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A Comprehensive Review on Silver Nanoparticles

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Abstract

Herbal drugs are herbal goods that are fully plantbased. Phytopharmaceuticals are made of a huge number of ingredients, making it impossible to study one ingredient at a time. It is important to understand the chemical properties of herbal products, botanical phytochemicals in order to conduct relevant research. Plant-pharmaceuticals are regarded as medicinal products. A diverse assortment of botanical therapies is often gone by various names, including phototherapeutics, photomedicine, and photomedicine. Healing compounds and extracts can be used in the plant-based method. Leaves, bark, rhizomes, roots, rhizomes and seedlings are commonly used materials. This short review is going to express the key note on Nanophytopharmaceuticals.

Keywords: Phytochemicals, Nanopharmaceuticals, Phytomedicine, Plant-pharmaceuticals.

1 Introduction

Phytocannabinoids are found in dried plant materials, or extracted from parts of plants. Herbal medications are quickly and rapidly metabolized by the body's chemistry. As a consequence, their performance is reduced or their adverse effects are avoided. "Chem" medications, on the other hand, are potentially toxic. There should also be inclusion of the drug's active principles with other phytochemicals, which improve or impede the actions of the drug in different ways¹. The relationship between frequently-used medications and dosages does not always match. The method of further processing yields dried extracts, powders, tablets, and other dosage forms. They are often called intermediate or semi-finished products, and can be further processed to yield higher technological improvements. Gum tragacanthin and deexamylated acacia are both defined as phytopharmaceuticals^{2,3}. A developis applied to discover the existence ment and/quantification of a substance, to ascertain the composition or structure of a solution or mixture, or to determine the molecular makeup⁴.

The World Health Organization (WHO) has made it possible to justify the concept of phytomedicine as the practice of herbs, herbal ingredients, herbal formulations and manufactured herbal extracts which comprises mainly therapeutically active, plant-insulating phytocomponents or combinate plant extracts⁵. In general, these herbal components, including roots, stalks, blooms, bulbs, seeds or by-products (gum, resins and much else), are isolated in the plant parts⁶. According to the literature and the studies published, nearly 50,000 plant species exhibit medicinal activity. Roughly 80% of the world's populations, particularly in developing countries, are still the first-choice medicinal products for the management of a variety of illnesses and therefore play a vital role in today's health wellness. Furthermore, as opposed to other traditional drugs, both phytomedicines and doctors have been able to improve bioavailability due to their imminent therapeutic effects and less harmful effects. Phytomedicine-based treatment can effectively be used to prevent and treat different malignancies as they concentrate primarily on natural chemical entities from different plant components. In comparison to chemotherapy, the health outcomes of sick people have improved dramatically, with decreased adverse effects⁷⁻⁹.



Fig 1. Nanophytomedicines for treating various disorders

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In qualitative analysis, you learn about the composition or the number of species and in quantitative analysis you learn about the number of components. As the NPs have excellent solubility and biological availability, these drugs tend to remain in the blood stream longer. Medicinal nanotechnology has the potential to solve other forms of plant problems¹¹.

Nanophytomedicines and Their Bioavailability Studies

Herbal medications have been used in many cultures all over the world for different illnesses because of their comparatively benign side effects as opposed to today's pharmaceuticals. Biologically effective plant-derived chemicals have already been discovered in the field of phytotherapy. Tannins, i.e., flavonoids, and terpenoids showed a poor absorption in water. They also have high molecular weights, which limited their bioavailability and absorption. due to the increased molecular weight and low solubility, most plant medicines were found to be poorly absorbed by fat cells, and were thus ineffective¹²⁻¹⁵.

The recent progress in the nanoherbalization of the herb's bioavailability and efficacy have made it more difficult for herbs to be water-soluble. reviews of Apigenin, e.g., and taxifolin and curcumin have been done Since the delivery method has a large effect on the amount of a drug that gets into the bloodstream, the nanomedicines' advantages in the increasing solubility and accessibility are impressive¹⁶.

