

## Nanoemulsion: A Promising and Novel Nanotherapeutic Vehicle for Transdermal Drug Delivery Application

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**Key words:** Nano-emulsion, skin, bioavailability, surfactants, transdermal, drug delivery, stratum corneum

**Abstract:** Since, transdermal delivery of therapeutics have been identified by scientific researchers as an alternative choice of drug administration, owing to its unique and tremendous benefit over conventional and oral administrations, special formulation that is capable of meeting up and overcoming its associated challenges is required. Nano-emulsion is one of the nano-technological formulations that have gained wide exposure as a novel drug delivery system that solves pharmaceutical challenges as well as improves dermal and transdermal drug delivery. The objective of this review is to create the basic understanding of the skin as an alternative transdermal delivery route while presenting nano-emulsions as a unique therapeutic vehicle. In this study, improvement in drug delivery application using nano-emulsions with particular attention to drug delivery transdermally is reviewed. The basic understanding and considerations in nano-emulsion generation including the required understanding of the skin as a possible transdermal drug delivery route is highlighted. The identity of nano-emulsions reflects a positive and successful application in the delivery of nano-therapeutics to the systemic circulation through the skin.

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### INTRODUCTION

In recent years, there has been an overwhelming interest in the dermal and percutaneous delivery of

pharmaceuticals as well as bioactives and the use of nanoemulsion as the vehicle reflects a good future in drug delivery<sup>[1]</sup>. Transdermal drug delivery is an innovative research area that is becoming prominent globally as a

result of its unique merits over other routes<sup>[2]</sup>. This system uniquely achieves the therapeutic concentration of drug through skin with proper delivery at a controlled rate, making it very conducive for the treatment of both chronic and acute disorders<sup>[3, 4]</sup>.

Transdermal drug delivery system constitutes a novel research area of drug delivery that is successfully emerging as it helps to overcome drawback associated with parenteral routes<sup>[5]</sup>. Its ability to promote therapeutic efficacy can be based on its improving bioavailability of drug, non-invasive nature, patient's accessibility, pain free nature, ability to avoid hepatic first pass metabolism and facilitation of self-medication in patients<sup>[6-8]</sup>. In addition, extensive investigation shows this route as a substitute to oral route of drug transport because the rich supply of blood by the dermis ease the unswerving passage of drug into the blood for systemic circulation and effects<sup>[9-12]</sup>.

In contrast, the biophysical properties of the skin impose a barrier towards enhanced permeation of drug through the stratum corneum<sup>[13]</sup>. Also, limitations are associated to percutaneous delivery of semi-solid formulations as a result of their large particle size, slow skin entry, fast volatilization of highly vaporous compounds, environmental disintegration, etc.<sup>[8]</sup>. In order to promote drug permeation through this significant barrier, chemical and physical methods such as iontophoresis, microneedle array, laser and thermal ablation including chemical penetration enhancers have been explored<sup>[9, 11, 14]</sup>. However, these techniques pose limitations such as irritation and damage to the skin barrier. Higher dosage of chemical enhancers could also lead to increased toxicity<sup>[14]</sup>. Extensive investigation have also been carried out in the area of nanotechnology to be able to deliver less soluble, highly volatile as well as photosensitive and charged surface<sup>[15]</sup>.

In view of the best technique to display safe and improve drug delivery approach as evident from the literature, nanoemulsion-based technology is expected to fit by Shakeel *et al.*<sup>[16]</sup>. Nanoemulsion based technology allows the combined possibility of enhancing penetration across the skin by altering the lipid bilayer and at the same time, acts as tiny storage channel for drugs<sup>[17]</sup>. Considering its flowing (fluidic) nature, notable interrelation with skin cells, adequate permeation ability, small globule size, ability to protect and deliver even irritant as well as compound of high volatility and molecular weight makes it the best<sup>[1]</sup>. Nanoemulsion droplet are able to overcome the barrier posed by the skin by penetrating the skin pores and arrive the systemic circulation, getting successfully channelized for effective delivery of drugs<sup>[18]</sup>. This delivery vehicle also offers advantages such as great solubilizing ability of both hydrophilic and hydrophobic drugs, reduced cost of

preparation, reduced viscosity with Newtonian behavior, thermodynamic and increased storage stability and protection of bioactive ingredients against oxidation and hydrolysis<sup>[7, 13, 19]</sup>.

