

Liquid Crystals Pharmaceutical Application: A Review

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Abstract:

Liquid crystals (LC) are substances that flow like liquids but maintain some of the ordered structure characteristics of crystalline solids. Based on the ways that LCs are generated, they can be classified into two types 1) Thermotropic LCs and 2) Lyotropic LCs. The Thermotropic is generated by temperature variation in the liquid state, while the Lyotropic is formed by dissolving the compound in certain solvents. Either way offers the systems sufficient large amplitude molecular mobility so that molecules can change positions and reorient themselves forming LC phases. The thermotropic LCs usually are one-compound systems, while lyotropic LCs are always solutions consisting of multiple compounds containing solute and solvent. The aim of this review article is to scientifically highlight Pharmaceutical Liquid crystals, its importance in current development of targeted drug delivery system. Objective of this review is also to provide in depth information of pharmaceutical liquid crystal technology which include its latest & advanced technology development.

Keywords: Pharmaceutical liquid crystals, Mesomorphs, Liquid crystal Emulsions,

Introduction:

Liquid Crystals are intermediate state between solid and liquid state. It is often called a mesomorphic state which is state of matter in which the degree of molecular order is intermediate between the perfect three-dimensional, long-range positional and orientational order found in solid crystals and the absence of long-range order found in isotropic liquids, gases, and amorphous solids¹ it is also called as meso intermediate. The liquid crystals are of thermotropic and lyotropic types lyotropic liquid crystals are induced by the presence of solvent, and have been extensively described elsewhere in the context of emulsion technology; however, other pharmaceutical examples are emerging. Thermotropic liquid crystals are induced by a change in temperature and are essentially free of solvent, where more pharmaceutical applications appear in the literature².

Liquid crystals (LC) are liquids featuring a certain level of orientational order. Specifically, molecules in LCs tend to point to a certain direction, while they still have translational (positional) freedom. Although they are best known for their application in displays, liquid crystals are also an essential part of all life forms. Lyotropic liquid crystals are essential organic substances, DNA, lipids of cellular

Membranes and proteins are some examples of well known liquid crystals. In liquid crystals drug delivery crystalline solids exhibit short as well as long-range order with regard to both position and orientation of the molecules. Whereas liquids are amorphous in general but may show short-range order with regard to position and/or orientation. Liquid crystals show at least orientational long-range order and may show short-range order, whereas positional long-range order disappears.

Accordingly, liquid crystalline phases represent intermediate states and are also called mesophases.⁸

Applications:

Liquid Crystal Emulsion

A large part of cosmetic products are made in the form of emulsions, a form that allows the simultaneous use of lipophilic and hydrophilic ingredients in the required dosages. A product in the form of an emulsion also has the advantage of having the most convenient appearance and texture that also facilitates its application. They can be formulated to be liquid, milk type emulsions of variable consistency, creams, or even super liquid spray able emulsions. Finally, we should also consider the fact that an emulsion is the best carrier for active ingredients and functional substances.

The theory of stabilising an emulsion through the formation of a network of liquid crystals is different to the HLB theory or the Schulman couple.s theory. The gelification of the water phase obtainable with hydrosolvatable polymers or with emulsifiers that are able to form a reticular organised structure in liquid crystal form, eliminates the need to use waxy components in large quantities and consistency factors that are no longer in harmony with the modern conception of light and easy to spread emulsions

Advantage of Liquid crystals in emulsions:

LCs (mesophases) provides the following advantages to emulsion.

1. Increased stability
 2. Prolonged hydration
 3. Controlled drug delivery
- 1) **Stability:** Emulsion stability of the multilayers around the oil droplets act as a barrier to coalescence. If oil droplets coalesce emulsion breaks. This barrier for coalescence acts as increased stability property of the emulsion
- 2) **Prolonged hydration:** Lamellar liquid crystalline and gel network contain water layer, which shows that 50% of the water of oil in water (o/w) emulsion can be bound to such structures. Such water is less prone to evaporation when applied to the skin and

permits a long lasting moisturisation / hydrating effect, necessary for drug entry.

