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### A REVIEW: - NOVEL HERBAL DRUG DELIVERY SYSTEM AND ITS APPLICATION

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### **ABSTRACT:**

Herbal medicine is closer to traditional medicine than other traditional or alternative medicine. They contain many ingredients that are known for their healing properties and have been used for centuries. Conventional products have unique advantages, such as rich treatments and their many chemical and organic effects, when used by scientists to create new medicines. Scientists have isolated, characterized and analyzed the chemical activity of drug molecules from various herbal sources. However, plant molecules must be properly designed to support their goals in the body and provide better results. This evaluation should• examine data on natural preparations for further development of existing and new green medicine and examine methods that can deliver them to order

**KEYWORLD :-** *Natural goods; liposomes; phytosomes nanoparticles; bioactive compounds; new herbal drug delivery systems (NHDDS)* 

## **INTRODUCTION**:

Natural remedies have been found in nature Product with many medicinal properties mad e from plant products for centuries. Medicinal plants are increasingly used to improve health and well- being and can equally be used alone or in combination with existing medications to treat medical problems<sup>[1]</sup>. It is an important part of the new lead model for different treatments Novel delivery systems (NHDDS) are important for the delivery of bioactive molecules to active sites. Many new types of herbal medicines, such as polymeric nanoparticles, liposomes, phospholipases, Nano emulsions, microspheres, transsomes, and , are thought to use bioactive atoms. In fact, NHDDS is a very powerful tool to d deliver herbal medicines through molecules, and natural product scientists are working to choose the best delivery method.<sup>[2]</sup> As new As new medication delivery methods are developed, it is more crucial than ever to quickly identify those that work. NHDDS represents a recent innovation in the field of herbal medicine. This review discusses natural materials and how various NHDDS encapsulations are used to distribute them.<sup>[3]</sup> The knowledge gleaned from this article will assist in selecting the best delivery method for compounds produced from PLANTs.

# The source of a plant's bioactive substances :

based structure is still widely used and maintained in many communities, and it plays a significant role in the provision of healthcare. Response to the author of this articles university Institute of pharmaceutical Sciences, Turn to nature to cure human illnesses as long-term usage of manufactured medications might result in numerous adverse effects.<sup>[4]</sup> Herbal remedies' intricacy, structural variety, great selectivity, and distinct biological activity drugs derived from molecules make them interesting for drug production.<sup>[5]</sup> Although it is a small part Since 1805It has been confirmed that morphine, the first chemical drug, is recorded for allergies and that this is the poison of the plant. The plant has many dynamic components that can be valuable in improving purification.<sup>[6]</sup> Knowledge of phytochemicals and the evidence and isolation of individual substances in medicinal plants or plant materials are important for the emergence of peace. According to the analysis, there are at least 15 important phytochemicals in plants, such as Each category has distinct chemical features, including flavonoids, alkaloids, glycosides, essential oils, tars, phytochromes, natural acids, amino acids, tannins, proteins, compounds, and the following: polysaccharides, mineral salts, etc.

[7].

A combination of aggravating substances may be present in the natural preparation. Therefore, it is important to use this combination to improve herbal medicine ability.<sup>[8]</sup> Plants are not yet used to treat distinct illnesses, but they provide a solid foundation for finding bioactive compounds that may be used to generate new drugs.

# Justifications for Novel Plant-Derived Molecule Drug Delivery Systems. :-

The medication's therapeutic and bioavailability at the site of action determine its pharmacological impact.

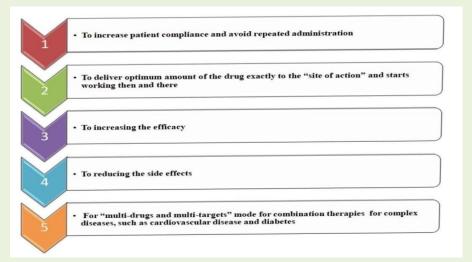


Fig.No 1:-Explains the rationale behind the unique medication delivery method for herbal origin compounds

Thus, the genesis and growth into a sensible natural detail for the transportation of certain bioactive particles are crucial.<sup>[9]</sup> The development of a novel treatment for herbal remedies has received a lot of attention in the few preceding decades<sup>[10]</sup>. The analysts in the field of natural preparation deal with problems such as inadequate utilization of the starting point of herbal particles. strength in the stomach, a high level of first-pass metabolism, inability to dissolve fats easily, or an incorrect atomic size or both, resulting in low bioavailability and retention, and so on. Even if phyto compounds move remarkably in vitro, they show little to no activity in vivo. In this sense, NHDDS takes on a big role in overcoming these problems.<sup>[11]</sup> These approaches have the potential to increase the patient's consistency. Measurement frameworks have advanced along with medicine delivery systems have evolved from simple mixtures and tablets to incredibly sophisticated innovations, concentrated medicine delivery systems, or NHDDSs<sup>[12]</sup>. These devices can administer the medication at the necessary pace and for a prolonged amount of time by channeling the bioactive compounds derived from herbs to the site of action. Consequently, NHDDS has a bright future in enhancing therapeutic activity and resolving issues supported by natural medications.

