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Novel Drug Delivery Systems of Herbal Medicine

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ABSTRACT:

Over the past several years, great advances have been made on development of novel drug delivery systems (NDDS) for plant actives and extracts. The various type of novel herbal formulations such as polymeric nanoparticles, nanocapsules, liposomes, phytosomes, animations. microsphere, transfersomes, ethosomes has been reported using proactive and plant selections. The novel formulations are reported to have remarkable 0advantages over conventional formulations of plant actives and extracts which include ha cement of solubility, bioavailability, protection from toxicity, enhancement pharmacological activity, enhancement of stability, improved tissue macrophages distribution, sustained delivery, and protection from physical and chemical degradation.Phytosome is a patented technology developed by a leading maker of drugs and nutraceuticals, to incorporate standardized plant extracts or water-soluble phytoconstituents in to phospholipids to produce lipid - compatible molecular complexes. The herbal drugs can be used in a more upright course with enhanced efficacy by incorporating them into moderndosage forms. The present review highlights the current status of the development of novel herbal formulationsand summarizes their method of preparation, type of active ingredients, size, entrapmentefficiency, route of administration, biological activity and applications of novel formulations.

KEYWORDS: Liposomes, Nanoparticles, Phytosome, Ethosomes, Emulsion, Marketed Herbal Novel Drug Delivery Formulations'

I. INTRODUCTION:

In the past few decades, considerable attention has been Concentrated on the evolution of a novel drug delivery. The novel carriers should ideally Fulfill two prerequisites. Firstly, it should deliver the

drug at a Rate directed by the needs of the body, over the period to Treatment. Secondly, it should channel the active entity of Herbal drug to the site of action. Conventional dosage forms Including prolonged release dosage forms are unable to meet None of these. In phyto formulation research, developing Nano System (NDDS) for herbal drugs. Dosage forms (polymeric nanoparticles and nanocapsules, Liposomes, solid lipid nanoparticles, phytosomes and Nano-Emulsion etc.) have a number of advantages for herbal drugs, Including enhancement of solubility and bioavailability, pro-M Tection from toxicity, enhancement of pharmacological activity, enhancement of stability, improving macrophages Distribution, sustained delivery, protection from physical and Chemical degradation etc. Thus the nano sized novel drug Delivery systems of herbal drugs have a potential future for Enhancing the activity and overcoming problems associated With plant medicines. Liposomes, which are biodegradable And essentially non-toxic vehicles, can encapsulate both hydrophilic and hydrophobic materials [1]. Liposome based drug Delivery systems offer the potential to enhance the therapeutic Index of anti-cancer agents, either by increasing the drug con-Centration in tumor cells and/or by decreasing the exposure In normal tissues exploiting enhanced permeability and retention effect phenomenon and by utilizing targeting strategies[2]. The main advantages using liposomes include: i) the biocompatibility, ii) the easiness of preparation, iii) the Chemical versatility that allows the loading of hydrophilic, Amphiphilic, and lipophilic compounds, iv) the simple Modulation of pharmacokinetic properties by changing the chemical composition of the bilayer components [3]. Delivery of agents to the reticuloendothelial system (RES) is Easily achieved, since most conventional liposomes are trapped By the RES [1]. The application of novel approaches can also Improve the efficacy of herbal