The prospect significance of this area of research have extended the hand of researchers into the challenging issues associated with the formulation of nanoemulsions and its applicability in the treatment of crucial and challenging issues associated with transdermal delivery route. In this manuscript, some of the vital findings in various literatures were represented, thereby discussing nanoemulsions and their association with the skin barrier. The skin physiology and mechanism of transdermal drug transport is also reviewed.

### Nanoemulsion basics

**Nanoemulsion and its classification:** Nanoemulsions or submicron emulsions are optically isotropic system composed of a mixture of two immiscible liquids (mainly oil and aqueous medium) resulting in a fine dispersion of drugs in nanodroplets with small size distribution<sup>[20-22]</sup>. They are kinetically stabled colloidal particulate system with uniform size distribution stabilized by an interfacial layer of suitable surfactant<sup>[8, 23]</sup>.

Nanoemulsions can be classified based on their 'components' and 'surface charge' over the nanodroplets. For the former, nanoemulsions can be grouped into:

- (a) Oil-in-Water (O/W) nanoemulsions: it is a type of nanoemulsion formed when an oil phase is dispersed into an aqueous medium
- Water-in-Oil (W/O) nanoemulsions: it is a type of nanoemulsion formed when water is dispersed into an oil medium
- Bicontinuous nanoemulsion, i.e., interdispersion of micro-domains of oil and water phase<sup>[19, 24]</sup>

The latter classifies nanoemulsions into Neutral nanoemulsions, Anionic nanoemulsions and Cationic nanoemulsions-where, neutral, anionic and cationic surfactants are used, respectively<sup>[21, 25]</sup>.

Recent research have proven O/W nanemulsions to be of high popularity in medicine than the W/O counterpart owing to its compatibility with water, safe and wide use in drug delivery<sup>[26, 27]</sup>.

**Nanoemulsion composition:** Basically, nanoemulsions comprises oil, surfactants /emulsifiers including co-surfactants/co-emulsifiers and the aqueous phase<sup>[28]</sup>. Figure 1 represents an illustration of an O/W nanoemulsion including the general composition of a nanoemulsion. Oil Phase is one of the important adjuvants in nanoemulsions where its solubilizing ability in relation to drugs (phytoactives or synthetic drugs) of choice is an important factor in choosing oil. Depending on the chain

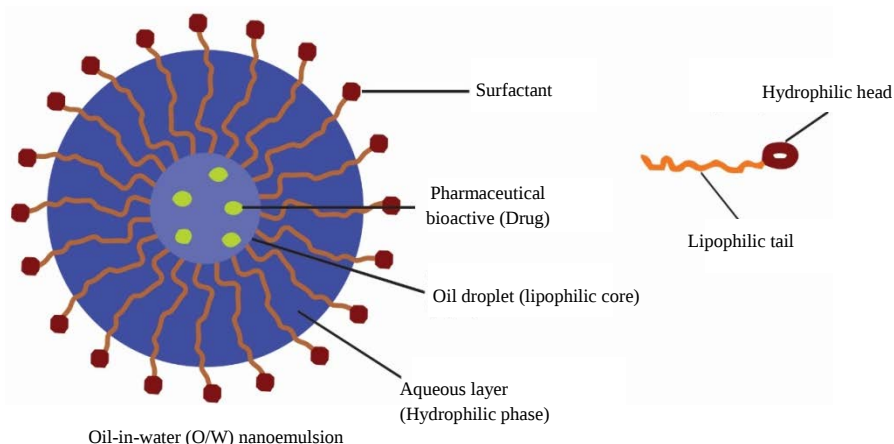


Fig. 1: An illustration of an O/W nanoemulsion including the general composition of a nanoemulsion

lengths, long chain triglycerides and medium chain triglycerides are used either alone or in combination for nanoemulsion formulation. Recent studies have reported that nanoemulsion can be employed to improve bioavailability of drugs with poor water solubility such as curcumin by solubilizing<sup>[29]</sup>. Also, permeability of cytotoxic drugs such as piplartine can be enhanced<sup>[30]</sup> into cancer cells.

