Review Article

Novel Drug Delivery System Used in Cosmetics: A Short Review

Siddhi Khanke, Shweta Kale

Department of Quality Assurance, Dada Saheb Balpande College of Pharmacy, Besa, Nagpur MS, India - 440037

Received: 21th Oct 21; Revised 25th Nov, 21, Accepted: 15th Dec, 21; Available Online: 25th Dec., 21

ABSTRACT

Delivery systems area unit chemical agents that carry the active compounds to the location of their action. they're utilized in cosmetics chiefly for his or her ability to enhance the soundness of sensitive actives, their higher incorporation into formulations, reduction of irritation potential, glorious penetration and sustained unleash properties. The foremost unremarkably used delivery systems embrace liposomes, noisome, microemulsions and Nano emulsions, small and nanoparticles, chemical compound micelles and cyclodextrin complexes. Their properties and effects area unit totally different and every system is appropriate for various kinds of compounds. Delivery systems area unit wide used for the incorporation of anti-aging compounds, like lightening agents, plant extracts, antioxidants and vitamins, similarly as for ultraviolet radiation filters and fragrances. Flavourer cosmetics area unit outlined because the product that ready by or enclosed plants and/or flavourer elements that area unit combination of the many natural molecules or compounds

INTRODUCTION

Novel Drug delivery System refers to the approaches, formulations, technologies, and systems for transporting a pharmaceutical compound within the body as required to securely win its desired therapeutic effects. Drug delivery systems (DDSs) square measure developed to deliver the specified variety of medicine effectively to acceptable target sites and to take care of the specified drug levels. analysis in newer drug delivery system is being disbursed in liposomes, nanoparticles, noisome, percutaneous drug delivery, implants, microencapsulation, and polymers¹. Drug delivery systems, nanoparticles as carriers have shown nice potential in recent years. The encapsulation of medicine in nanoparticles, together with micelles, liposomes, dendrimers, nano capsules, nanospheres et al., improves the therapeutic index and reduces the adverse facet effects Delivery, systems square measure chemical agents carrying active compounds to the location of their action. They typically contain anti-aging compounds (vitamins, change of colour agents, antioxidants), ultraviolet illumination filters or fragrances. They'll even be applied in hair care cosmetics as carriers of nutritive agents, dyes, conditioners, humectants, deodorants, and as antistatic agents²³. the foremost usually used delivery systems embody sac delivery systems (liposomes and noisome), emulsion delivery systems (microemulsions and nano emulsions), particulate systems (microparticles, nanoparticles, compound micelles and solid supermolecule nanoparticles)¹. The cosmetics are originated from plants in its historical development.

ancient use of plants for cosmetic functions supported perfuming and skin care within the kind of infusions, poultices and etc. within the last century, researchers are cantered on plants to research their effectivity and safety in cosmetics field.²³ Generally flavourer sources square measure wealthy with vitamins, antioxidants, oils (essential etc.) hydrocolloids, proteins, terpenoids and different bioactive compounds that have functions within the scope of cosmetics like anti-aging, anti-oxidant, emollient result etc². Herbs may be utilised for cosmetics in numerous forms as- a region of herb, total extract of the herb, extract of selective elements, specific molecules refined from extracts⁴⁶. within the scope of flavourer cosmetics Phyto cosmetics square measure outlined because the product that ready solely by plants and/or flavourer elements and chiefly included; plants, plant extracts, volatile oils, distillates, aromatic waters, juices, binary compound extracts, tinctures, resins, gums and being, flavoured oils lipids, waxes, mucilage's, plant carbohydrates or refined plant elements. vital activities just in case of Phyto cosmetics square measure generally; inhibitor activity, hymenopter on tyrosinase activity and antimicrobial activity⁴⁶. As the Phyto formulation could be a mixture of quite one active ingredient, care ought to be taken to the determination of the soundness profile for Phyto cosmetics/herbal cosmetics¹⁻³. **Ouantitative** standards of all the flavourer elements have to be compelled to determine in keeping with a globally acceptable reference like The Ayurvedic assemblage of Republic of India, Chinese formulary and etc. vital

ISSN: 0975-4873

parameters that have an effect on the ultimate quality and stability of flavourer cosmetics/Phyto cosmetics square measure the specifications of flavourer inputs, structure of formulations Associate in Nursing producing method Liposomes square measure sac delivery systems with a centre consisting of an binary compound cavity that is encircled by one or additional hydrophobic bilayer membranes composed of phospholipids. The diameter of those vesicles will vary in vary from 25 to 5000 nm². Microspheres square measure spherical microparticles that square measure usually between one and one,000 microns (1 mm) in diameter. By comparison, a person's hair is about 75 microns in diameter. Microspheres square measure usually brought up as spheres, balls, beads, small beads or micro balloons and square measure factorymade from a range of raw materials. The foremost common sorts square measure solid and hollow glass microspheres, solid and hollow chemical compound microspheres, and ceramic microspheres. Microemulsions represent a promising carrier system for cosmetic active ingredients thanks to their various blessings over the present typical formulations²⁹. Noisome square measure utilized in the sector of cosmetics since the first 1970's. they're stable with smart penetrating power and fewer irritating as compared to different, mixture carriers. Noisome, additionally known as non-ionic wetter vesicles square measure microscopic lamellar structures that square measure fashioned by the admixture of non-ionic wetter and cholesteric¹. Polymeric micelles, with typical diameters starting from ten to one hundred nm, square measure nanoscopic core-shell structures fashioned by amphiphilic block copolymers. The inner core consists of hydrophobic regions of amphiphiles, wherever the lipotropic medicine square measure being solubilized¹. Nano emulsions square measure clear, kinetically stable identical mixtures of oil, water, wetter and co-surfactant, with a driblet diameter of but one hundred nm. They're characterised by smart sensory and biophysical properties²⁷.