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cosmetic formulations on the Human body [4]. Similarly the other vesicular systems like Nanoemulsion, ethosomes and transferosomes are highly Useful assemblies and find various advantages in the delivery Of herbal medicines; some of them are summarized in present Article. The phytosome process has also been applied to many Popular herbal extracts including Ginkgo biloba, grape seed, Hawthorn, milk thistle [5], green tea, and ginseng. The Flavonoid and terpenoid components of these herbal extracts Lend themselves quite well for the direct binding to Phosphatidylcholine. Phytosome is produced by binding Individual components of herbal extracts to phosphatidyl Choline, resulting in a dosage form that is better absorbed and Thus, produces better results than the conventional herbal Extracts [6]. The results indicate that the absorption of silybin From silybin phytosome is approximately seven times greater Compared to the absorption of silybin from regular milk Thistle extract [5]. Drugs can be embedded or dissolved in Nanoparticles and can also be adsorbed or coupled on the Surface [7]. Encapsulating drugs within NPs can improve the Solubility and pharmacokinetics of drugs, and, in some cases, Enable further clinical development of new chemical entities That have stalled because of poor pharmacokinetic properties [8]. The major carrier materials of nanoparticles are synthetic Biodegradable high molecular polymer and natural polymer. The former usually includes poly-αcyanoacrylate alkyl Esters, polyvinyl alcohol, polylactic acid, and polylacticco-Glycolic acid, etc. The latter is usually divided into two classes:Proteins (albumin, gelatin and vegetable protein) and Polysaccharides (cellulose, starch and its derivatives, alginate, Chitin and chitosan, etc.) [9]. In this article, an attempt has been made to touch upon Different aspects related to the development of novel herbal formulations, including method of preparation, type of active Ingredient, entrapment efficiency, and applications etc

of Novel Herbal Drug **Types** Systems: Several approaches in case of new herbal drug delivery system include different types of expressions such as mouth-dissolving tablets, liposomes, phytosomes, pharmacosomes, museums, microspheres, nanoparticles, transfersomes, transdermal drug delivery system ethosomes, (TDDS), and proniosomes are discussed Mouth-dissolving tablets: Asoka Lifescience Limited launched Res-Q, the world's first Polyherbal mouth- dissolving

mouth-dissolving drug. It induces a new drug delivery system that imparts Efficacy.IntheAyurvedicmedicine segment, this is the inaugural Attempt to make medicines more effective in managing chronic Ailments. Res - Q is a polyherbal medicine highly effective for Lung problems and other respiratory ailments such as asthma. This unique mouth-dissolving drug delivery system ensures That the drug reaches the blood right away and the first- pass Metabolism is bypassed. It dissolves in mouth by mixing with The saliva and get absorbed. This Res-Q provides relief from Respiratory distress within 15 min. Hence the product shows a Great resemblance with the efficacy of Sorbitrate, a revolutionary Mouth-dissolving drug used in cardiac distress.[10]

LIPOSOMES: The liposomes are spherical particles that encapsulate a Fraction of the solvent, in which they freely diffuse (float) into Their interior. They can have one, several or multiple Concentric membranes. Liposomes are constructed of polar Lipids which are characterized by having a lipophilic and Hydrophilic group on the same molecules [11]. Upon Interaction with water, polar lipids selfassemble and form Self-organized colloidal particles. Simple examples are deter-Gents, components form micelles, while polar lipids with Bulkier hydrophobic parts cannot associate into micelles with High curvature radii but form bilayers which can self-close Into liposomes or lipid vesicles. A cross-section of a liposome(Fig. 1) depicts the hydrophilic heads of the amphiphile Orienting towards the compartment while the lipophilic tails orient away from the water towards the center of The vesicle, thus forming a bilayer. Consequently, water Soluble compounds are entrapped in the water compartment And lipid soluble compounds aggregate in the lipid section. Uniquely, liposomes can encapsulate both hydrophilic and Lipophilic materials. Liposomes usually formed from phospholipid's, have been used to change the pharmacokinetics Profile of, not only drugs, but herbs, vitamins and enzymes. A Variety of herbal liposomal formulations has been studied Which are summarized in Table 1. Because of their unique Properties liposomes are able to enhance the performance of Products by increasing ingredient solubility, improving Ingredient bioavailability, intracellular uptake and enhanced pharmacokinetics and biodistribution [12] and in vitro And in vivo stability. Liposomes as a drug delivery system can Improve the therapeutic activity and safety of drugs, mainly By delivering them to



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their site of action and by maintaining Therapeutic drug levels for prolonged periods of time [13–14]. Milk thistle (Silybummarianum) is one of the few herbal Drugs whose excellent pharmacological profile readily lends Itself to proof of clinical efficacy [15]. Meanwhile, silymarin is Poorly absorbed (20–

50%) from the gastrointestinal tract [16]That causes the effects of silybin, one of the main active Flavonoids commonly found in the dried fruits of silymarin, to Be greater after parenteral than oral administration [17].

