology and Biochemistry*, 66(5), 1009–1014. <https://doi.org/10.1271/bbb.66.1009>
3. Melinte, V., Stroe, L., & Chibac-Scutaru, A. L. (2019). Polymer nanocomposites for photocatalytic applications. *Catalysts*, 9(12), 986. <https://doi.org/10.3390/catal9120986>
4. Qanash, H., Al-Rajhi, A. M. H., Almashjary, M. N., Basabrain, A. A., Hazzazi, M. S., & Abdelghany, T. M. (2023). Inhibitory potential of *Rutin* and *Rutin* nano-crystals against *Helicobacter pylori*, colon cancer, hemolysis and Butyrylcholinesterase in vitro and in silico. *Applied Biological Chemistry*, 66(1). <https://doi.org/10.1186/s13765-023-00832-z>
5. Javed, H., Khan, M., Ahmad, A., Vaibhav, K., Ahmad, M., Khan, A., Ashafaq, M., Islam, F., Siddiqui, Safhi, M., & Islam, F. (2012). *Rutin* prevents cognitive impairments by ameliorating oxidative stress and neuroinflammation in rat model of sporadic dementia of Alzheimer type. *Neuroscience*, 210, 340–352. <https://doi.org/10.1016/j.neuroscience.2012.02.046>
6. Shanmugasundaram, D., & Roza, J. M. (2022). Assessment of anti-inflammatory and antioxidant activity of quercetin-*Rutin* blend (SophorOx™) – an invitro cell based assay. *Journal of Complementary and Integrative Medicine*, 19(3), 637–644. <https://doi.org/10.1515/jcim-2021-0568>
7. Shanmugasundaram, D., & Roza, J. M. (2022). Assessment of anti-inflammatory and antioxidant activity of quercetin-*Rutin* blend (SophorOx™) – an invitro cell based assay. *Journal of Complementary and Integrative Medicine*, 19(3), 637–644. <https://doi.org/10.1515/jcim-2021-0568>
8. Chen, L., Huang, C., Liao, C., Chang, H., Kuan, Y., Tseng, T., Yen, K., Yang, K., & Lin, H. (2020). Effects of *Rutin* on wound healing in hyperglycemic rats. *Antioxidants*, 9(11), 1122. <https://doi.org/10.3390/antiox9111122>
9. Sun, X., Zhang, Z., Chen, C., Wu, W., Ren, N., Jiang, C., Yu, J., Zhao, Y., Zheng, X., Yang, Q., Zhang, H., Li, J., & Li, Z. (2018). The C-S-A gene system regulates hull pigmentation and reveals evolution of anthocyanin biosynthesis pathway in rice. *Journal of Experimental Botany*, 69(7), 1485–1498. <https://doi.org/10.1093/jxb/ery001>
10. Martins, R. M., De Siqueira Martins, S., Barbosa, G. L. F., Fonseca, M. J. V., Rochette, P. J., Moulin, V. J., & De Freitas, L. a. P. (2022). Photoprotective effect of solid lipid nanoparticles of *Rutin* against UVB radiation damage on skin biopsies and tissue-engineered skin. *Journal of Microencapsulation*, 39(7–8), 668–679. <https://doi.org/10.1080/02652048.2022.2156631>
11. Sharma, A., Comejo, C., Mihalic, J., Geyh, A., Bordelon, D. E., Korangath, P., Westphal, F., Gruettner, C., & Ivkov, R. (2018). Physical characterization and in vivo organ distribution of coated iron oxide nanoparticles. *Scientific Reports*, 8(1). <https://doi.org/10.1038/s41598-018-23317-2>
12. Lu, B., Huang, Y., Chen, Z., Ye, J., Xu, H., Chen, W., & Long, X. (2019). Niosomal Nanocarriers for Enhanced Skin Delivery of Quercetin with Functions of Anti-Tyrosinase and Antioxidant. *Molecules*, 24(12), 2322. <https://doi.org/10.3390/molecules24122322>
13. Carvalho, L., Bezerra, C., Da Rocha, C., De Oliveira, L.,

- Vasconcelos, L., & Santana, S. (2023). Controlled Release of Rutin from Babassu Coconut Mesocarp Starch Films. *Journal of the Brazilian Chemical Society*. <https://doi.org/10.21577/0103-5053.20230045>
14. Sengupta, P., Das, D., Bhattacharya, S., Sur, R., Bose, A., & Sen, K. (2023). A pH-driven method for liposomal encapsulation of dietary flavonoid Rutin: Sustained release and enhanced bioefficacy. *Food Bioscience*, 52, 102392. <https://doi.org/10.1016/j.fbio.2023.102392>