HISTORY

- The 1st skin patch was approved in 1981 to forestall the nausea and physiological reaction related to ill.
- The bureau has approved, till 2003, over thirty-five skin patch products, spanning thirteen molecules IN USA).
- The U.S.A. transdermic market approached \$1.2 billion in a pair of 001 it had been supported eleven drug molecules: anodyne, nitro-glycerine, oestradiol, ethinylestradiol, norethindroneacetate, androgen, clonidine, nicotine, lidocaine, prilocaine, and hyoscine.λ

- Two new, recently approved skin patch product (a contraceptive patch containing ethinylestradiol and nor elgestromin, and a patch to treat hyperactive bladder, containing oxybutynin.
- 1965 1st description of closed supermolecule bilayer vesicles.
- 1967 introduction of the term liposomes to explain closed supermolecule bilayer vesicles.
- 1972 liposomes 1st used as delivery systems of drug.
- 1974 1st patients to be injected with liposomes.
- 1979 liposomes 1st used as delivery systems of nucleic acids to cells 1980 1st being opposing body targeted liposomes termed immunoliposome

AIM

- The aim of Novel Drug Delivery System is to supply a therapeutic quantity of drug to the acceptable web site within the body to accomplish promptly then maintain the specified drug concentration.
- The drug- delivery system ought to deliver drug at a rate management by the essentially of the body over a mere term of treatment.
- Novel drug delivery system in cosmetics business aims for durable effects.
- To increase stability.
- To minimize drug degradation.
- To increase bioavailability

Objective

- To cut back dosing frequency.
- To delay delivery to the colon to attain high native concentrations within the treatment of diseases of the distal gut.
- To delay delivery to a time acceptable to treat acute phases of sickness (chronotherapy).
- To deliver to a vicinity that's metabolically less hostile, e.g., to facilitate absorption of acid and enzymatically labile materials, particularly amide

Scope

- In dermatology
- In dermal applications liposomes are used as protective systems for active ingredients
- In modern cosmetic science
- In herbal formulations for cosmetics
- Liposomes deliver nutrients directly to aging cells and would improve skin hydration and texture, reduce fine lines and diminish wrinkles

Liposomes

Liposome's area unit spherical vesicles during which their central binary compound section is close to by one or additional of a bilayer membrane (Lamella) that's continuously fenced by aquatic environments. These vesicles area unit shaped once amphiphilic lipids oppose with binary compound environment. They will vary in size from fifteen nm to several microns. Liposomes is Greek words means that 'lipo' mean 'Fat' and 'Some's' mean 'Body'. Liposomes were initial factory-made in European country in 1961 by Alec D. Bingham within the last thirty years, the employment of vesicle has been expanded from drug delivery to the cosmetic field and it's the foremost wide well-known cosmetic delivery system these days. (4) thanks to their special structure, liposomes are often used as a delivery system, carrying hydrophilic agents through their fenced binary compound section, and oleophilic substances the non-ionic tails of the bilayer section. Liposome's area unit sac delivery systems with a centre consisting of associate degree binary compound cavity that is close one or additional hydrophobic bilayer membranes composed of phospholipids. The diameter of those vesicles will vary in vary from 25 to 5000 nm.³¹ The most reason why liposomes area unit widespread in cosmetics is their ability to encapsulate each hydrophilic and hydrophobic molecules, the development of active ingredient absorption by the skin, and also the simple technique of their preparation. additionally, they will

additionally offer controlled unleash profiles for several substance Liposomes also are ready to improve the steadiness of sure ingredients, as shown by Lee C.H. et al. with astaxanthin, a strong inhibitor utilized in cosmetics as anti-aging agent. They encapsulated this sensitive compound into liposomes that improved its light-weight and thermal stability and augmented its doable application in cosmetics Liposomal product in cosmetics aren't restricted to skin care and for hair care, in 1989 a liposomal formulation was made. However, not several alternative liposomal products used for hair made to the market since then. The primary product containing liposomes for make-up product was a powder launched in 1988 followed by war paint and completely different foundations9. The first liposomal cosmetic product introduced into the industrial market was the Capture anti-aging cream by firm Christian Dior in 1986, that has been followed by several alternative products

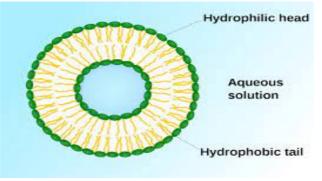
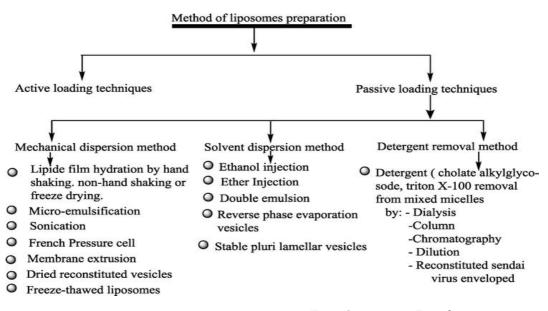


Figure 1: Structure of liposome





Types of Cosmetic Liposomes

• Transferosomes: Transferosomes square measure extremely deformable, reactive, and economical

liposomes applied thus far for direct transcutaneous drug delivery¹⁴.

- **Photosomes:** Photosomes act by cathartic photolysis enzymes extracted from the marine plant Anacystinidulans they're extensively utilized in sunscreens¹⁴.
- Ethosomes: These sorts of liposomes square measure soft and versatile multilayer vesicles composed of lipoid phosphatidylcholine, water, and 2 hundredth five hundredth ethyl alcohol. Ethosomes square measure non-invasive carriers that change the element penetrate deeply into the skin layers or enter circulation¹⁴.
- Ultrasomes: Ultrasomes square measure a singular class of liposomes that square measure shaped by denial of the nuclease extracted from Micrococcus luteus. they assist observe actinic radiation damage to the skin and increase the speed of treatment by up to four times¹⁴.