An increased interest have aroused in the use of natural product in the field of drug delivery due to their preferred advantages. These include low toxicity and widespread therapeutic benefits<sup>[31]</sup>. The use of essential oils as oil phase in nanoemulsion formulation is not only safe and biocompatible but are volatile natural compounds that are susceptible to oxidation, thermal and photo degradation<sup>[32]</sup> with complex fatty acids composition known to have diverse benefits<sup>[33]</sup>.

Gundel *et al.*<sup>[32]</sup>, recently developed nanoemulsion that contains Basil oil. The objective is to determine its stability as well as cytotoxicity in healthy cells of human with the antioxidant including antimicrobial potential evaluation. Result indicated reduced activity of the free oil and the antioxidant. Also, cytotoxicity was not represented but the antimicrobial activity proves effective<sup>[32]</sup>.

Ge *et al.*<sup>[34]</sup>, prepared a complex comprising a phospholipid and Dabigatran Etexilate (DE) nanoemulsion matrix. The formulation displayed an improved dissolution in stimulated intestinal fluids. This is carried out by containing DE into an oil phase. DE, (a direct thrombin inhibitor for treatment of thromboembolism) in a phospholipid complex (DE-PC) showed significant reduction ( $p < 0.05$ ) in drug leakage. Relatively, bioavailability of DE-PC was found to increase to 147.3 and 606.6%<sup>[34]</sup>.

Tumtibsri *et al.*<sup>[35]</sup> also carried out a recent research on spearmint oil (SMO) which was incorporated into a nanoemulsion. Spearmint oil acts as a cytotoxic agent to

averse oral carcinoma cell line using MTT assay. SMO is mostly used in oral hygiene products and it has numerous interesting potentials, especially its anticancer property. Unfortunately, its application in combating cancer is reduced because it is naturally insoluble in water. In this study, nanoemulsion was chosen as the alternative vehicle for SMO to impede oral cancer cells. Surfactant amount and type, oil loading as well as the proportion of SMO in relation to Virgin Coconut Oil (VCO) were studied. The SMO-VCO nanoemulsions showed notable cytotoxic effect against oral carcinoma (KON) cell line using MTT assay. The investigation reveals the improved feasibility of SMO-VCO nanoemulsions as potential unique carrier for oral cancer treatment<sup>[35]</sup>.

Aqil<sup>[36]</sup> investigated the development and optimization of clove oil with olmesartan nanoemulsion formulation in transdermal application using the Box-Behnken design. Result suggested that the developed nanoemulsion carrier of the omelsartan provides  $53.11 \pm 3.13$  nm as the particle size,  $0.335 \pm 0.008$  as the polydispersity index with a transdermal flux of  $12.65 \pm 1.60 \mu\text{g cm}^{-3} \text{h}^{-1}$ . Confocal laser scanning microscope unveiled an improved entry of the Rhodamine-B loaded nanoemulsion, extending far down into the deeper layers of the skin<sup>[36]</sup>.

Barabbas *et al.*<sup>[37]</sup>, developed and characterized o/w Hydrogel Thickened Nanoemulsion (HTN). The transdermal delivery of 8-methoxysalen (8-MOP) encapsulated in an HTC comprising sweet fennel oil as well as clove essential oil as the oil phase was studied. The incorporation of 8-MOP into two nanoemulsions that encompasses either sweet fennel or clove essential oil as oil phase led to the development of formulation that brings possibility in the modulation of drug delivery transdermally and skin retention. This formulation proves to be promising for photo-chemotherapy and also guaranteed increased efficacy and reduced systemic exposure<sup>[37]</sup>.

Surfactants are amphiphilic molecules which absorb readily at oil and water interface thereby providing steric or electrostatic stabilization<sup>[17]</sup>. Surfactant renders a repulsive surface force while stabilizing the droplets against aggregation and coalescence<sup>[38, 39]</sup>. In nanoemulsion, lower concentration of surfactants are required, exposing nanoemulsions to thermodynamic instability<sup>[17, 40, 41]</sup> but they end up being more kinetically stable due to reduction from macroemulsion to nanoemulsion after external application of shear/energy<sup>[42]</sup>. Since, lower surfactant concentration is required, little or no toxicity is accompanied with the generated nanoemulsion.