Various Nanophytomedicines

Clinical research results were obtained with a camptothecin, which is derived from the Camptothecaacuminata stem tissues. The anticancer efficacy in a rat lung cancer model was found to be greatly enhanced through the pulmonary administration of this drug an orthoptic animal model of lung cancer (naked rats) was successfully applied in this research, and they were able to produce tumour about one hundred percent of the time. Fivefold increase in plasma concentrations were found in rats when 9-nitaminoicotinotride polymeric nanoparticles were used as a model for cytotoxicity in a trial¹⁷. An extract rich in flavonoids, is Silymarin. In a rat model, it was achieved to promote antioxidant activity as well as cleanse the internal GTH stores. A significant advantage of its position as the hepatocyte membrane stabilizer and permeability regulator was that it shielded hepatotoxins from getting into the hepatocytes. Translation will encourage the development of ribosomal RNA in the same experiment¹⁸.

However, the solubility of this drug is extremely low in water, necessitating the use of an acidic medium. The solubility of the drug in polyvinylpyrrolidone nanoparticles was roughly three times as great as in its non-loaded commercial unmodified form (Culbokon®). They developed a hepatotoxicity-related pathway; however, no bioavailability studies were conducted on the animal model. It was loaded into polymers to enhance the bioavailability. Danshen, a highly regarded Chinese plant medication, is derived from Radix salvia. Cardiac activities have been observed in the course of this nanomedical treatment. Using a highspeed centrifugal extractor, this herbal product's solubility and bioavailability improved markedly. Another concern is that my study has addressed is the consistency of drug release. This plant medicine was tested in volunteers with no disease or nutritional deficiencies, and found to have a fivefold increase in its bioavailability when micronized¹⁹.

The extract from PhylliumSchumacharie and Thulium has historically been used to improve liver function. This plant medicine is in its current form requires an excessive amount of the extract and lengthy treatment durations. To be administered for an extended period of time, the ethanolic extract of this plant was incorporated in sodium alginate for better absorption and uptake in the intestines. A rodent model was used to investigate the hepatoprotective behavior of a nanoparticle ethanolic extract of Amargosa spinosa (E=Phenolic; P=NPA). The oral NPA administration was found to be more effective and safer than PA that had lower levels of hepatotropic activity. Cuscuta is named after the species from which it is derived. Both the type of this medication, which is taken orally, has a low solubility and slow absorption rate. The solubility of this drug has been enhanced in nanoparticle form²⁰. This will improve liver function while at the same time helping the cells resist oxidation and scavenging free radicals. The hepatoprotrophic plant's toxicity was improved while its ethanolic extract was not bioavailable to rats; however, no biological activity was detected²¹.

Nano-Phytomedicine for Improved Target Ability

Minority groups, who are in desperate need of care, may be targeted higher the target concentration of the drug, the better the results. These mechanisms concenpharmacologicalactive/active trate the ly/locally/specifically and prevent the accumulation elsewhere. They transport, retain, or sequester the drug in target cells. Specific active locations may be an area of focus, a capillary bed, a specific cell type of cell, or an intracellular location. To achieve an objective, a carrier must be constructed that contains an agent that will only recognize the chosen target. To keep cancer from recurring, the transporter must be able to identify cancerous cells, rather than only normal cells²².

Ligands assist in recognizing the targets, and facilitate targeted delivery. Exogenous or naturally occurring hormones are some of the examples of ligands. A primary goal of the medications is disease control or prevention. Many pharmaceutical factors have been selected when designing delivery systems for drugs with these characteristics, including instability in the stomach, low solubility, large volume of distribution, high plasma protein binding, short biological half-life, low specificity, pharmacokinetic/dynamic, pharmacodynamic, short half-life, pharmacodynamic and low therapeutic index²³. The principal objective of this strategy is to reduce side effects on other tissues. In order to be efficient, delivery systems should be biologically safe, non-immunogenic, not distributable, and stay within the target tissue, be physical, restricted, and exhibit stability²⁴. The control should be able to provide an optimal dose of the drug, and the results should be welldefined. Carriers are made to minimize the presence of drugs when in transit. It is as critical if polymers used for the production of carriers are biodegradable and readily excreted by the body, even if they are not so in your body. They should be simple to make or already on hand, and cheap to purchase. The specificity of these isolators, the design of the markers, or the efficiency of transporting of the markers are taken into consideration when isolators are created²⁵⁻²⁷.