Application of Liposomes

Liposomes will play the role as each vehicle of cosmeceutical materials and as active agents themselves. once skin is suffering from skin condition or broken because of lack of wet, empty liposomes will extremely move with skin lipids, proteins, and carbohydrates serving to during this approach the skin to come to traditional state and creating the horny layer perform its defensive perform properly¹⁰

- Facilitate the Penetration¹³
- Overcome solubility limitations¹³
- Increase Stability
- Overcome solubility limitations⁹
- Cause Longer Effect⁹
- Target Selective¹⁵
- Make the merchandise Economic¹⁰
- Separating element from External Milieu¹⁴

Advantages

- Biocompatibility
- capacity for self-assembly
- ability to hold massive drug payloads

Disadvantages

- Less stability.
- Batch to batch variation.
- Difficult in massive scale producing and sterilization Noisome

Niosomes are artificial microscopic vesicles consisting of associate degree binary compound core closed in an exceedingly metallic element layer consisting of sterol and one or additional non-surfactants Noisome are employed in the sector of cosmetics since the first 1970's. they're stable with smart penetrating power and fewer

irritating as compared to alternative, mixture carrier. Noisome, additionally known as non-ionic wetter vesicles are microscopic lamellar structures that are shaped by the admixture of non-ionic wetter and sterol⁶. They're shaped by the self-assembly of amphiphilic molecules into closed bilayers. Since they need associate degree amphiphilic bilayer structure, they will entrap each hydrophilic further as hydrophobic medication. Acceptable mixtures of surfactants and charge causation agents provide thermodynamically stable vesicles. Alternative factors tributary to the formation of noisome embrace HLB worth of the amphiphilic molecule, binary compound bed, lipide chain-length, chain-packing and membrane. Noisome are used for the dermatologic purpose in cosmetic trade. The cosmetic whole that initial developed and proprietary niosomes was L'Oréal. Later the merchandise 'Noisome Plus's associate degree antiageing cream was developed. Niosomes offers many blessings in cosmetic and skin care product thanks to their ability to extend the steadiness of entrapped medication with improved bioavailability of poorly absorbed ingredients so enhancing skin penetration²¹. But skin acts a serious barrier for topical formulations, horny layer being the biggest barrier. Hence, there's associate degree urge to possess a correct carrier to deliver the medication through the skin which may be consummated victimization novel delivery systems. The foremost wide used delivery systems are liposomes and that they are getting used in an exceedingly form of skin care rejuvenating product. Liposomes are capable of encapsulating numerous anti-ageing active ingredients and deliver them deep into the cells. The primary liposomal anti-ageing cream to enter the market was "Capture" launched by Christian Dior in 1986¹⁵.

Types of Noisome

The different types of niosomes are proniosomes, as pasomes, deformable niosomes, vesicles in water and oil systems (v/w/o)

- Pro-niosomes Pro-niosomeare dry granular product which get converted to liposomal suspension after subsequent hydration. They are more stable as compared to niosomes⁶
- Aspasomes These vesicles are formed from a mixture of ascorbic palmitate, cholesterol and highly charged lipid. In order to form niosomes, aspasomes are first hydrated and then sonicated. Aspasomes enhance the transdermal permeation of drugs⁶
- ² Deformable niosomes these vesicles are also called as elastic niosomes as they are flexible in nature. They are composed of surfactants, ethanol and water.⁶

• Vesicles in water and oil system (v/w/o) An aqueous suspension of niosomes is emulsified into the oily

phase at 60°C to form vesicles in water in oil emulsion (v/w/o). 6

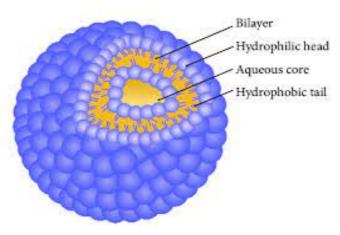
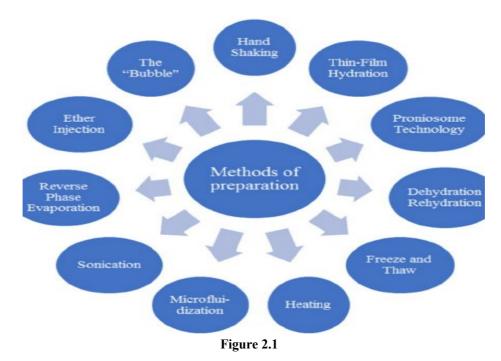


Figure 2 : Structure of Noisome

Application of Niosome

- Anti-neoplastic treatment
- Transdermal Drug Delivery Systems
- Noisome enhance the uptake of drugs through the skin.
- Cosmetics: Topical use of noisome entrapped antibiotics to treat acne is done.

METHOD OF PREPARATION OF NOISOME



Advantages

• Enhance the skin porosity of medication once applied locally

• they're osmotically active and stable

• the surfactants used and additionally the ready noisome area unit perishable, biocompatible and non-immunogenic

Disadvantages

- Time consuming
- Requires specialized equipment
- Charge inducing molecule
- inefficient drug loading

MICROEMULSIONS

Microemulsions square measure clear, thermodynamically stable and frequently low viscous dispersions of water, oil and frequently wetting agent and co-surfactant. The dimensions of particles range between ten and one hundred ums. Therefore, microemulsions seem as isotropous, optically clear liquids or gels. They're a lot of stable than common emulsions, putting them among enticing cosmetic delivery systems. On the opposite hand, a better concentration of surfactants and co-surfactants (20-25 percent) is required for his or her preparation, which might cause accrued irritation of the skin¹⁷. Microemulsions represent a promising carrier system for cosmetic active ingredients because of their various blessings over the prevailing typical formulations. They're capable of solubilizing each deliquescent and

lipotropic ingredients with comparatively higher encapsulation. Microemulsions could more be used for making ready DE make-up removers; face cleansing merchandise, as an example, face cleansing foam for shiny, sensitive and/or inflammatory disease skin; degreasing lotion for shiny hair or dandruffs; lotions for moistened handkerchiefs; shampoo formulas, among alternative cosmetic formulations. The operate of this cosmetic microemulsion is to scrub the hair by causative on the removal of shampoo residues maintained thereon and, on the scalp, getting used as a pre-shampoo, that is, before applying the standard shampoo, while not inflicting the scales of the hair to open, which might harm the cuticle^{18,19}.