## Distinctive Herbal Medicine Delivery Systems (NHDDS) :-

Dosage formulations, including sustained release, cannot maintain the medicinal ingredients in the different proportions needed by the body during the healing period, and Botanical For maximum healing response components are taken to their targets<sup>[13]</sup>. Herbal preparation is a quantitative preparation containing at least one plant or herbs processed in a certain amount to have health benefits, healing and different qualities, for the purpose of analysis, treatment or modification in the person's body.<sup>[14]</sup> To improve the effectiveness of herbal medicine, polymer innovation, pharmaceuticals, immunology, subatomic science, etc. New drug delivery systems, which are the integration of many research areas such as, play an important role.<sup>[15]</sup> The underlying recommendation of NDDS includes a focus on sedative delivery, which can reduce dose recovery and improve solubility and uptake while reducing clearance (Figure 2). Various NDDS Drug delivery methods have been employed, such as various organisational structures for regulated and focused sedative administration. Numerous

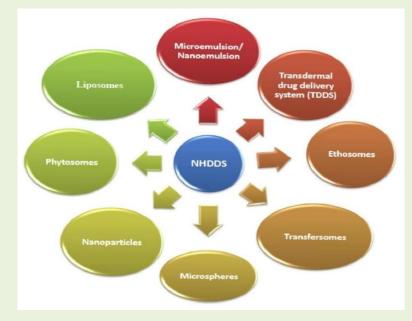
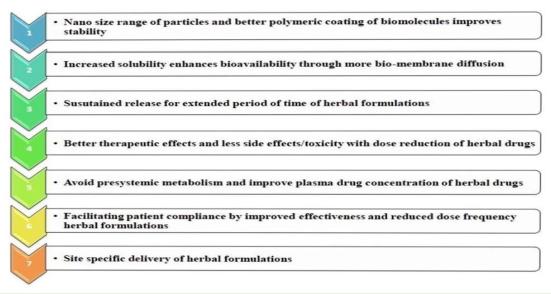


Fig 2: An overview of the use NHDDS

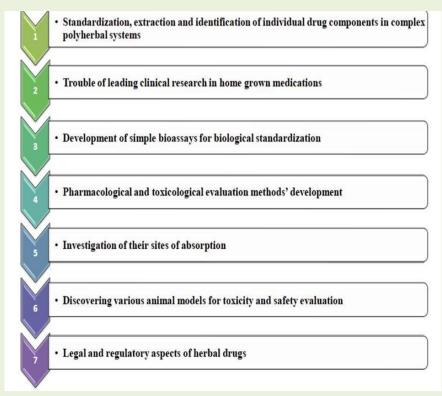
NHDDS are employed to transfer herbal medication from molecules to the site of action, as seen in Figure (2). Table 1 provides an overview of the use of NHDDS for the delivery of therapeutic compounds in various disorders. Here, an attempt has been made to investigate the varied benefits of various NHDDS (Figure 1). 3. One of the special qualities of liposomes is their ability to store material.<sup>[16]</sup>



### Fig. 3: Quiet characteristics of NHDDS

Phytosome is a patented technology that encapsulates standardized extracts or botanical ingredients into phospholipids to create molecular complexes that enhance bioavailability, permeability, and stability—particularly when there is a significant propensity for selfaggregation and good solubility characteristics.<sup>[17]</sup> Little dosages are needed since it improves the active components' absorption. Additionally, the phospholipid complex improves the solubility of herbal constituents in bile and their absorption, enabling the liver to receive the herbs.<sup>[18]</sup> Among the many benefits of nanoparticles are their ability to increase biomolecules' solubility, decrease drug intake, improve plant absorption, and encapsulate medications within nanoparticles to deliver them straight to the site of action.<sup>[19]</sup>

Drugs or Compounds (Class)	Plant Origin (Part)	Activity	Drug Delivery System	Chemical Structure	Ref
Glycyrrhizie acid (Saponín)	Glycyrrhiza gla- bra Family:Fabaceae (Root)	Anti- hypertensive, Anti- inflamma- tory	Nanoparticle	-topper	[37, 3
Hypocrellins (Pigments)	Shiraia bambusi- cola Family: Hypocreaceae (Fruit)	Antiviral	Nanoparticle		[39, 4
Magnolol (Lignan)	Magnolia officinalis Family: Magnoliaceae (Bark)	Vascular smooth muscle proliferation inhibition	Liposome		[41, 42
Matrine (Alkaloid)	<i>Sophora</i> <i>flavescens</i> Family: Fabaceae (Root)	Anti- inflammatory, Anticancer, Anti- rheumatism, Antibacterial, Antioxidant	Ethosomes, Nancemulsion		[43-45
Naringenin (Trihydroxy fla- vanone)	Lycopersicum esculentum Family: Solanace- ae (Fruit)	Anticancer, Anti- inflammatory, Hepato -protactive	Phytosome, Nanoparticle		[46]
Oxymatrine (Quinolizidinalkaloid)	Sophora Jlavescens Family: Fabaceae (Root)	Antiviral	Phytosome		[45, 47]