 15. Gayoso, L., Claerbout, A., Calvo, M. I., Caverio, R. Y., Astiasarán, I., & Ansorena, D. (2016). Bioaccessibility of Rutin, caffeic acid and rosmarinic acid: Influence of the in vitro gastrointestinal digestion models. *Journal of Functional Foods*, 26, 428–438. <https://doi.org/10.1016/j.jff.2016.08.003>
 16. Popović, B. M., Uka, D., Alioui, O., Pavlović, R. Ž., & Benguerba, Y. (2022b). Experimental and COSMO-RS theoretical exploration of Rutin formulations in natural deep eutectic solvents: Solubility, stability, antioxidant activity, and bioaccessibility. *Journal of Molecular Liquids*, 359, 119266. <https://doi.org/10.1016/j.molliq.2022.119266>
 17. Mechchate, H., Es-Safi, I., Haddad, H., Bekkari, H., Grafov, A., & Bousta, D. (2020). Combination of Catechin, Epicatechin, and Rutin: Optimization of a novel complete antidiabetic formulation using a mixture design approach. *The Journal of Nutritional Biochemistry*, 88, 108520. <https://doi.org/10.1016/j.jnutbio.2020.108520>
 18. Souto, E. B., Fernandes, A. R., Martins-Gomes, C., Coutinho, T. E., Durazzo, A., Lucarini, M., Souto, S. B., Silva, A. M., & Santini, A. (2020). Nanomaterials for skin delivery of cosmeceuticals and pharmaceuticals. *Applied Sciences*, 10(5), 1594. <https://doi.org/10.3390/app10051594>
 19. Souto, E. B., Fernandes, A. R., Martins-Gomes, C., Coutinho, T. E., Durazzo, A., Lucarini, M., Souto, S. B., Silva, A. M., & Santini, A. (2020b). Nanomaterials for skin delivery of cosmeceuticals and pharmaceuticals. *Applied Sciences*, 10(5), 1594. <https://doi.org/10.3390/app10051594>
 20. Sharma, Y., Bhardwaj, R., Kaur, A., Singh, G., Kulkarni, S., Bhatia, A., & Bala, K. (2024). Nano-gel formulation of polyphenolic fraction of tobacco stem for wound healing and its inhibitory efficacies against the receptors of chronic wound development. *Journal of Biomolecular Structure and Dynamics*, 1–15. <https://doi.org/10.1080/07391102.2024.2317986>
 21. Silva, A. R., Taofiq, O., Ferreira, I. C., & Barros, L. (2020). Hypericum genus cosmeceutical application – A decade comprehensive review on its multifunctional biological properties. *Industrial Crops and Products*, 159, 113053. <https://doi.org/10.1016/j.indcrop.2020.113053>
 22. Perumalla, A., & Hettiarachchy, N. S. (2011). Green tea and grape seed extracts — Potential applications in food safety and quality. *Food Research International*, 44(4), 827–839. <https://doi.org/10.1016/j.foodres.2011.01.022>
 23. Pandian, S. R. K., Pavadai, P., Vellaisamy, S., Ravishankar, V., Palanisamy, P., Sundar, L. M., Chandramohan, V., Sankaranarayanan, M., Panneerselvam, T., & Kunjiappan, S. (2020). Formulation and evaluation of Rutin-loaded solid lipid nanoparticles for the treatment of brain tumor. *Naunyn-Schmiedeberg Archives of Pharmacology*, 394(4), 735–749. <https://doi.org/10.1007/s00210-020-02015-9>
 24. Wu, H., Su, M., Jin, H., Li, X., Wang, P., Chen, J., & Chen, J. (2020). Rutin-Loaded silver nanoparticles with antithrombotic function. *Frontiers in Bioengineering and Biotechnology*, 8. <https://doi.org/10.3389/fbioe.2020.598977>
 25. Sharma, S., Rabbani, S. A., Narang, J. K., Potttoo, F. H., Ali, J., Kumar, S., & Baboota, S. (2020). Role of Rutin nanoemulsion in ameliorating oxidative stress: Pharmacokinetic and Pharmacodynamics studies. *Chemistry and Physics of Lipids*, 228, 104890. <https://doi.org/10.1016/j.chemphyslip.2020.104890>
 26. Kajbafvala, A., & Salabat, A. (2021b). Microemulsion and microemulsion gel formulation for transdermal delivery of Rutin: Optimization, in-vitro/ex-vivo evaluation and SPF determination. *Journal of Dispersion Science and Technology*, 43(12), 1848–1857. <https://doi.org/10.1080/01932691.2021.1880928>