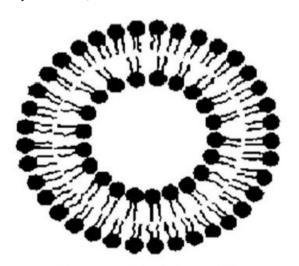


Fig. 1. Cross-section of a liposome [4].

Plants/constituents	Therapeutic category	Applications with respect to liposomal technology	Reference
Ampelopsin	Anticancer	Improved therapeutic efficacy	18
Capsaicin	Analgesic	Prolong action, permeation enhancement	19
Curcumin	Anticancer	Long systemic residence time and high entrapment efficiency	20
Paclitaxel	Anticancer	pH sensitivity and improved entrapment efficiency	21
Usnic acid	Antimycobacterial	Prolong action and solubility enhancement	22
Wogonin	Anticancer	Prolong duration of action	23
Quercetin and rutin	Hemoglobin	Enhancement of Hemoglobin binding	24
Garlicin	Lungs	Increase efficiency	25
Catechins	Antioxidant and chemopreventive	Increased permeation through skin	26
Breviscapin	Cardiovascular diseases	Sustained delivery of breviscapine	27

Table 1: Herbal formulations in liposomal drug delivery systems.

Nanoparticles: In recent year, the nanonization of herbal medicines has Attracted much attention; [28] some of them are illustrated in Table 2. Nanoparticles and Nano emulsions (Fig. 2) are Colloidal systems with particles varying in size from 10 nmTo 1000 nm [29,30]. Nanoparticle systems with mean particle Size well above the 100 nm standard have also been

reported In literature, including nanonized curcuminoids [31], pacli-Taxel [32] and praziquantel [33] which have a mean particle Size of 450, 147.7, and even higher than 200 nm, respectively. IN addition, nanoparticles could also be defined as being Submicronic (b1 lm) colloidal systems [34]. The nanospheres Have a matrix type

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structure in which the active ingredient is Dispersed throughout (the particles), whereas the Nano capSules have a polymeric membrane and an active ingredient Core. Nanonization possesses many

advantages, such as Increasing compound solubility, reducing medicinal doses, And improving the absorbency of herbal medicines compared With the respective crude drugs preparations [35].

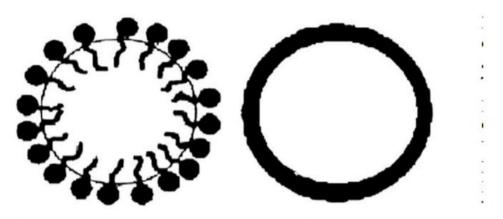


Fig. 2. Cross-section of (a) nanoemulsion and (b) biopolymeric nanoparticle [4].

Plants/constituents	Therapeutic category	Applications Wrt Nanotechnology	Reference
Triptolide nanoparticle	Anti- inflammatory	Enhance the penetration of drugs through the stratum corneum	36
		by increased hydration	
Artemisinin nanocapsules	Anticancer	Sustained drug release	37
Texel nanoparticle	Anticancer	Enhance the bioavailability and sustained drug release	38
Berberine nanoparticle	Anticancer	Sustained drug release	39
Curcuminoids solid lipid	Anticancer	Prolonged release of the curcuminoids	40
Nanoparticle	A	D 1 111 1 1 1 1 1	4.1
Camptothecin encapsulated	Anticancer	Prolonged blood circulation and high accumulation in tumors	41
Nanoparticle			
Naringenin nanoparticle	Hepatoprotective	Improved the release of NAR and improved its solubility	42
Breviscapine nanoparticle	Cardiovascular and	Prolong the half- life and decrease RES uptake	43
	cerebrovascular		
Tetrandrine nanoparticle	Lung	Sustained drug release	44
Glycyrrhizic acid nanoparticle	Anti- inflammatory, antihypertensive	Improve the bioavailability	45

Table 3: Herbal Nanoparticulate Drug Delivery System.