Surfactants used in nanoemulsion are collected considering their toxicity profile, solubility in nanoemulsion phases, i.e., the oil and water, together with their emulsification ability, i.e., Hydrophile-Lipophile (HLB) value. Nonionic surfactants are the preferred choice of surfactants for therapeutic purpose because of their reduced toxicity as well as less irritant compared to the ionic counterparts<sup>[8, 28]</sup>. Also, the use of O/W nanoemulsions for medicinal purpose requires non-ionic surfactants and this is likely to render *in-vivo* stability<sup>[43]</sup>. Choosing suitable surfactant is a critical factor. To improve stability of nanoemulsions a right blend of low and high HLB surfactants should be considered upon dilution with water. Selection of a surfactant blend not only show great effect on size and stability of nanoemulsion but sometimes also determines its toxicity, pharmacokinetics including pharmacodynamics<sup>[44, 45]</sup>. Surfactants that has their HLB values between 4 and 6 generally find application in stabilizing W/O emulsions while surfactants that has their HLB values between 8 and 18 stabilizes O/W nanoemulsions. Recently, sugar based natural surfactants are employed because of their skin and environmental friendliness<sup>[46]</sup>.

Co-surfactants in nanoemulsion could also be added in low concentration to further reduce the interfacial tension by complementing structurally weaker areas<sup>[47]</sup>. Independent use of single chain emulsifier might not be prove enough ability to bring down the interfacial tension to a large extent, hence, it's use with a co-surfactant that possesses an amphiphilic character is also encouraged. A co-emulsifier is able to further penetrate the interfacial layer thereby reducing the fluidity and elevating the entropy of the emulsion system<sup>[48]</sup>.

Pengon *et al.*<sup>[26]</sup>, investigated the role surfactants play in the physical properties a nanoemulsion prepared with coconut oil. Different amount of surfactants that includes, polyoxyethylene sorbitan monostearate (POS), Polyethylene glycol Hydrogenated Castor oil (PHC), Polyethylene Glycol phenyl ether (PGO), Sodium Lauryl Sulphate (SLS) and poloxamer 407 (PLX) were used. From the result, it was indicated that Coconut oil

nanoemulsion exhibited small percent creaming index when the surfactants employed were: POS, PGO and PHC<sup>[26]</sup>. Yin *et al.*<sup>[48]</sup> also conducted a research on biocompatible nanoemulsion using hemp oil with less surfactant for the oral delivery of baicalein with enhanced bioavailability. Aside the major role of surfactants in the preparation and stability of nanoemulsions, surfactants can also upset the skin barrier function and play a penetration enhancing role to these systems<sup>[46]</sup>.

**Methods of nanoemulsion preparation:** Formulation of nanoemulsion is an important and crucial aspect that needs to be considered before the formation of a nanoemulsion. Various methods of nanoemulsion development are discussed in this review in order to help scientist for proper selection of the suitable method of formulation. Numerous methods have been developed for nanoemulsion formulation but can be grouped into high and low energy methods<sup>[27, 49, 50]</sup>.

**The high energy methods:** Here, high shear forces are required. The nanoemulsion prepared using this technique requires external energy in the form of a high shear stress to generate a nanosized emulsion<sup>[41, 51]</sup>. These methods employs specially designed mechanical devices to break the intermolecular forces that exist between the liquids<sup>[16]</sup> by intense shear turbulence and cavitation flow profile<sup>[49]</sup>. The most common techniques employed in the high energy/intensity emulsification are Microfluidization; High Pressure Homogenization (HPH); Sonication Method and production with High Magnitude Ultrasound<sup>[52-54]</sup>.

In the Sonication method, a sonication mechanism is employed to reduce the droplet size of conventional emulsion to form small batches of nanoemulsions<sup>[55]</sup>. Here, the system is exposed to substantial amplitude of sound wave in the presence of a sonicator probe. Cavitation occurs and during this process, there exists generation of sound waves and energy is supplied to the emulsion droplets which are coarse. Due to these sound waves, globule size becomes reduced to nanosized droplets with optimum frequency and time<sup>[41]</sup>.

In High pressure Homogenization (HPH), a very intense pressure is supplied into the emulsion system that contains oily phase, aqueous (water) phase, surfactants and co-surfactants with the aid of high pressure homogenizer. O/W nanoemulsion that contains <20% oil can be prepared with this method<sup>[56]</sup>. And the fluid is finally, discharged as an homogenized nanoemulsion<sup>[57]</sup>. Limitation of this method aside its selectivity also includes deterioration of active ingredients due to exposure of the emulsion system to a large amount of heat<sup>[17, 53, 54, 58]</sup>.