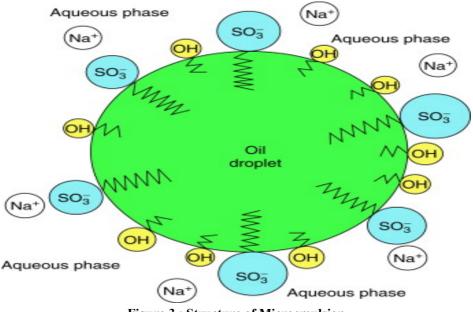


Figure 3 : Structure of Microemulsion

Types of Microemulsions

- O/W Microemulsion
- W/O Microemulsion
- Bi continuous Microemulsion

Method of Preparation of Microemulsions

- Phase titration method
- Phase inversion method Omega College of pharmacy.

Application of Microemulsions

- To delivery of deliquescent similarly as lipotropic drug as drug carriers as a result of
- Improved drug solubilization capability
- Long period
- Easy of preparation
- Improvement of bio-availability Omega faculty of pharmacy

Advantages

- Increase the speed of absorption
- Increase bio-availability

- Helpful in style masking
- Eliminates variability in absorption

Disadvantages

- Use of enormous concentration of wetter and cosurfactant necessary for the stabilising small droplets.
- Limited solubilizing capability for prime melting substances.
- Microemulsion stability is influenced by environmental parameters like, temperature & pH

Nano Emulsion

Nano emulsions area unit clear, kinetically stable isotropic mixtures of oil, water, wetter and co-surfactant, with a driblet diameter of but one hundred nm. they're characterised by smart sensory and biophysical properties Nano emulsions as vehicles to be utilized in dermo cosmetics and cosmetics for skin and hair applications³⁶. Nano emulsions area unit pseudo ternary systems, deeprooted by ingredients of ancient cosmetics (water, oils, and surfactants) however, in contrast to emulsions, these formulations area unit extremely stable, have low driblet size, and permit a straightforward flow over the skin with no creaming and shiny coating The potentialities of nano emulsions in terms of incorporation of bioactive molecules, final stipulation form, and therefore the pleasant sensory properties were self-addressed²⁷.

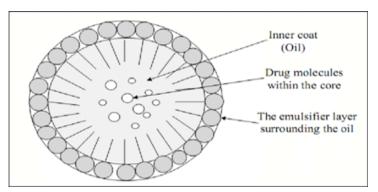
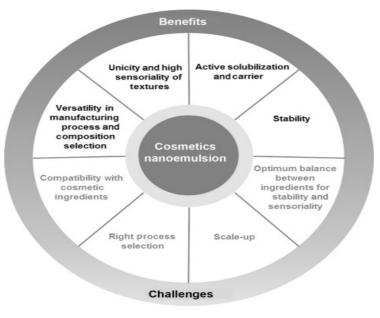


Figure 4: Structure of Nanoemulsion

Benefits of Nano emulsion





Application of Nano Emulsion

• small drop size (in the vary 20-200 nm) with high surface space,

- transparent or semi-transparent look,
- high solubilization capability

• low viscousness • high kinetic stability thanks to alluviation,

• flocculation, and in some cases, the coalescency.

• Nano emulsion area unit a lot of appropriate delivery system for the transport of oleophilic compounds as they support the skin penetration of active ingredients and therefore increase their concentration within the skin that plays a vital role in cosmetics product formulations^{27,34}.

Advantages

• Nano emulsion has small-sized droplets having bigger extent providing bigger absorption.

• It will be developed in form of formulations like foams, creams, liquids, and sprays.

• It provides higher uptake of oil-soluble supplements in cell culture technology.

- It helps to solubilize oleophilic drug.
- Protection of drug.
- Enhance drug solubility

Disadvantages

• Use of an outsized concentration of chemical agent and co-surfactant necessary for helpful the nanodroplets.

• Limited solubilizing capability for high-melting substances.

• The chemical agent should be nontoxic for exploitation pharmaceutical applications.

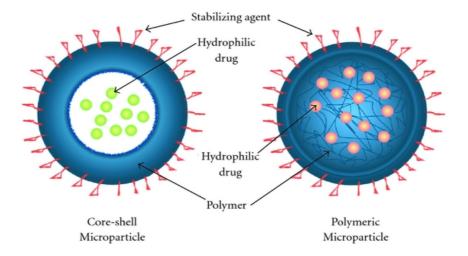
- Stability
- Solubility

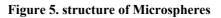
• Expensive **MICROPARTICLES**

Microparticles square measure solid chemical compound particles go in size from one to a hundred um. they're shaped from a core and membrane consisting of organic polymers, fats, proteins, polysaccharides, etc. they will be employed in cosmetics for the protection of sensitive compounds from the setting, as proven by Scalia and Mizzens with pic unstable agents and for compatibility improvement and unfavourable-odour reduction This result preserves the protecting capability of the actinic ray filters by holding them on the skin surface, and conjointly limits potential nephrotoxic reactions. (24) furthermore, encapsulation implies that a lower concentration of actinic ray filters is needed within the formulation. Microparticles of atomic number 11 alginate and atomic number 11 alginate with starch square measure employed in the developed formula as abrasive agents that exfoliate dead skin cells. The obtained microparticles have regular, spherical form, that minimizes the danger of skin irritation throughout application of the peel¹⁹.

Types of Microspheres

- Bio adhesive microspheres
- Floating microspheres
- Radioactive microspheres
- Magnetic microspheres
- Polymeric microspheres i) Biodegradable polymeric microspheres ii) Synthetic polymeric microspheres





Method of Preparation

- Single emulsion technique
- Double emulsion technique
- Solvent evaporation
- Phase separation coacervation technique
- Spray drying and spray congealing
- Solvent extraction
- Polymerization

Application of Microparticles

- Scrubs and Exfoliating Agent
- Enhances the tactile expertise of a cosmetic product
- To free pores and exfoliate dead cell.

Advantages

• Particle size reduction for enhancing solubility of the poorly soluble drug

- Provide constant and prolonged therapeutic impact.
- Provide constant drug concentration in blood there by increasing patent compliance.
- Decrease dose and toxicity.