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Phytosome: Over the past century, phytochemical and phytipharmacological sciences established the compositions, biological Activities and health promoting benefits of numerous plant Products. Most of the biologically active constituents of plants Are polar or water soluble molecules. However, water soluble Phytoconstituents (like flavonoids, tannins, terpenoids, etc.)

Are poorly absorbed either due to their large molecular size Which cannot absorb by passive diffusion, or due to their poor Lipid solubility; severely limiting their ability to pass across The lipidrich biological membranes, resulting poor bioavailability [46]. It has often been observed that the isolation and Purification of the constituents of an extract may lead to a Partial or total loss of specific bio-activity for the purified Constituent the natural constituent synergy becomes lost. Very often the chemical complexity of the crude or partially Purified extract seems to be essential for the bioavailability of The active constituents. Extracts when taken orally some

Constituents may be destroyed in the gastric environment. As Standardized extracts are established, poor bioavailability Often limits their clinical utility due to above said reasons. It Has been observed that complexation with certain other Clinically useful nutrients substantially improves the bioavailability of such extracts and their individual

constituents. The nutrients so helpful for enhancing the absorption are the Phospholipids. Phytosome is a patented technology developed by a leading manufacturer of drugs and nutraceuticals, To incorporate standardized plant extracts or water soluble Phytoconstituents into phospholipids to produce lipid compatible molecular complexes, called as phytosomes and so Vastly improve their absorption and bioavailability [47] (Table 3). In liposomes no chemical bond is formed; the Phosphatidylcholine molecules surround the water soluble Substance. There may be hundreds or even thousands of Phosphatidylcholine surrounding the water soluble compound. In contrast, with the phytosome process the Phosphatidylcholine and the plant components actually form A 1:1 or a 2:1 molecular complex depending on the substance (s) complexed, involving chemical bonds (Fig. 3). Phospholipids are complex molecules that are used in all known life Forms to make cell membranes. In humans and other higher Animals the phospholipids are also employed as natural Digestive aids and as carriers for both fat-miscible and water Miscible nutrients. They are miscible both in water and in Lipid environments, and are well absorbed orally. Phytosomes Are more bioavailable as compared to conventional herbal Extracts owing to their enhanced capacity to cross the lipoidal Bio membrane and finally reaching the systemic circulation.

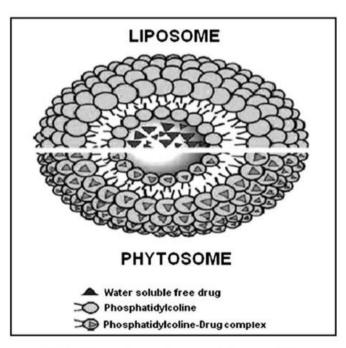


Fig. 3. Difference between liposome and phytosome [58].



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Plants/constituents	Therapeutic category	Applications WrtPhytosomal technology	Reference
Ginkgo biloba	Cardioprotective, antioxidant activity	Flavonoids of GBP stabilize the ROS	48
Ginsenosides	Nutraceutical, immunomodulator	Increase absorption	49
Curcumin	Antioxidant, anticancer	Increase antioxidant activity and increase bioavailability	50
Quercetin	Antioxidant, anticancer	Exerted better therapeutic efficacy	51
Epigallocatechin	Nutraceutical, systemic antioxidant, anticancer	Increase absorption	52
Naringenin	Antioxidant activity	Prolong duration of action	53
Silybin	Hepatoprotective, antioxidant for liver and skin	Absorption of silybin phytosome from silybin is approximately 7 times greater	54