Phytosome

The efficacy of any herbal drug depends on the potency of the medicinal active compound. When administered orally or topically they are seriously restricted in their bioavailability. Phytosome formulations that are more absorbed than extracts have been launched recently. "Phyto" is a plant and "some" a cell-like term. Over the past century, the nature, biologic activity and health sciences of several botanicals have been created in the field of phytochemistry and phytopharmacology. Polar or water-soluble molecules are mostly biologically active ingredients of plants. However, water-losablePhyto-constituents (such as flavonoid substances, tannins, glycosidic aglycons, etc.) are not properly absorbed either by their large molecule sizes, which cannot be absorbed by passive diffusion, or due to their low lipid solubility. Pharmacokinetic (tissue distribution) and activity experiments in animals and humans have shown that the phytosomes have increased bioavailability over easier, non-complex plant extracts. The established health behavior of the phospholipids itself has an additional dimension to phytosomes. Often known as Herbosomes 5, Phytosome is also referred to. The profile of phytosomes is stronger than traditional herbal extracts for pharmacokinetic and pharmagodynamic use²⁸⁻³⁰.

Herbosome

The phyto-chemical extracts from the flavonoid and terpenoids include phosphatidylcholine for the direct complex. Herbosome results from a reaction to the standardize extract or polyphenolic constituents in a non-polar solvent with stoichiometric phospholipid (phosphatidylcholine) amounts. Phosphatidylcholine (phosphatidyl) is a bi-functional movement which is lipophilic and the hydrophilic motivation is the choline movement. In particular, the phosphorous choline head binds to these compounds while the body and tail section, which then envelops the choline-bound content, is a lipid soluble phosphatidyl component. Thus, a lipid molecular complex consistent with phospholipids, also called a complex Phyto phospholipid, constructs the Phytoconstituents³¹.

Ethosomes

Ethosomes are vesicles consisting of high ethanol phospholipids. Due to the high levels of ethanol in the ethosomes, the fluidisation of the skin lipids improves their permeability through your skin. These carriers can penetrate the skin in depth and even into blood circulation, contributing to increased medicines. Triptolide ethosomes are used to supply the medication topically. In comparison to other formulations, the ethosomal formulation showed a greater bioavailability among rats due to increased accumulation and decreased erythema³².

Dendrimers

Dendrimers are extremely ramified, inner-core nanostructures. Drugs are integrated in the inner center both covalently and in electrostatic mode with the branched surface. Dendrimers, however, are affected by various limitations, such as poor/unstable hydrophobic drug loadings and ineffective drug release. Recent evolution to address this issue is new class of molecules known as dendronized polymers which are linear polymers that have dendrons on each repetitive unit. Another solution is the use of dendrimers with a degradable connection to monitor release of the drug. Chang et al, for example, are preparing a medication releases device based on folic acid (FA) conjugated to doxorubicin (DOX) and superparamagnetic iron oxide poly (ethylene glycol) (PEG)-modified dendrimers (PAMAM)³³.

Conclusion

The composition of the mixture needs to be determined when creating new herbal products containing blends other than pure contents, which express the optimal individuality for which the substance has been produced. Phytopharmaceuticals have been generally recognized as both medical and dietary agents. In nanotechnology and nanoproduct clinical trials, the safety, pharmacokinetics and ethics of human subjects also present new and specific problems. While nanotechnologies are able to overcome the limitations of traditional approaches, they face problems. But more challenges are still required to be overcome. Nano-structured systems of delivery have enormous impending in the provision of phytomedicine for therapeutic purposes. This is because they have enormous solutions, pharmacokinetic parameters such as absorption, delivery, metabolism and excretion, bioavailability, objective capacity, effectiveness and protection. They also use different medicinal advantages themselves. Phytomedicines along with therapeutic nanocarriers for their double effect have also been studied. The invention of nanocarrier systems based on phytomedicine has been done to resolve the numerous unsatisfactory built to date can be found that help phytomedicines to the nearest site due to their unique ability to alter physical and chemical properties.