• Protect the drug from catalyst and photolytic cleavage therefore found to be best for drug

Disadvantages

• The prices of the materials and process of the controlled unleash preparation, area unit well more than those of ordinary formulations.

• The fate of compound matrix and its impact on the atmosphere.

• The fate of compound additives like plasticizers, stabilizers, antioxidants and fillers.

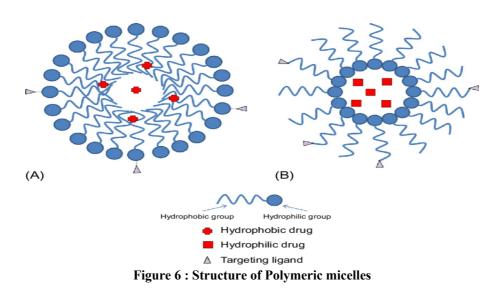
• Reproducibility is a smaller amount.

• Process conditions like modification in temperature, pH, solvent addition, and evaporation/agitation might influence the soundness of core particles to be encapsulated.

• The environmental impact of the degradation product of the compound matrix created in response to heat, hydrolysis, oxidation, radiation or biological

Polymeric micelles

Polymeric micelles, with typical diameters starting from ten to a hundred nm, are nanoscopic core-shell structures shaped by amphiphilic block copolymers. The inner core consists of hydrophobic regions of amphiphiles, wherever the lipotropic medication is being solubilized. (30) The core region is enclosed by a palisade or corona composed of deliquescent blocks of amphiphiles. chemical compound micelles are typically composed of polyesters or poly amino acids covalently secured to a biocompatible deliquescent block, usually PEG (polyethylene glycol) The chemical compound micelles of oleanolic acid ready during this study were stable and effective at assuaging wrinkles in humans because the principal active ingredient. supported these findings, it's expected that chemical compound micelles of oleanolic acid may be wide employed in cosmetic applications. (31)



Method of Preparation

- Direct method
- Dialysis method
- Indirect method using organic solvent
- Solution casting method ^(30,31)

Application

- oleanolic acid in *cosmetic* products as the main ingredient is limited by its poor aqueous solubility
- in dermatological products of cosmetics

Advantages

- Specific ability to encapsulate hydrophilic drugs.
- These carriers can enhance the therapeutic efficacy and minimize the systemic side effects of the drugs.

Disadvantages

- High cost of preparation and Drug loading
- Deformation

Solid Lipid Nanoparticles

In associate degree liquid medium and square measure stabilised by zero. 5-5 pp. wetting agent. Secondgeneration nanoparticles square measure referred to as nanostructured macromolecule carriers (NLC). They were developed to beat potential limitations related to SLN, that square measure lower loading capability for active compounds, higher water content of the particle suspension or inflated potential to expulsion of active compounds throughout storage⁵². In cosmetics, NLC give controlled unleash profiles for several substances. Because of their macromolecule composition, they exhibit low toxicity and toxicity, that interprets into wonderful tolerability. Their tiny size ensures shut contact with the stratum and may increase the number of the drug penetrated into the skin. Moreover, they're ready to improve the chemical stability of compounds sensitive to light-weight, chemical reaction and chemical reaction47.

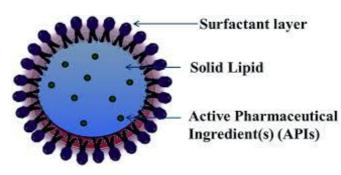


Figure 7: Structu0re of Solid Lipid Nanoparticles

METHOD OF PREPARATION

The lipoid supermolecule nanoparticles were ready from a liquid nanophase containing water and a water mixable organic solvent wherever each lipoid and desoxyribonucleic acid area unit severally dissolved by removing the organic solvent, stable and homogeneously sized lipid-nucleic acid nanoparticle (70-100 nm) were shaped. (34)

Application

• Solid lipid nanoparticles for skin and drug delivery

Advantages

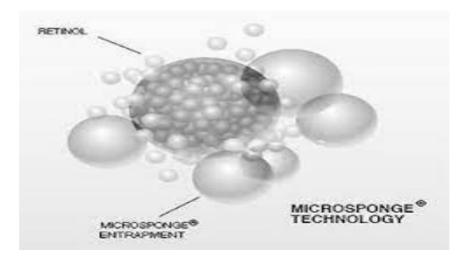
- Low toxicity
- high bioavailability of drug
- versatility of incorporation of hydrophilic and oleophilic medication
- feasibility of large-scale production

Disadvantages

- Low drug-loading capacities,
- The chance of super cooled melts that cause stability.
- Presence of different mixture structure

Micro sponges

The small sponge Delivery System (MDS) may be a proprietary compound system consisting of porous microspheres. Square measure they're} little sponge like spherical particles that carries with it a myriad of interconnecting voids within a no collapsible structure with an oversized porous surface through that active ingredient are discharged in an exceedingly controlled manner. Small sponges square measure compound delivery systems composed of porous microspheres³⁸. They're little sponge-like spherical particles with an oversized porous surface. Moreover, they will enhance stability, scale back facet effects and modify drug unleash favourably. Small sponge technology has several favourable characteristics, that build it a flexible drug delivery vehicle. Microsponge Systems square measure supported microscopic, polymer-based microspheres which will suspend or entrap a good form of substances, and may then be incorporated into a developed product like a gel, cream, liquid or powder³.