- 3) **Controlled Drug delivery:** Liquid crystals prevent the fast release of the drug dissolved in the oil phase of an emulsion. This is attributed to the lamellar liquid crystalline multilayer, which reduces the interfacial transport of a drug dissolved within the oil droplets. Microscopic observations under polarized light show the exceptional thickness of liquid crystalline lamellar layer around the oil droplets

Function & properties of the liquid crystal system

LCs When present at the oil/water interface, the liquid crystals help give the system rigidity and, by limiting the fluctuation of the components at the interface, give the emulsion great stability. Furthermore, the liquid crystal system enhances the moisturising ability of the emulsion; in this special network. The quantity of inter-lamellar water can be extremely high and become immediately available when the cream is applied to the skin.

For these reasons these emulsions have a shiny surface, a fresh and original feel and they leave a light and pleasant sensation on the skin.

In recent years, the moisturising effect of creams and lotions has become increasingly more important and cosmetic chemists are constantly searching for better methods of retaining water in the superior layers of the skin. The evaporation of the bonding water in emulsions containing anisotropic lamellar phases is slower and permits a hydroretentive action that prolongs the moisturising effect. The associations that are formed because of the excess water are particularly interesting; in these cases the ability of the crystalline phase to swell is strictly linked to the stability and the behaviour of the emulsion because, in a liquid crystal system, the quantity of inter-lamellar water and of hydrophile elements can amount to 70% of the total external phase

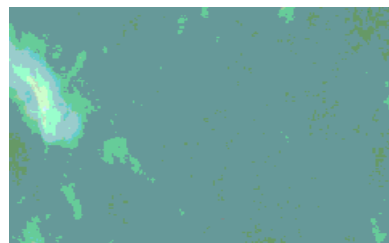


Fig. 1 Image of Liquid crystal

Controlled Release of Bioactive Materials:

The release of the active substance from liquid crystalline delivery systems is often controlled by diffusion, and some systems using the photoinduced or thermal phase transition of the liquid crystals as the release for bioactive materials¹⁴

Drug loading

According to the nature of the drug, it can be added in both the aqueous as well as oil phase. Loading totally depends on solubility of active constituents and their partition between existing phases.

Drug can be incorporated in liquid crystal technology:¹⁵

Drugs	
Cefazolin	Cefuroxime
Clomethiazole	Clindamycin phosphate
4- Phenylbutylamine, prilocaine	Oestriol
Isosorbide mononitrate	Insulin
Indomethacin	Clotrimazole gramicidin
Nitroglycerin	Lidocaine Hydrochloride

Latest Development in Pharmaceutical Liquid crystal technology

Kent State University, Summa Health System and IC-MedTech have developed a new

paradigm in drug discovery based on the pharmacologic properties of Liquid Crystal Pharmaceuticals (LCPs). LCPs are a unique class of lyotropic liquid crystals that represent novel drug candidates for the treatment of a wide range of diseases. Representatives of this research project recently filed applications for two new patents: one for a new LCP-based anti-tumor drug called **Tolecine**TM and another for a formulation that combines Tolecine and another LCP, Apatone.⁶

Olecine is a new pre-clinical anti-tumor LCP that also has antiviral and antibacterial applications. It is tumor-cell selective and exhibits a strong anti-neoplastic activity (it counteracts abnormal proliferation of cells in a tissue or organ). In addition, it has been shown to be more effective than the current standard of care for herpes.

Apatone is a clinical phase investigational new drug for late-stage prostate cancer. It is currently under consideration for clinical study for potential applications such as augmentation of chemotherapy to allow lower, less toxic doses of common chemotherapeutic agents. Apatone is made from two non-toxic compounds, a liquid crystal compound and a sugar, that selectively bio-concentrate within cancer cells and produce a free radical. Formation of the strong, short lived free radical is a concentration driven intracellular reaction and therefore only takes place in cells with sufficient sugar concentrations (such as cancer cells). The reaction results in oxidative stress that weakens the targeted cells from within. It is carried out quickly only within the cancer cells, with no toxic reaction by-products that might harm adjacent healthy cells.