Ethosomes: Newer advancements in the patch technology have led to The development of ethosomal patch, which consists of Drug in ethosomes. Ethosomal systems are made up of soya Phosphatidylcholine, ethanol and water. They may form Multilamellar vesicles and have a high entrapment capacity For particles of various lipophilicities. The elastic vesicles transfersomes have also been used as drug carriers for a Range of small molecules, peptides, proteins vaccines.[55] Ethosome has a deformability and entrapment efficiency And can penetrate through the skin completely and improve Drug delivery through the skin. Likened to other liposomes, the Physical and chemical properties of ethosomes make the legal Transfer of the drug through the stratum corneum into a deeper Skin layer efficiently or even into the blood circulation.[56] This property is very important as the topical drug carrier and Transdermal delivery system. Moreover, the ethosomes carrier Also can provide an efficient intracellular delivery for both Hydrophilic and lipophilic drugs,[57] percutaneous absorption Of matrine an anti-inflammatory herbal drug is increased,[58] It also permits the antibacterial Peptide to penetrate into the Fibrocyte easily.[59] From the review of literature it has been observed that, Only three clinical trials have been conducted on ethosomal Systems in human volunteers. Horwitz et al. carried out a Pilot, double-blind, randomized clinical study to compare The efficacy of an ethosomal acyclovir preparation and Commercially available acyclovir cream (Zovirax®) in treating Recurrent herpes labialis in 40 human volunteers. The results Revealed that the ethosomal acyclovir preparation performed Betterthan Zovirax cream and showed significant improvement In all the evaluated clinical parameters, such as the time of crust Formation and disappearance and pain parameters. The efficacy Of ethosomal gel of clindamycin phosphate and salicylic acid Was evaluated in a pilot clinical trial of 40 acne patients treated With the gel twice daily for 8 weeks. Volunteers treated with Ethosomal gel showed considerable improvement in acne Condition, with a decreased number of comedies, pustules, And total number of lesions compared to placebo. Ethosomal Preparation of prostaglandin E1 Was evaluated in a pilot clinical Study in patients with erectile dysfunction. It was observed that 12 of 15 tested patients had improved peak systolic velocity And penile rigidity. Erection duration was 10-60 min. There Was no reported adverse skin reactions associated with the Treatment in any of the aforementioned clinical trials. [60] Table 7 showing the Clinical data of ethosomes.[61]



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Plants/constituents	Applications category	Therapeutic Wrt	Reference
		Nanotechnology	
Capsaicin transfersomes	Analgesic	Increase skin penetration	62
Colchicine transfersomes	Antigout	Increase skin penetration	63
Vincristine transfersomes	Anticancer	Increase entrapment	64
		efficiency and skin	
Matrine ethosomes	Anti-inflammatory	Improve the	65
		percutaneous permeation	
Ammonium	Anti-inflammatory	Increase of the in vitro	66
glycyrrhizinate		percutaneous permeation	
ethosomes			

EMULSION: Emulsion refers to a homogeneous dispersion system That is composed of two kinds of liquids unable to dissolve each Other, and one of which disperse in the other one in a form of Droplets [67]. Generally, emulsion is composed of oil phase, Water phase, surfactant and sub-surfactant. Its appearance is Translucent to transparent liquid. Emulsion can be classified Into ordinary emulsion (0.1–100 µm), micro-emulsion (10- 100 nm), sub-micro-emulsion (100-600 nm), etc. (Table 4). Among them, the micro-emulsion is also called Nano emulsions, And the sub-microemulsion is also called lipid emulsion. As a Drug delivery system, emulsion distributes in vivo in the Targeted manner due to its affinity to the lymph. In addition, the Drug can be sustained release in a long time because the drug is Packaged in the inner phase and kept off direct touch with the Body and tissue fluid [68]. After the oily drugs or lipophilic Drugs being made into O/W or O/W/O emulsion, the oil droplets Are phagocytosised by the macrophage and get a high Concentration in the liver, spleen, and kidney in which the Amount of the dissolved drug is