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References

 J. W. Alexander, "History of the medical use of silver," Surgical Infections, vol. 10, no. 3, pp. 289– 292, 2009.

- [2] K. M. M. Abou El-Nour, A. Eftaiha, A. Al-Warthan, and R. A. A. Ammar, "Synthesis and applications of silver nanoparticles," Arabian Journal of Chemistry, vol. 3, no. 3, pp. 135–140, 2010.
- [3] P. Mohanpuria, N. K. Rana, and S. K. Yadav, "Biosynthesis of nanoparticles: technological concepts and future applications," Journal of Nanoparticle Research, vol. 10, no. 3, pp. 507–517, 2008.
- [4] S. Poulose, T. Panda, P. P. Nair, and T. Theodore, "Biosynthesis ´ of silver nanoparticles," Journal of Nanoscience and Nanotechnology, vol. 14, no. 2, pp. 2038–2049, 2014.
- [5] M. Vijayakumar, K. Priya, F. T. Nancy, A. Noorlidah, and A. B. A. Ahmed, "Biosynthesis, characterisation and anti-bacterial effect of plantmediated silver nanoparticles using Artemisia nilagirica," Industrial Crops and Products, vol. 41, no. 1, pp. 235–240, 2013.
- [6] J. G. Parsons, J. R. Peralta-Videa, and J. L. Gardea-Torresdey, "Chapter 21 Use of plants in biotechnology: synthesis of metal nanoparticles by inactivated plant tissues, plant extracts, and living plants," Developments in Environmental Science, vol. 5, pp. 463–485, 2007.
- [7] M. Ghaffari-Moghaddam, R. Hadi-Dabanlou, M. Khajeh, M. Rakhshanipour, and K. Shameli, "Green synthesis of silver nanoparticles using plant extracts," Korean Journal of Chemical Engineering, vol. 31, no. 4, pp. 548–557, 2014.
- [8] M. A. Faramarzi and A. Sadighi, "Insights into biogenic and chemical production of inorganic nanomaterials and nanostructures," Advances in Colloid and Interface Science, vol. 189- 190, pp. 1–20, 2013.
- [9] S. Azizi, F. Namvar, M. Mahdavi, M. B. Ahmad, and R. Mohamad, "Biosynthesis of silver nanoparticles using brown marine macroalga, Sargassum muticum aqueous extract," Materials, vol. 6, no. 12, pp. 5942–5950, 2013.
- [10] T. N. V. K. V. Prasad, V. S. R. Kambala, and R. Naidu, "Phyconanotechnology: synthesis of silver nanoparticles using brown marine algae Cystophora moniliformis and their characterisation," Journal of Applied Phycology, vol. 25, no. 1, pp. 177–182, 2013.
- [11] R. Das, S. S. Nath, D. Chakdar, G. Gope, and R. Bhattacharjee, "Synthesis of silver nanoparticles and their optical properties," Journal of Experimental Nanoscience, vol. 5, no. 4, pp. 357–362, 2010.
- [12] M. Gilaki, "Biosynthesis of silver nanoparticles using plant extracts," Journal of Biological Sciences, vol. 10, no. 5, pp. 465–467, 2010.
- [13] R. Vaidyanathan, K. Kalishwaralal, S. Gopalram, and S. Gurunathan, "Nanosilver-the burgeoning therapeutic molecule and its green synthesis," Biotechnology Advances, vol. 27, no. 6, pp. 924–937, 2009.