Patents with Liquid Crystals Applications:

Sr. No	Title of Patent	Patent /Date	No Abstract
1	Liquid crystals containing cosmetic and pharmaceutical compositions and methods for utilizing such compositions	US 4,999,348 / Mar.12, 1991	Cosmetic and pharmaceutical compositions and methods comprising delivery systems for the controlled release and enhanced penetration of biologically active material (e.g. Vitamin A) to the skin. The delivery systems comprise cholesteric liquid crystals wherein the active material is retained within the lamellar molecular structure (i.e. between the molecular sheets) of the cholesteric liquid crystals ³

2	Liquid crystals emulsion type pharmaceutical composition containing cyclosporine and therapeutic method of treating cutaneous disease therewith	US 2010/190695 A1/ July 29,2010	A dermal external pharmaceutical composition that excels in feeling at application or after application and that by enhancing of the transdermal absorption of cyclosporine exerts medicinal benefits at low concentration. There is provided a liquid crystals emulsion- type pharmaceutical composition comprising cyclosporine a hydrophilic nonionic surfactant, a lipophilic nonionic surfactant, an oil a fatty acid that is insoluble in the oil at room temperature, a solid fatty alcohol that is insoluble in the oil at room temperature and a water soluble polyhydric alcohol that is immiscible with the oil at room temperature, and a method of testing cutaneous diseases with the use of the pharmaceutical composition. ⁴
3	Drug delivery system utilizing liquid crystal structure	US 5,891,845/ April 6,1999	Vitamin E TPGS/drug composition and method are provided which obviate the need for surfactants or non evaporated co solvents because the active drug component is dissolved directly into vitamin E TPGS to form a true molecular solution –not an emulsion or a micro emulsion the invention provides a slowly dissolving TPGS/Drug matrix that absorbs gastrointestinal fluid into the matrix at crystal is formed. This gel front forms a liquid crystals boundary where drug dissolution is highest. At this liquid crystals/GI fluid boundary, synchronization takes place in which the rate of formation of liquid crystals equals the dissolution rate of liquid crystals at the water interface thereby giving controlled order release of the drug into the GI tract. The rate of dissolution is also controlled by the geometry of the dosage form. The solid vitamin E TPGS/ drug matrices of the inventions can be solidified and compressed into tablets or filled into capsules, with other excipients, binder and /or fillers. The solid TPGS/drug solution of the invention also can be made into an immediate release liquid formulation upon addition of water or into a controlled release system solid tablet by the use of impermeable or semi permeable barriers or coating surroundings portions of the tablet. ⁵
4	Transdermal pharmaceutical composition	10/575,145/June 1, 2010	The invention relates to a liquid crystal gel containing polyoxyethylene-glyceryl-trioleate, propylene-glycol, isopropyl myristate and a hyaluronic acid salt or complex for use in the manufacture of transdermal pharmaceutical compositions and healing cosmetics. The invention also relates to transdermal pharmaceutical composition consists of an estrogen and a progestin component as well as a liquid crystal gel containing polyoxyethylene-

5	Silicone Liquid Crystals, Vesicles, And Gels	US 6608126 / 8/19/2003	glyceryl-trioleate, propylene-glycol, isopropyl myristate and a hyaluronic acid salt or complex. The invention can be applied for transdermal hormone replacement therapy and for other transdermal depending on the active principles included. ⁷
6	Imine Based Liquid Crystals For The Controlled Release Of Bioactive Materials	20090306196	Silicone liquid crystals, silicone vesicles, and silicone gels are formed by combining (i) an amine functional silicone or an organic amine compound, (ii) an organic epoxide containing at least two epoxy groups or an epoxy functional silicone containing at least two epoxy groups, and (iii) a fluid such as water, a silicone fluid other than a silicone used in (i) and (ii), and a polar organic compound or nonpolar organic compound other than organic compounds used in (i) and (ii) in which the polar and nonpolar organic compound are present at a concentration to provide a level of solids less than 40 percent by weight based on the weight of (i)-(iii). These silicone liquid crystals, silicone vesicles, and silicone gels are useful in personal care for preparing various skin, hair, and underarm products. ¹⁰ The present invention relates to a delivery system based on a film of an imine forming liquid crystalline phase, mixed with at least with one biologically active substance, to which a constant or variable electric field can be applied. The delivery system can be used as a delivery system for biologically active substances such as flavors, fragrances, bactericides, fungicides, insecticides, insect attractants or repellents, agrochemicals or pharmaceuticals ¹⁴

Conclusion:

Liquid crystals technique for drug delivery can be an effective and useful for delivering the drug with desired target. This technique can be utilized largely in topical delivery of the drug as it posses the advantages of smooth feel and drug loading of incompatible molecule. Day by day various patents are coming in this liquid crystals technique. This technique of drug loading is a new & immerging techniques requires attention further for its practice in real scientific industries to deliver a quality outcome for a society

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References:

- 1) IUPAC Compendium of Chemical Terminology IUPAC Gold book 2005.
- 2) Stevenson CL, Bennett DB, Lechuga-Ballesteros D Pharmaceutical liquid crystals: the relevance of partially ordered systems J Pharm Sci. 2005 Sep; 94 (9):1861-80.
- 3) Gheorge Cioca, Lake Grove, James Hayward, Port jeffeson, Manuel L.Tan, Glen Clove, Morris Herstein scarsdae, N.Y.walter, Stamford, conn

- “Liquid crystals containing cosmetic and pharmaceutical compositions and methods for utilizing such compositions” US 4,999,348 March 12, 1991
- 4) Ryo Akamastu, Masahiro Fujii, Tomoki Sakaguchi, Eijiro Horisawa Japan “Liquid crystals emulsion type pharmaceutical composition containing cyclosporine and therapeutic method of treating cutaneous disease therewith” US 2010/190695 A1/ July 29, 2010
- 5) Garry Myers, Kingsport, Tenn. “Drug delivery system utilizing liquid crystal structure” US 5,891,845/ April 6, 1999
- 6) <http://thefutureofthings.com/news/1058/liquid-crystals-battle-cancer-and-other-diseases.html>
- 7) Eros, et al. “Transdermal pharmaceutical composition” 10/575,145/June 1, 2010
- 8) Christel C. Mueller-Goymann, “Drug Delivery: Liquid Crystals in” Encyclopedia of Pharmaceutical Technology, Chapter 75, Third Edition Informa Healthcare
- 9) Imran K. Tadwee, Sourabh Gore, Prashant Giradkar “Advances in Topical Drug Delivery System: A Review” Int. J. of Pharm. Res. & All. Sci. 2012; Volume 1, Issue 1, 14-23
- 10) Michael Salvatore Ferritto; Zuchen Lin; William James Schulz Jr. “Silicone Liquid Crystals, Vesicles, And Gels” US 6608126 / 8/19/2003
- 11) J. C. Shah et al., Adv. Drug. Delivery Rev., 2001, 47, 229-250,
- 12) C. C. Muller-Goymann, Eur. J. Pharm. Biopharm., 2004, 58, 343-356
- 13) B. J. Boyd, Surfact. Sci. Ser., 2005, 127, 285-305
- 14) Jean-Marie Lehn, Andreas Herrmann Nicolas Giuseppone “Imine Based Liquid Crystals For The Controlled Release Of Bioactive Materials” Patent App. No 20090306196
- 15) P. V. Patel, J. B. Patel, R. D. Dangar, K. S. Patel, K. N. Chauhan “Liquid Crystal Drug Delivery System” International Journal of Pharmaceutical and Applied Sciences/1 (1)/2010 118-123.
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Liquid crystals (LCs) are a state of matter which has properties between those of conventional liquids and those of solid crystals. For instance, a liquid crystal may flow like a liquid, but its molecules may be oriented in a crystal-like way. There are many different types of liquid-crystal phases, which can be distinguished by their different optical properties (such as textures). The contrasting areas in the textures correspond to domains where the liquid-crystal molecules are oriented in different Pharmaceutical co-crystals - a review. 104. other drug molecules as co-crystallization partners has been reported 53. Scientists have observed that liquid-assisted grinding of two enantiomorphous co-crystals resulted in to either formation of a centrosymmetric three-component co-crystal, consisting of the left- and right-handed co-crystal former molecules and the model API or the model API along with the racemic form of the co-crystal former 74. Applications: Liquid Crystal Emulsion. A large part of cosmetic products are made in the form of emulsions, a form that allows the simultaneous use of lipophilic and hydrophilic ingredients in the required dosages. A product in the form of an emulsion also has the advantage of having the most convenient appearance and texture that also facilitates its application. They can be formulated to be liquid, milk type emulsions of variable consistency, creams, or even super liquid spray able emulsions. Finally, we should also consider the fact that an emulsion is the best carrier for active ingredient... Imran Tadwee, Dr. Sadhana Shahi, Vivek Ramteke, Syed Iftakar

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