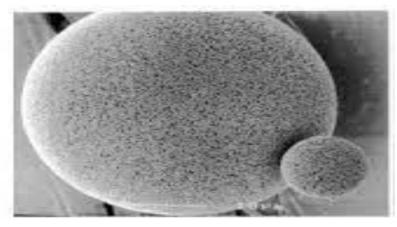


Figure 8. Structure of Micro sponges³⁹

Method of Preparation of small sponge

- Liquid-liquid suspension chemical change
- Quasi-emulsion solvent diffusion³⁷

Applications of small sponges

- Micro sponge for topical deliver
- For retention of indefinite quantity kind on skin

Advantages

- Enhanced product performance.
- Extended unharness.
- Reduced irritation and thence improved patient Compliance.
- Improved product elegancy
- Improved oil management because it will absorb oil up to six times its weight while not drying
- Improved formulation flexibility.
- **Disadvantages**
- Greasiness,
- Requires high quantity of drug formulations
- Stickiness

Advantages

- Hydrophilic and hydrophobic drug are often delivered.
- Liposome botanical medicine acts as a carrier for little cytotoxic molecules and as vehicle for macromolecules as cistron.
- Sustained and controlled unleash of formulation are often potential Increase effectiveness of the drug.
- Site specific delivery.
- Decreased toxicity/side effects.
- Increased convenience.
- Viable treatments for antecedental incurable diseases.
- Potential for prophylactic applications.
- Better patient compliance
- Enhancement of solubility.
- magnified bioavailability.
- Protection from toxicity.
- Improvement of pharmacologic activity
- Improvement of stability.
- Improved tissue macrophages distribution.

- Sustained delivery
- Protection from physical and chemical degradation **Disadvantages**

Disadvantages

- Poor patient compliance
- Increased probabilities of missing the dose of a drug with short half-life that frequent administration is critical
- The ineluctable fluctuation of drug concentration might result in underneath medication or over medication
- A typical peak depression plasma conc time profile is obtained that create attainment of steady state condition troublesome
- The fluctuations in drug levels might result in precipitation of adverse effects particularly of a drug with tiny therapeutic index whenever over medication occur
- Daily dose of over 10mg isn't doable.
- Local irritation may be a major downside.
- Drug requiring high blood levels are unsuitable.
- Drug with long half- life can't be developed in TDDS
- Uncomfortable to wear. Might not be economical.
- Barrier operate changes from person to person and inside a similar person
- Heat, cold, sweating (perspiring) and showering stop the patch from projecting to the surface of the skin for over in some unspecified time in the future.

• A new patch has got to be applied daily.

Future Prospect and Application

According to a recent estimate, the event of a replacement drug for human use in volves around \$800 million and 10-12years of analysis inputs. However, nine out of ten medications fail in their clinical study section inflicting Brobdingnagian loss to the investigation organization. For NDDS development, America estimate shows the event value be around \$40 million and amount between three months to three years The burden on company's pecuniary resource is kind of less with smart probabilities of ensured returns sometimes, because of little development cycle, multiple pipe-line NDDS product may well be contemplated guaranteeing an even bigger market-share in an exceedingly specific section. The variety of novel seasoning formulations like chemical compound nanoparticles, nano capsules, liposomes, phytosomes, nano emulsions, microsphere, transferosomes, and ethosomes has been reportable victimization bioactive and plant extracts. The novel formulations square measure reportable to own outstanding blessings over typical formulations of plant actives and extracts that embrace sweetening of solubility, bioavailability, protection from toxicity, sweetening of medicine activity, sweetening of stability, improved tissue macrophages distribution, sustained delivery, and protection from physical and chemical degradation. The present review highlights the present standing of the event of novel seasoning formulations and summarizes their technique of preparation, form of active defence ingredients, size, potency, route of administration, biological activity and applications of novel formulation. The usage of seasoning formulations for NDDS is additional beneficious and advantageous as against others. The usage of cyst, emulsion, phytosomes, microsphere, and powerful lipoid nanoparticles of seasoning formulation have improved the remedial impacts of plant extricates. With the usage of these, directed delivery of the formulation is accomplished, attributable to that the formulation exhibits impact on the location, and therefore the bioavailability of the formulation is likewise enlarged. With these novel medication conveyance frameworks, the actives and concentrates that square measure utilized as a part of natural formulations exhibit sustained unleash of formulation, sweetening in stability, improved therapeutic effectiveness, and protection from toxicity

CONCLUSION

The use of assorted varieties of delivery systems in cosmetics is changing into ever a lot of in style, totally on account of growing efforts to realize the best potential impact of the active compounds. Developments during this space square measure creating fast headway Efforts square measure specializing in the event of entirely new systems and on innovating or combining some existing systems. Thus, combining the resultant impact of the loaded substances is a lot of exaggerated. One among the best edges of the delivery systems is that they're able to penetrate the corneum within the case of carrying a vigorous compound that has to reach deeper layers of the skin. At identical time, they need the capability to retain different substances that don't seem to be meant to penetrate the skin's surface. Herbal cosmetics that square measure designed with novel drug delivery systems have bestowed some blessings like providing high effectiveness, increased stability, cut back undesirable effects and higher aesthetic look of merchandise. This article explains the constituents of liposomes; however, they need been discovered and entered the cosmetic field, also as their definition. Later, it introduces differing kinds of cosmetic liposomes that may be utilised in numerous cosmetic formulations counting on their specific properties and at last, the advantages of application of liposomes in cosmetics square measure taken into

thought. By mistreatment liposomes, we tend to square measure able to overcome some restrictions like low penetration, solubility, stability, length of impact and high aspect effects or prices, and improve another characteristic. The small sponge delivery technology of controlled unharness system during which active pharmaceutical ingredient is loaded within the macro porous beads and initiates reduction in aspect effects with improved therapeutic effectiveness. Micro sponge is effectively incorporated into topical drug delivery system for retention of indefinite quantity kind on skin, and conjointly use for oral delivery of medicine mistreatment bio erodible polymers, particularly for colon specific delivery and controlled unharness drug delivery system so rising patient compliance by providing web site specific drug delivery system and prolonging indefinite quantity intervals. This technology is being employed presently in cosmetics, over-the-counter skin care, sunscreens, and prescription merchandise. This sort of drug delivery technology could result in a more robust understanding of the healing of many diseases. Hence, the small sponge-based drug delivery technology is probably going to become a valuable drug delivery matrix substance for numerous therapeutic applications within the future