- [14] V. K. Sharma, R. A. Yngard, and Y. Lin, "Silver nanoparticles: green synthesis and their antimicrobial activities," Advances in Colloid and Interface Science, vol. 145, no. 1-2, pp. 83–96, 2009.
- [15] A. M. Fayaz, K. Balaji, M. Girilal, R. Yadav, P. T. Kalaichelvan, and R. Venketesan, "Biogenic synthesis of silver nanoparticles and their synergistic effect with antibiotics: a study against gram-positive and gram-negative bacteria," Nanomedicine: Nanotechnology, Biology, and Medicine, vol. 6, no. 1, pp. e103– e109, 2010.
- [16] R. Singh and D. Singh, "Chitin membranes containing silver nanoparticles for wound dressing application," International Wound Journal, vol. 11, no. 3, pp. 264–268, 2014.
- [17] G. Habiboallah, Z. Mahdi, Z. Majid et al., "Enhancement of gingival wound healing by local application of silver nanoparticles periodontal dressing following surgery: a histological assessment in animal model," Modern Research in Inflammation, vol. 3, no. 3, pp. 128–138, 2014.
- [18] D. Nambiar and Z. P. Bhathena, "Use of silver nanoparticles from Fusarium oxysporum in wound dressings," Journal of Pure and Applied Microbiology, vol. 4, no. 1, pp. 207–214, 2010.
- [19] J. Tian, K. K. Y. Wong, C.-M. Ho et al., "Topical delivery of silver nanoparticles promotes wound healing," ChemMedChem, vol. 2, no. 1, pp. 129–136, 2007.
- [20] J. Kaur and K. Tikoo, "Evaluating cell specific cytotoxicity of differentially charged silver nanoparticles," Food and Chemical Toxicology, vol. 51, no. 1, pp. 1–14, 2013.
- [21] A. Schrofel, G. Krato "`sova, I. 'Safa ``r´ık, M. Safa ``r´ıkova, I. Ra ´`ska, and L. M. Shor, "Applications of biosynthesized metallic nanoparticles—a review," Acta Biomaterialia, vol. 10, no. 10, pp. 4023–4042, 2014.
- [22] N. Kulkarni and U. Muddapur, "Biosynthesis of metal nanoparticles: a review," Journal of Nanotechnology, vol. 2014, Article ID 510246, 8 pages, 2014.
- [23] O. V. Kharissova, H. V. Rasika Dias, B. I. Kharisov, B. O. Perez, ´ and V. M. J. Perez, "The greener synthesis of nanoparticles," ´ Trends in Biotechnology, vol. 31, no. 4, pp. 240–248, 2013.
- [24] A. K. Mittal, Y. Chisti, and U. C. Banerjee, "Synthesis of metallic nanoparticles using plant extracts," Biotechnology Advances, vol. 31, no. 2, pp. 346–356, 2013.
- [25] S. Prabhu and E. K. Poulose, "Silver nanoparticles: mechanism of antimicrobial action, synthesis, medical applications, and toxicity effects," International Nano Letters, vol. 2, article 32, 2012.
- [26] V. Kumar and S. K. Yadav, "Plant-mediated synthesis of silver and gold nanoparticles and

their applications," Journal of Chemical Technology and Biotechnology, vol. 84, no. 2, pp. 151–157, 2009.

- [27] V. Arya, R. Komal, M. Kaur, and A. Goyal, "Silver nanoparticles as a potent antimicrobial agent: a review," Pharmacologyonline, vol. 3, pp. 118– 124, 2011.
- [28] T. M. Tolaymat, A. M. El Badawy, A. Genaidy, K. G. Scheckel, T. P. Luxton, and M. Suidan, "An evidence-based environmental perspective of manufactured silver nanoparticle in syntheses and applications: a systematic review and critical appraisal of peer-reviewed scientific papers," Science of the Total Environment, vol. 408, no. 5, pp. 999–1006, 2010.
- [29] C. Marambio-Jones and E. M. V. Hoek, "A review of the antibacterial effects of silver nanomaterials and potential implications for human health and the environment," Journal of Nanoparticle Research, vol. 12, no. 5, pp. 1531– 1551, 2010.
- [30] K. K. Y. Wong and X. Liu, "Silver nanoparticlesthe real silver bullet in clinical medicine," MedChemComm, vol. 1, no. 2, pp. 125–131, 2010.