#### **Figure 4: Difficulties in Formulating Herbal Medicines**

It is simple to alter the characteristics of nanoparticles to accomplish precise and regulated medication administration<sup>[20]</sup>. Hydrophilic and hydrophobic compounds may be delivered with efficiency using nanoparticles. Alcohol creates a platform for the transportation of various herbs and aids in the penetration of medications via the skin.<sup>[21]</sup> In certain partial models, alcohol has been utilised to increase patient compliance<sup>[10]</sup>. Because of the buildup and decrease of erythema, the body's ethanol concentration exhibits an experiment in bioavailability, with a faster rate of rise in bioavailability than for other contents.<sup>[22]</sup> It is advantageous for the local dispersion of alkaloids and also makes the stratum conium more permeable.<sup>[23]</sup> The advantages of transdermal drug delivery systems (TDDS) include simplicity of use, fewer adverse effects, enhanced bioavailability, and regulated drug administration.<sup>[24]</sup> With less than zero-level kinetics, this distribution prevents firstpass metabolism and allows for sustained drug delivery and consumption<sup>[25]</sup> Artefacts that mimic a human body are called transformation body vectors. Transdermal medication delivery issues are mostly caused by physical changes, such as the difficulty to transport big molecules, the cost-limiting step of penetration into the stratum conium, and the physicochemical properties of the child's reaction via the skin.<sup>[26]</sup> These pliable vesicles have the ability to pass through skin pores, which are frequently smaller than usual, and carry heavier molecules <sup>[27]</sup> 5. Obstacles in the Medical Field The development of herbal bio-molecular formulations remains a challenge. Designing a suitable conveyor system has some challenges.

Lipid emulsions are the term for sub-micro emulsions, while micro emulsions are also known as nano emulsions.<sup>[28]</sup> Because of its interaction with lymph fluid, the cream is dispersed to different regions of the body in accordance with drug distribution theory.<sup>[29]</sup> Herbal remedies in emulsion form improve the stability of hydrolyzed materials, intensify the effects of medications on the skin and mucosa, and lessen the effects of medications on tissues. It can eliminate the medication over an extended period of time since it becomes lodged inside and travels to various bodily tissues.<sup>[30-32]</sup> Because lipophilic medications generate O/W/O emulsions, the oil droplets are taken up by macrophages and concentrate in the kidneys, spleen, and liver, where they become valuable after a break. It was a triumph Water-soluble medications

are designed as W/O/W emulsions, but when injected intramuscularly or subcutaneously, they show promise in the lymphatic system.<sup>[33]</sup> Because microspheres may be swallowed or infused, altered to provide useful release profiles, and utilised for the transport of biomolecules, the organisation of microsphere formulations is essential. It can guarantee the drug's special properties and release the medication into the external phase for widespread dispersion.<sup>[34]</sup> Years ago, reports of magnetic microspheres and immunological microspheres also surfaced.

Because polymer microspheres are coated or adsorbed with antibodies and antigens, immune microspheres offer protection against illness.<sup>[35]</sup> Alcohol facilitates the penetration of drugs through the skin and provides a platform for the transport of many different herbs. Alcohol has been used in partial models to improve patient compliance<sup>[36]</sup>. Because of the buildup and decrease of erythema, the body's ethanol content exhibits an expansion in bioavailability, with a faster rate of growth in bioavailability than various content.<sup>[37]</sup> It is advantageous for the local dispersion of alkaloids and also makes the stratum conium more permeable<sup>[38]</sup>

The advantages of transdermal drug delivery systems (TDDS) include simplicity of use, fewer adverse effects, enhanced bioavailability, and regulated drug administration. With less than zero-level kinetics, this distribution prevents first-pass metabolism and allows for sustained drug delivery and consumption<sup>[39]</sup>.Transformation body vectors are artificial objects that resemble the body of a body. Physical changes are largely responsible for the problems caused by transdermal drug delivery for example, the inability the cost-limiting phase in the transportation of big molecules, the penetration into the stratum corn emulsion, and the physicochemical features of the child's response via the skin.<sup>[40]</sup> These pliable vesicles have the ability to pass through skin pores, which are frequently smaller than usual, and carry heavier molecules<sup>[41]</sup>.