very large. While water soluble Drug is produced into W/O or W/O/W emulsion, it can be easily Concentrated in the lymphatic system by intramuscular or Subcutaneous injection. The size of the emulsion particle has an Impact on its target distribution. Apart from its targeted sustained release, producing the Herbal drug into emulsion will also strengthen the stability of The hydrolyzed materials, improve the penetrability of drugs to The skin and mucous, and reduce the drugs' stimulus to tissues. So far, some kinds of herbal drugs, such as camptothecin, Brucea Javanica oil, coixenolide oil and zedoary oil have been made Into emulsion. For example, Zhou et al. [69] studied the Influence of the elemenum emulsion on the human lung Adenocarcinoma cell line A549 and protein expression. Results Showed that the elemenum emulsion has a significant Inhibition on the growth and proliferation of the A549 in Vitro and it showed a time and dose-dependent relationship. Elemenum emulsion is a type of new anti-cancer drug with Great application prospects. Furthermore, it has no marrow Inhibition and no harm to the heart and liver.

Plants/constituents	Applications category	Therapeutic Wrt	Reference
		Nanotechnology	
Self-nanoemulsifying	Hepatoprotection	Improved aqueous	70
Zedoary	anticancer		
essential oil	Anti-inflammatory	dispersibility, stability and oral	71
	-	dispersibility, bioavailability	
Triptolide microemulsion	Anti-inflammatory	Enhance the penetration of drugs	72
_	-	through the stratum corneum by	
		increased hydration	
Docetaxel submicron	Anticancer	Improve residence time	73
Emulsion			
Berberine nanoemulsion	Anticancer	Improve residence time and	74
		absorption	
Silybin nanoemulsion	Hepatoprotective	Sustained release formulation	75
Quercetin microemulsion	Antioxidant	Enhance penetration into stratum	76
		corneum and epidermis	

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Marketed Herbal Novel Drug **Delivery Formulations**: Two companies dominate the market for these systems, Namely, Cosmetochem and Indena. For herbal drug delivery, Cosmetochem launches Herbasec® technology in markets Which are actually liposomal preparations of various herbal Ingredients such as extracts of White tea, Green tea, white Hibiscus, Gurana, and Aloe Vera. These extracts are used in Cosmetics because of their anti-oxidant effects for prevention Of aging. Indena patented the technology of phytosomes® And launches many products in market under this having Diverse therapeutic benefits. Indena commercializes plant Constituents/extracts of liquorice (18ß-glycyrrhetinic acid), Ammi visnaga (visnadin), Centella asiatica (triterpenes), G. biloba (ginkgoflavonglucosides, ginkgolides, bilobalide), Hawthorn flower (vitexin-2"-O-rhamnoside), milk thistle (silymarin and Silybin), horse chestnut (escin β-sitosterol), Terminalia sericea (sericoside), Panax ginseng (ginsenosides), Grape seed (polyphenols), Green tea (polyphenols), etc.[77]

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II. Conclusion:

Herbal medications have been widely employed all over The globe since ancient times and have been acknowledged By doctors and patients for their better therapeutic value as They cause fewer adverse effects as compared with modern Medications. The drugs of Ayurvedic origin can be utilized in A more upright course with enhanced efficacy by incorporating In modern dosage forms. However, phytotherapeutics need a Scientific approach to render the components in a new way to Increase patient compliance and avoid repeated administration. This can be accomplished by designing NDDS for herbal Ingredients. NDDS not only reduce the repeated administration To overcome noncompliance, but also help to increase the Therapeutic value by reducing toxicity and increasing the Bioavailability and so on. Recently, pharmaceutical scientists.

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