REFRENCES

- 1. Delivery System Handbook for Personal Care and Cosmetic Products: Technology, Applications, and Formulations, Edited by Meyer R.R., Ed. William Andrew Publishing, Norwich, N.Y., USA. 2005.
- Winterhalter M, Hilty C, Bezrukov S M, Nardin C, Meier W, Fournier D. Controlling membrane permeability with bacterial porins. Applications to encapsulated enzymes. Talanta Arora N., Agarwal S., et al., IJPSDR, 2012; 4(3): 168-182.
- Lee VHL. Controlled Drug Delivery Fundamentals and Applications: Influence of drug properties on design. 2nd ed. Marcel Dekker, Inc. New York: 1987; 16-25 Patra vale V.B. and Mandawgade S.D., Int J Cosmetic Sci, 2008; 30(1): 19-33.
- Dragicevic N, Maibach HI. Percutaneous Penetration Enhancers Physical Methods in Penetration Enhancement. Springer; Sharma B. and Sharma A., Int J Pharm Pharma Sci, 2012; 4(3): 57-66.
- Makino K, Isayama R, Hisamitsu I, Hayamizu K, Tsuji T, Shibata M. Preparation of Liposome Film Pleasant Texture and Highwater Loss Suppression for Lip Care. Soc Colour Mater. 2013;86(7):243–6 Li D., Wu Z., et al, J Cosmetic Sci, 2011; 62(6), 549-563.
- Liu J.J., Nasal S., et al., J Cosmetic Sci, Moat A. de C., de Freitas Z.M., et al. Int J Nanomedicine 2013 ; 64(1) : 9-17 (8, 4689- 4701).
- 6. Pillaiyar T, Manickam M, Namasivayam V. Skin whitening agents: medicinal chemistry perspective of

tyrosinase inhibitors. J Enzyme Inhibit Med Chem. 2017

- Nacht, S. and Katz, M. The microsponge: a novel topical programmable delivery system. In: Topical Drug Delivery Formulations (Osborne, D.W. and Amann, A.H., eds). Marcel Dekker Inc., New York (1990). 322-323
- Park S.N., Lee M.H., et al., Beachem Biopsy's Res Common, Hayward, J.A. Potential of liposomes in cosmetic science. Cosmetic. Toil. 105, 47–54 (1990). 435(3), 361-366 (2013).
- Bale, S.; Khurana, A.; Reddy, A.S.S.; Singh, M.; Godugu, C. Overview on Therapeutic Applications of Microparticle Drug Delivery Overview on Therapeutic Applications of Microparticulate Drug Delivery Systems. Crit. Rev. Ther. Drug Carry. Syst. 2016, 4, 309–361.
- Verma P. and Pathak K., JAPTR, Tavano L., Muzzalupo R., et al., Colloids Surf, Paul B.K. and Moulik S.P., Curr Sci, B 1(3), 274-282 (2010). 114, 144-149 (2014).
- 12.Wang, B.H.; Longquin Hu, T.J.S. Drug Delivery to the Lymphatic System. In Drug Delivery Principles and Applications; Wang, B., Longquin Hu, T.J.S., Eds.; John Wiley and Sons Inc.: Hoboken, NJ, USA, 2016
- Acter, S.; Cho, J.; Kim, J.W.; Byun, A.; Park, K.H.; Kim, J.W. Synthesis and shape control of uniform polymer microparticles by tailored adsorption of poly (ethylene oxide)-b-poly(ε-caprolactone) copolymer. Bull. Korean Chem. Soc. 2015, 36, 1467–1473
- Carlotti M.E., Gallarate M., et al., J Cosmetic Sci, Tsai Y.H., Lee K.F., et al., Int J Pharm 54(5), 451-462 (2003). 388(1-2), 257-262 (2010).
- Redziniak, G. and Perrier, P. Cosmetic applications of liposomes. In: Microencapsulation: Methods and Industrial Applications (Benita, S., ed.), pp. 577–579. Marcel Dekker Inc., New York (1996)
- Spiclin P., Homar M., et al., Int J Pharm Mahdi E.S., Noor A.M., et al., Int J Nanomedicine,256 (1-2), 65-73 (2003). 6, 2499-2512 (2011)
- Schmidt, J.B., Binder, M., Macheines, U.V. and Bieglmager, C. New treatment of atrophic acne scars by iontophoresis with estriol and tretinoin. Int. J. Dermatol. 34, 57–62.
- Hu Z., Liao M., et al., Int J Nanomedicine, Jain S.K. and Jain N.K., Int J Cosmetic Sci 7, 5719-5724 (2012). 32(2), 89-98 (2010).
- Scalia S. and Mezzena M., AAPS PharmSciTech, 10(2), 384-390 (2009). 20. Jain S.K. and Jain N.K., Int J Cosmetic Sci, 32(2), 89-98 (2010)

- Suzuki, K. and Sakon, K. The applications of liposome to cosmetics. Cosmetic. ToilCiternesi, U. and Sciacchitano, M. Phospholipid/ active ingredient complexes. Cosmetic. Toil. 110, 57–68 (1995)
- 20. Scalia S., Mezzena M., et al., Skin Pharmacol Physiol, Lu Y. and Park K., Int J Pharm, 24453 (1), 198-214 (2013).
- Sonneville-Aubrun, Simonnet, J.T. and Alloret, F.L. Nanoemulsions: a new vehicle for skin care products. Adv. Colloid Interface Sci. 108–109, 145–149 (2004)
- Fox, C. Cosmetic and pharmaceutical vehicles: skin care, hair care, makeup and sunscreens. Cosmetic. Toil. 113, 45–56 (1998).
- Pardeike J., Hommoss A., et al., Int J Pharm, Arnett, J.M. and Scher, R.K. Nail cosmetics. Int. J. Dermatol. Yamazaki, H. Novel O/W type nail enamel. Fragrance J. 20, 86–88 (1992 366(1-2), 170-184 (2009). 31, 675 (1992).
- 24. Preat, V. and Vanbever, R. Topical delivery by iontophoresis. In: Handbook of Cosmetic Science and Technology (Barel, A.O., Paye, M. and Maibach, H.I., eds), pp. 211–219. Marcel Dekker Inc., New York Nikolić S., Keck C.M., et al., Int J Pharm, 414(1-2), 276-284 (2011).
- 25. Adriana RP, Leticia C, Graziela M, Leonardo US, Nadya PD and Silvia SG. Structural model of polymeric nanospheres containing indomethacin ethyl ester and in vivo antiedematogenic activity. Int J Nanotechnol, 4(5), 454–67. (2007)
- 26. Aiyar HN, Seshadri R, Raina G, Sen R and Rahul R. Study of Carbon Nano capsules (Onions) and Spherulitic Graphite by Stim and Other Techniques. In: Fullerene Sci Tech, 3 (6), 765-7. (1995)
- 27. Andrieu V, Fessi H, Dubrasquet M, Devissaguet JP, Puisieux F and Benita S. 1989. Pharmacokinetic evaluation of indomethacin nano capsules. Drug Des Delivery 4(4), 295- 302. (1989).
- Benita. S. 1998. Microparticulate drug delivery systems: release kinetic models. Microspheres, Microcapsules and Liposomes (the MML Series). R. Arshady (Ed.), Citrus Books, London, pp. Kaur N. Puri R., et al., AAPS Pharma SciTech, 11(2), 255-278 - 528-537 (2010).
- Gers-Barlag, H. and Muller, A. Emulsifier-free finely disperse systems of the oil-in-water and water-in-oil type. US Patent 6 703 032. Beiersdorf AG, Hamburg Numanoglu U., Sen T., et al., AAPS Pharma SciTech, 8(4), E1-E9 (2004), (2007)
- Laouini A, Jaafar-Maalej C, Limayem-Blouza I, far S, Charcosset C, Fessi H. Preparation, characterization and applications of liposomes: state