# Challenges Facing Medicine The development of herbal bimolecular formulations remains a challenge:-

Production of biomolecules from plants remains a challenging process. There are some difficulties in designing a suitable conveyor system as shown in the figure Further research is ongoing to find delivery systems to deliver herbal medicines from bioactive molecules to their site of action and increase their effectiveness and bioavailability. Some ways to improve the bioavailability of new herbal medicines are deriving active biomolecules to increase bioavailability. Mixing different electronic products using different. The mixture works according to the analysis. It helps to promote dynamic particle uptake, stabilize natural molecules and promote intestinal assimilation, improving pharmacokinetic properties by forming components such as soy lecithin<sup>[42-49]</sup>. Determining the adequacy or safety of most facilities is still not fully resolved. Herbs, whether alone or in combination, can have adverse effects due to differences that may cause the drug to be incompatible. Negative clinical in vitro and in vivo analyzes of herbal extracts often support treatment. But there is still a lack of welldefined and agreed-upon meetings in the exam. Research on the pharmacokinetics and bioavailability of herbal medications encounter comparable issues.. These are due to their uncertain or unknown chemical composition<sup>[50]</sup>. According to medical evidence, in the development of preclinical drug problems, the quality of medicinal herbs can influence the therapeutic effect and influence their successful integration into clinical trials.<sup>[51]</sup> The quality of botanical medicine can be very good due to natural and other factors. Different species, apparent physical characteristics, and Occasionally, natural variables such as variety might affect the amount and make-up of the biomolecules that plants generate.<sup>[52]</sup> Impurities that affect the properties of medicinal herbs, including ecological conditions, development and work in the field, health insurance, capacity, production goods, contamination, change and negative thinking<sup>[53-57]</sup>. Therefore, quality will vary from country to country, and in similar countries quality will vary from product label to product label and even within similar products. Traditional medicine is represented by the Medicines and The 1940 Cosmetics

Act and the 1945 Medicines and Cosmetics Act.<sup>[58-63]</sup> The Drugs and Cosmetics Act was modified by the Indian government in 1959 to cover medications derived from Indian medicine.<sup>[64]</sup>Treatment plans for all well conditions must adhere to the 1993 Regulation on the Safety and Efficacy of Natural Medicines, which was created by an expert panel. This applies to all medications and herbal remedies.<sup>[65]</sup> Natural elements are unregulated in various nations. Various recommendations are published by the World Health Organisation (WHO) for the clinical assessment of natural and traditional medicines.<sup>[67]</sup> The 1940 Cosmetics Act and the 1945 Medicines and Cosmetics Act. The Drugs and Cosmetics Act was modified by the Indian government in 1959 to cover medications derived from Indian medicine. Treatment plans for all well conditions must adhere to the 1993 Regulation on the Safety and Efficacy of Natural Medicines, which was created by an expert panel. This applies to all medications and herbal remedies. Natural elements are unregulated in various nations. Various recommendations are published by the World Health Organisation (WHO) for the clinical assessment of natural and traditional medicine. Treatment plans for all well conditions must adhere to the 1993 Regulation on the Safety and Efficacy of Natural Medicines, which was created by an expert panel. This applies to all medications and herbal remedies. Natural elements are unregulated in various nations. Various recommendations are published by the World Health Organisation (WHO) for the clinical assessment of natural and traditional medicines.<sup>[68-73]</sup>

# Actions needed prior to clinical evaluation of herbal drugs:-

- **A.** Collect facility details; verify identification; examine taxonomy, microscopy, and polymerase chain reaction (PCR); examine pesticides, antibiotics, and heavy metals.
- B. Create or Choose the Right Bioassays
- C. Conduct bioassays using several extract kinds.• In vitro; In vivo (if appropriate or feasible)
- **D.** The chemical characteristics and isolation of the active components are demonstrated by bioanalysis.
- E. Prepare vegetable and biological samples; carry out focused investigation
- **F.** Product in vitro investigations; Metabolism (including p450 interaction)

Mechanism of action, toxicity, and pharmacokinetics preliminary research on the production of natural botanical Able to perform Extend these. To illustrate the applicability of natural products in clinical studies, the use of a single, stable set of formulations is recommended.<sup>[74]</sup> Several techniques have to be applied in order to regulate the plant's properties Considerations like legality, correctness, and idea significance should be made while selecting science and technology.<sup>[75]</sup> Chemical analysis has evolved into an optional process with the introduction of contemporary techniques like gas chromatography (GC), high-performance liquid chromatography (HPLC), and gas chromatography-mass spectrometry (GC-MS). Regulatory agencies are currently in charge of examining how botanical preparations are administered, evaluating their efficacy, and promoting their advancement to the clinical trial stage<sup>[76-77]</sup>.

# **CONCLUSION :**

Researchers need to understand more about novel medicine delivery methods and natural ingredients. The pharmaceutical industry will use plants as a global standard for the development of novel medications. Plants have produced a wide range of synthetic materials that are isolated for use in repair or synthesis using natural crystals. Thus, the data gathered in traditional medicine plays a significant part in enhancing the availability of native medical practices. Natural goods, specialized medications, and emerging technology have always been crucial to several medical advancements and research initiatives. NHDDS facilitates the release of natural drugs by adsorbing and transporting drugs at active sites, thus limiting the serious effect of increased bioavailability of the drug. In this direction the future offered by the features of the project holds great potential, providing a great ability to analyze valuable information regarding new drug models and their new transport models determined by new drug development. Case-by-case studies should not be the only kind of study conducted on the preclinical and clinical performance of these activities.. Recently, pharmaceutical scientists have turned their attention to using methods to prepare drug delivery systems for natural medicines. The NHDDS framework will

not only boost the natural medicine market but also play an important role in providing better and safer treatment to people.

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### **REFERENCES: -**

- 1. Bruno JJ, Ellis JJ. Herbal use among US elderly: 2002 National Health Interview Survey. Ann Pharmacother 2005; 39(4): 643-8.
- Cragg GM, Newman DJ. Natural products: A continuing source of novel drug leads. Biochim Biophys Acta 2013; 1830(6): 3670-95. http://dx.doi.org/10.1016/j.bbagen.2013.02.008 PMID: 23428572.
- 3. Pan SY, Zhou SF, Gao SH, , et al. New perspectives on how to discover drugs from herbal medicines: CAM's outstanding contribution to modern therapeutics. Evi Based Compl Altern Med 2013;1-25.
- Yuan H, Ma Q, Ye L, Piao G. The traditional medicine and modern medicine from natural products. Molecules 2016; 21(5): 1-18. http://dx.doi.org/10.3390/molecules21050559 PMID: 27136524.
- 5. Ajazuddin SS, Saraf S. Applications of novel drug delivery system for herbal formulations. Fitoterapia 2010; 81(7): 680-9. http://dx.doi.org/10.1016/j.fitote.2010.05.001 PMID: 20471457.
- 6. Mandal SC, Mandal M. Current status and future prospects of new drug delivery system. Pharm Times 2010; 42: 13-6.
- 7. Musthaba SM, Baboota S, Ahmed S, Ahuja A, Ali J. Status of novel drug delivery technology for phytotherapeutics. Expert Opin Drug Deliv 2009;6(6): 625-37.
- 8. Sarangi MK, Padhi S. Novel herbal drug delivery system: An overview. Arch Med Health Sci 2018; 6(1): 171-9.
- 9. Chanchal D, Swarnlata S. Novel approaches in herbal cosmetics. J Cosmet Dermatol 2008; 7(2): 89-95.
- 10. Mukherjee PK, Harwansh RK, Bhattacharyya S. Bioavailability of herbal products: approach toward improved pharmacokinetics. Evi Based Valid Herbal Med 2015; 7: 217-45.
- 11. Wen Z, Liu B, Zheng Z, You X, Pu Y, Li Q. Preparation of Liposomes entrapping essential oil from Atractylodes macrocephala Koidz by modified RESS technique. Chem Eng Res Des 2010; 88:1102-7.
- Du Q, Cai W, Xia M, Ito Y. Purification of (+)-dihydromyricetin from leaves extract of Ampelopsis grossedentata using high-speed countercurrent chromatograph with scale-up triple columns. J Chromatogr A 2002; 973(1-2): 217-20. http://dx.doi.org/10.1016/S0021-9673(02)01092-0 PMID:12437181.
- Shariff A, Pk M, Klk P, MM. Entrapment of andrographolide in Ycross-linked alginate pellets: I. Formulation and evaluation of associated release kinetics. Pak J Pharm Sci 2007; 20(1): 1-9. PMID: 17337420.
- Rajani M, Shrivastava N, Ravishankara MN. A rapid method for isolation of andrographolide from Andrographis paniculata nees (Kalmegh). Pharm Biol 2000; 38(3): 204-9. http://dx.doi.org/10.1076/13880209(200007)3831-SFT204 PMID:21214463.
- 15. Liu M, Li H, Luo G, Liu Q, Wang Y. Pharmacokinetics and biodistribution of surface modification polymeric nanoparticles. Arch Pharm Res 2008; 31(4): 547-54. http://dx.doi.org/10.1007/s12272-001-1191-8 PMID: 18449515.