of the art. J Colloid Sci Biotechnol. 2012;1(2):147-68

- Kurapati S. The current role of nanomaterials in cosmetics. J Chem Pharm Res. Arora N., Agarwal S., et al., IJPSDR. Patravale V.B. and Mandawgade S.D., Int J Cosmetic Sci, 30(1), 19-33 (2008). 2016;8(5):906–144(3), 168-182 (2012).
- Reva T, Vaseem AA, Satyaprakash S, Md. Khalid JA. Liposomes: The novel approach in cosmaceuticals. World J Pharmacy Sci. 2015;4(6):1616–40.
- 33. Singh A, Malviya R, Sharma PK. Nova some-a breakthrough in pharmaceutical technology a review article. Adv Boil Res. 201 1;5: 184–9.
- 34. Karimi N, Ghanbarzadeh B, Hamishehkar H, KEYVANI F, Pezeshki A, Gholian MM. Phytosome and liposome: the beneficial encapsulation systems in drug delivery and food application. J Apple Food Biotechnol. 2015;2(3):17–26.
- Tapas KP, Oli M. Prospect of nanotechnology in cosmetics: benefit and risk assessment. World J Pharm Res. 2014;3(2):1909–13.
- 36. 1. Shivani Nanda, Mandeep Kaur, Nikhil Sood, Sahil Nagpal, Microsponge drug delivery system: an overview, World Journal of Pharmacy and Pharmaceutical Sciences, Volume 2, Issue 3, 1032-1043
- 37. 3. Chad war, V. and J. Shaji, Microsponge delivery system. Curr Drug Deliv, 2007. 4(2): p. 9-123. 4.
 N.H. Anorak, A.S. Kulkarni, D.J. Ingle and R.A. Patil, Micro sponges as Innovative Drug Delivery Systems, International Journal of pharmaceutical Sciences and Nona technology, Volume 5, Issue 1, April – June 2012.
- Anderson D.L., Cheng C.H., Nacht S (1994). Flow Characteristics of Loosely Compacted Microporous Microsponge(R) Polymeric Systems. Powder Technol78: 15-18.
- Barkai A., Pathak V., Benita S (1990). Polyacrylate (Eudragit retard) microspheres for oral controlled release of nifedipine. I. Formulation design and process optimization. Drug Dev Ind Pharm 16: 2057-2075.
- 40. Namrata Jadhav, Vruti Patel, Siddhesh Mungekar, Manisha Karpe, Vilasrao Kadam, Microsponge

delivery system: an updated review, current status and future prospects, World Journal of Pharmacy and Pharmaceutical Sciences, Volume 2, Issue 6, 6463-6485.

- 41. Embil K. Nacht S. The Micro sponge Delivery System (MDS): A topical delivery system with reduced irritancy incorporating multiple triggering mechanisms for the release of actives. J. Microencapsul.1996; 3(5), 575-588.
- 42. Nacht S, Kuntz M. The micro sponge: A novel topical programmable delivery system. Top Drug Deliv Syst. 1992; 42:299-325.
- 43. Won R: Method for delivering an active ingredient by controlled time release utilizing a novel delivery Pyrogen vehicle which can be prepared by a process utilizing the active ingredients as a. 1987; US Patent No. 4690825
- Burlando B, Verotta L, Cornara L, Bottini-Massa E. Herbal Principles in Cosmetics: properties and mechanisms of action. New York: CRC Press; 2010. 56-97.
- 45. Plants in Cosmetics. Council of Europe Publishing, Strassbourg Cedex; 2001.
- Alğın Yapar E, İnal, Erdal S. Design and in vivo evaluation of emulgel formulations including green tea extract and rose oil. Acta Pharmaceutica 2013;63(4):531-44.
- Masand S, Madan S, Balian S. K. Modern concept of storage and packaging of raw herbs used in Ayurveda. Int J Res Ayurveda Pharm. 2014; 5(2):242-5
- 48. D'Amelio FS. Botanicals: a Phyto cosmetic desk reference. USA: CRC Press; 1999.
- 49. Seidell JC, osterlee A, Deurenberg P, et al. Abdominal fat depots measured with computed tomography: Effects of degree of obesity, sex, and age. EurJClinNutr 1988; 42:805-815.
- 50. Arul jothy M et al. An Overview of noisome as carrier in dermal drug delivery. Journal of pharmaceutical sciences & research 2015; 7 :923-927.
- 51. Ghanashyam S et al. Review of current &novel trends for anti-ageing formulations. International journal of pharmaceutical, chemical and biological sciences 2014; 4 :118-125.