- 16. Dahnum D, Abimanyu H, Senjaya A. Isolation of Artemisinin as Antimalarial Drugs from Artemisia annua L. Cultivated in Indonesia. Int J Basic Appl Sci 2012; 12(04): 90-5.
- 17. Chang CH, Huang WY, Lai CH, et al. Development of novel nanoparticles shelled with heparin for berberine delivery to treat Helicobacter pylori. Acta Biomater 2011; 7(2): 593-603. http://dx.doi.org/10.1016/j.actbio.2010.08.028 PMID: 20813208.
- 18. Li L, Wang DK, Li LS, Jia J, Chang D, Ai L. preparation of docetaxel submicron emulsion formation for intravenous administration. J Shenyang Pharm Univ 2007; 12: 736-9.
- 19. Pradhan D, Biswasroy P, Suri KA. Isolation of berberine from Berberis vulgaris Linn. and standardization of aqueous extract by RP-HPLC. Int J Herb Med 2013; 1(2): 106-11.
- Min KH, Park K, Kim YS, et al. Hydrophobically modified glycol chitosan nanoparticlesencapsulated camptothecin enhance the drug stability and tumor targeting in cancer therapy. J Control Release2008; 127(3): 208-18. http://dx.doi.org/10.1016/j.jconrel.2008.01.013 PMID: 18336946.
- Zeng XH, Li YH, Wu SS, et al. New and highly efficient column chromatographic extraction and simple purification of camptothecin from Camptotheca acuminata and Nothapodytes pittosporoides. Phytochem Anal 2013; 24(6): 623-30. http://dx.doi.org/10.1002/pca.2441 PMID: 23722924.
- 22. Awasthi DN, Singh BP. Isolation and identification of capsaicin and allied compound in chilli. 1972.
- 23. Pandey S, Goyani M, Devmurari V, Fakir J. Transferosomes: A Novel approach for transdermal drug delivery. Pharm Lett 2009; 1:143-50.
- Ellington E, Bastida J, Francesc V, Codina C. Supercritical carbon dioxide extraction of Colchicum autumnale L. Phytochem Anal 2003; 14(3): 164-9. http://dx.doi.org/10.1002/pca.702 PMID: 12793464
- 25. Hong W, Chen DW, Zhao XL, Qiao MX, Hu HY. Preparation and study in vitro of longcirculating nanoliposomes of curcumin. Zhongguo Zhongyao Zazhi 2008; 33(8): 889-92. PMID: 18619344
- 26. Bisht S, Feldmann G, Soni S, et al. Polymeric nanoparticleencapsulated curcumin ("nanocurcumin"): A novel strategy for human cancer therapy. J Nanobiotechnology 2007; 5: 3. http://dx.doi.org/10.1186/1477-3155-5-3 PMID: 17439648
- 27. Maiti K, Mukherjee K, Gantait A, Saha BP, Mukherjee PK. Curcumin-phospholipid complex: Preparation, therapeutic evaluation and pharmacokinetic study in rats. Int J Pharm 2007; 330(1-2): 155
- 28. Patel R, Singh SK, Singh S, Sheth NR, Gendle R. Development and characterization of curcumin loaded transferosomes for transdermal delivery. J Pharm Sci 2009; 1: 71-80.
- 29. Kumar RS, Kumar M, Ganesh GNK. Formulation and evaluation of pectin hydroxyl propyl methylcellulose coated curcumin pellets for colon delivery. Asian J Pharm 2009; 3: 138-42.
- Verghese J. Isolation of curcumin from Curcuma longa L. rhizome. Flavour Fragrance J 1993; 8(6): 315-319
- 31. Pathan R, Bhandari U. Preparation and characterization of embelinphospholipid complex as effective drug delivery tool. J Incl Phenom Macrocycl Chem 2011; 9: 13947. http://dx.doi.org/10.1007/s10847-010-9824-2
- 32. Pundarikakshudu K, Joshi H, Shah P, Panchal S. A simple, facile method for isolation of embelin from fruits of Embelta ribes Burm.f (vidang). Indian drugs 2016; 53(02): 23-7.
- 33. Bhattacharya S. Phytosomes: Emerging strategy in delivery of herbal drugs and nutraceuticals. Pharm Times 2009; 41: 9-12.

- Tu L, Sun H, He S, Zhu Y, Yu M, Sun X, et al. Isolation of epigallocatechin gallate from green tea and its effects on probiotics and pathogenic bacteria. Curr Top Nutraceutical Res 2019; 17(1): 69-77.
- 35. Semalty A, Semalty M, Singh D. Supramolecular phospholipid polyphenolics interaction: The phytosome strategy to improve the bioavailability of phytochemicals. J Incl Phenom Macrocycl Chem 2010; 67: 253-60. http://dx.doi.org/10.1007/s10847-009-9705-8
- 36. Liang Q, Zhang J, Su X, Meng Q, Dou J. Extraction and separation of eight ginsenosides from flower buds of Panax Ginseng using aqueous ionic liquidbased ultrasonic-assisted extraction coupled with an aqueous biphasic system. Molecules 2019; 24(4): 778. http://dx.doi.org/10.3390/molecules24040778 PMID: 30795582
- 37. Hou J, Zhou SW. Formulation and preparation of glycyrrhizic acid solid lipid nanoparticles. ACTA Academiae Medicinae Militaris Tertiae 2008; 30: 1043-45.
- Tian M, Yan H, Row KH. Extraction of glycyrrhizic acid and glabridin from licorice. Int J Mol Sci 2008; 9(4): 571-7. http://dx.doi.org/10.3390/ijms9040571 PMID: 19325770
- Wang F, Zhou L, Gu F. Characterization of anticancer hypocrellin A encapsulated with silica nanoparticles. J Therm Anal Calorim 2010; 80: 213-8. http://dx.doi.org/10.1007/s10973-009-0630-2
- 40. Fang LZ, Qing C, Shao HJ, et al. Hypocrellin D, a cytotoxic fungal pigment from fruiting bodies of the ascomycete Shiraia bambusicola. J Antibiot (Tokyo) 2006; 59(6): 351-4. http://dx.doi.org/10.1038/ja.2006.49 PMID: 16915819
- 41. Chen C. Inhibiting the vascular smooth muscle cells proliferation by EPC and DPPC liposome encapsulated magnalol. J Chin Inst Chem Eng 2008; 39: 407-11. http://dx.doi.org/10.1016/j.jcice.2008.04.005
- 42. Chan SSK, Zhao M, Lao L, Fong HHS, Che CT. Magnolol and honokiol account for the antispasmodic effect of Magnolia officinalis in isolated guinea pig ileum. Planta Med 2008; 74(4): 381-4. http://dx.doi.org/10.1055/s-2008-1034320 PMID: 18484527
- 43. Zhaowu Z, Xiaoli W, Yangde Z, Nianfeng L. Preparation of matrine ethosome, its percutaneous permeation in vitro and antiinflammatory activity in vivo in rats. J Liposome Res 2009; 19(2):155-62. http://dx.doi.org/10.1080/08982100902722381 PMID:19241204
- 44. Cao FH, Ouyang WQ, Wang YP, Dong HB. Study of preparation of matrin nanoemulsion and its antioxidation on mice. J Nortwest A F University 2007; 3: 61-4.
- 45. Chen H, Luo S, Zheng X, Fan H. Separation of matrine and oxymatrine from Sophora flavescens extract through cation exchange resin coupled with macroporous absorption resin. Pol J Chem Technol 2016; 18(2): 10-5. http://dx.doi.org/10.1515/pjct-2016-0026
- 46. Yen FL, Wu TH, Lin LT, Cham TM, Lin CC. Naringenin-loaded nanoparticles improve the physicochemical properties and the hepatoprotective effects of naringenin in orally-administered rats with CCl(4)-induced acute liver failure. Pharm Res 2009; 26(4):893-902. http://dx.doi.org/10.1007/s11095-008-9791-0 PMID: 19034626
- 47. Yue PF, Yuan HL, Li XY, Yang M, Zhu WF. Process optimization, characterization and evaluation in vivo of oxymatrine-phospholipid complex.Int J Pharm 2010; 387(1-2): 139-46. http://dx.doi.org/10.1016/j.ijpharm.2009.12.008 PMID: 20005937
- 48. Rane S, Prabhakar B. Formulation and evaluation of pH-sensitive, long circulating liposomes for paclitaxel delivery. Int J Pharm Tech Res 2009; 1: 914-7.
- 49. Trickler WJ, Nagvekar AA, Dash AK. A novel nanoparticle formulation for sustained paclitaxel delivery. AAPS PharmSciTech 2008; 9(2): 486-93. http://dx.doi.org/10.1208/s12249-008-9063-7 PMID: 18431660
- 50. Dong X, Mattingly CA, Tseng MT, et al. Doxorubicin and paclitaxel-loaded lipid-based nanoparticles overcome multidrug resistance by inhibiting Pglycoprotein and depleting ATP.

Cancer Res 2009; 69(9): 3918-26. http://dx.doi.org/10.1158/0008-5472.CAN-08-2747 PMID:19383919

- 51. Ketchum REB, Luong JV, Gibson DM. Efficient extraction of paclitaxel and related taxoids from leaf tissue of taxus using a potable solvent system. J Liq Chromatogr Relat Technol 1999;22(11): 1715-32. http://dx.doi.org/10.1081/JLC-100101762
- 52. Emiko EF, Nonaka GI, Nishioka I, Hayashi K. Isolation and structures of procyanidins (condensed tannins) from Rhaphiolepis umbellata. Agric Biol Chem 1986; 50(8): 2061-7. http://dx.doi.org/10.1080/00021369.1986.10867679
- 53. Rong G, Juqun X. Studies on molecular interaction between puerarin and PC liposomes. Chin Sci Bull 2007; 52: 2612-7. http://dx.doi.org/10.1007/s11434-007-0395-6
- 54. Li P, Lu Y, Du S, et al. Extraction and purification of flavonoids from Radix puerariae. Trop J Pharm Res 2013; 12(6): 919-27. http://dx.doi.org/10.4314/tjpr.v12i6.9
- 55. Priprem A, Watanatorn J, Sutthiparinyanont S, Phachonpai W, Muchimapura S. Anxiety and cognitive effects of quercetin liposomes in rats.Nanomedicine 2008; 4(1): 70-8. http://dx.doi.org/10.1016/j.nano.2007.12.001 PMID: 18249157
- 56. Wu TH, Yen FL, Lin LT, Tsai TR, Lin CC, Cham TM. Preparation, physicochemical characterization, and antioxidant effects of quercetin nanoparticles. Int J Pharm 2008; 346(1-2): 160-8. http://dx.doi.org/10.1016/j.ijpharm.2007.06.036 PMID: 17689897
- Vicentini FT, Simi TR, Del Ciampo JO, et al. Quercetin in w/o microemulsion: In vitro and in vivo skin penetration and efficacy against UVB-induced skin damages evaluated in vivo. Eur J Pharm Biopharm 2008; 69(3): 948-57. http://dx.doi.org/10.1016/j.ejpb.2008.01.012 PMID: 18304790
- 58. Horbowicz M. Method of quercetin extraction from dry scales of onion. Veget Crops Res Bull 2002; 57: 119-24.
- 59. Xiao L, Zhang YH, Xu JC, Jin XH. Preparation of floating rutinalginate-chitosan microcapsule. Chin Tradit Herbal Drugs 2008; 2:209-12.
- 60. Al-Mahdawe MM, Al-Mallah MK, Ahmad TA. Isolation and identification of rutin from tissues cultures of Ruta graveolens L. J Pharm Sci Res 2018; 10(6): 1517-20.
- 61. Garg R, Gupta GD. Gastro retentive floating microspheres of Silymarin: Preparation and in vitro evaluation. Trop J Pharm Res 2010; 9: 59-66. http://dx.doi.org/10.4314/tjpr.v9i1.52037
- 62. Wianowska D, Wisniewski M. Simplified procedure of silymarin extraction from Silybum marianum L. Gaertner. J Chromatogr Sci 2015; 53(2): 366-72.PMID: 24895445
- 63. Mazumder A, Dwivedi A, du Preez JL, du Plessis J. In vitro wound healing and
- 64. cytotoxic effects of sinigrin-phytosome complex. Int J Pharmaceu 2016; 498(1-2): 283-93. http://dx.doi.org/10.1016/j.ijpharm.2015.12.027 PMID: 26706438
- Al Shahawany AW, Al Hatta ZN, Al Tahhan SF. Qualitative and Quantitative Analysis of Sinigrin in Different Parts in vitro and in vivo of Brassica nigra Plants. Biomed Biotechnol 2016; 4(1): 19- 24.
- 66. Xiaoyan A, Jun Y, Min W, et al. Preparation of chitosan-gelatin scaffold containing tetrandrineloaded nano-aggregates and its controlled release behavior. Int J Pharm 2008; 350(1-2): 257-64.
- 67. Xie Z, Xu X, Xie C, Liang Z. Preparative isolation of tetrandrine and fangchinoline from Radix Stephania tetrandra using reversedphase flash chromatography. J Liq Chromatogr Relat Technol 2013; 37(3): 343-52. http://dx.doi.org/10.1080/10826076.2012.745139
- 68. Chen JG, Liu YF, Gao TW. Preparation and anti-inflammatory activity of triptolide ethosomes in an erythema model. J Liposome Res 2010; 20(4): 297-303.
- 69. Shen J, He C. Isolation and purification of triptolide from the leaves of Tripterygium wilfordii Hook F. Chin J Chem Eng 2010; 18(5): 750-4. http://dx.doi.org/10.1016/S1004-9541(09)60124-5

- 70. Li S, Ji Z, Zou M, Nie X, Shi Y, Cheng G. Preparation, characterization, pharmacokinetics and tissue distribution of solid lipid nanoparticles loaded with tetrandrine. AAPS PharmSciTech 2011;12(3): 1011-8. http://dx.doi.org/10.1208/s12249-011-9665-3 PMID: 21811889
- 71. Lira MCB, Ferraz MS, da Silva DGVC, et al. Inclusion complex of usnic acid with βcyclodextrin: Characterization and nanoencapsulation into liposomes. J Incl Phenom Macrocycl Chem 2009; 64: 215-24. http://dx.doi.org/10.1007/s10847-009-9554-5
- 72. Stark JB, Walter ED, Owens HS. Method of isolation of usnic acid from Ramalina reticulate. J Am Chem Soc 1950; 72(4): 1819-20. http://dx.doi.org/10.1021/ja01160a118
- Lu Y, Hou SX, Chen T, Sun YY, Yang BX, Yuan ZY. [Preparation of transfersomes of vincristine sulfate and study on its preutaneous penetration]. Zhongguo Zhongyao Zazhi 2005; 30(12): 900-3.PMID: 16124605
- 74. Hadagali A, Hegde P, Manasa KH, Madihalli C, Pradeep S, Shettihalli AK. Isolation and detection of vinca alkaloids from endophytes isolated from Catharanthus roseus. Euro J Biomedi Pharm Sci 2017; 4(10): 675-83.
- 75. Ke X, Xu Y, Yan F, Ping QN. The liposomes of wogonin & rats in vivo pharmacokinetics. Zhongguo Yaoke Daxue Xuebao 2007; 38:502-6.
- Delange DM, Rico CLM, Canavaciolo VG, Cuellar AC, Oliver ES. Selective and high yield isolation of pure Wogonin from aerial parts of Scutellaria havanensis Jacq. Int J Pharm Sci Rev Res 2015; 30(2): 104-8.
- 77. Mohanraj VJ, Chen Y. Nanoparticles: A review. Trop J Pharm Res 2006; 5(1): 561-73.