Microfluidization of an emulsion system is a technique that also generates nanosized droplets through

its mechanism of flow that occurs through microchannels<sup>[41]</sup>. It is a modified or revised high pressure homogenization that works with a positive displacement pump of high pressure (500-20,000 psi) that pressurize the liquid to pass along micro-channels arranged in it. The materials passing through the micro-channels breaks down<sup>[59]</sup> into very fine particle. The liquid phases (oil phase and aqueous phase) give a coarse emulsion in an inline homogenizer after processing. This coarse emulsion then undergoes further processing in the microfluidizer for generation of stable nanoemulsions of submicron range<sup>[7, 55]</sup>.

High amplitude ultrasound is a technique that requires high frequency sound waves supported with significant amplitude, hence, the process "Ultrasonication". The system creates uneven air current in the solution with pressure fluctuation being generated for vibration at specific frequency<sup>[19, 60]</sup>. In this technique, adjusting of experimental conditions such as temperature, amplitude, frequency as well as sonication time, nanoemulsions with appropriate nanosized droplets can be obtained. However, this technique is not acceptable for industrial application because of variation in particle size during large scale manufacturing. Also, the technique is capable of decomposing surfactants in surfactant based formulations due to the effect of the high frequency sound waves<sup>[42, 61, 62]</sup>.

From this review, the advantages and disadvantages of high energy method can be summarized. High energy methods can be said to be of advantages in respect to the fact that nanoemulsions are produced from a flexible sample of materials and they can easily be scaled to industrial quantities. However, their limitation can be associated to their exposure to an intense external energy input requirements and hence, highly expensive to operate. Also, newly created interface suffer from stabilization due to insufficient surfactant<sup>[38]</sup>.

**Low energy methods:** This method, unlike high intensity (energy) methods, rely on spontaneous generation of small droplets whenever the system composition experience a phase change or inversion as a way of responding to alterations generated from changes in composition of surfactants or temperature of the system<sup>[63, 64]</sup>. In this method, significantly less input of energy density is needed since energy required can readily be attained in a simple magnetic stirring system<sup>[63]</sup>. These methods are typically named as 'Phase inversion' method because it usually involves the generation of thermodynamically-stable nanoemulsion by altering the conditions (temperature and composition) which affects the HLB of the emulsion system<sup>[65]</sup>. Alteration in physicochemical parameters could lead to changes in properties of surfactants which enforce phase transformation making them either fat loving or hydrophilic molecules. The two most important low

energy methods includes: Phase Inversion Composition or Emulsion Inversion Point (PIC or EIP, respectively) and Phase Inversion Temperature (PIT)<sup>[19, 59]</sup>.

In PIT method, the solubilization pattern of the surfactants (hydrophilic to lipophilic) is changed by increasing the temperature of the emulsion system which forms bicontinuous microemulsions followed by emulsion inversion<sup>[7]</sup>. The principle of this technique is dependent on the specific properties of temperature sensitive surfactants which have the capability of changing its distribution affinity towards water and oil (HLB) with temperature change. Nonionic surfactants, usually the polyethoxylated surfactants are mostly chosen in this regard because of their temperature sensitivity. At low temperature, the surfactant tends to be hydrophilic because of the presence of hydrated polar groups, leading to O/W system. When temperature increases, dehydration of polyoxyethylene group makes it hydrophobic because of the presence of aliphatic groups<sup>[7, 66]</sup>. At HLB temperature, there is a common affinity for water and the oil, hence, kinetically stable nanoemulsion with narrow droplet size distribution and low polydispersity<sup>[42]</sup>. The main challenge associated to this particular method is that thermo-labile agents and other excipients cannot be employed as temperature plays major role in PIT<sup>[59, 67]</sup>.

Emulsion Inversion Point (EIP) is also known as Phase Inversion Composition (PIC) method since the composition of surfactant plays an important role in the phase transformation of the system<sup>[42]</sup>. At a constant temperature, phase transitions are produced by the composition of dilution of dispersed phase to convert<sup>[68]</sup>. In PIC, W/O macroemulsions are generated at ambient temperature, followed by slow dilution using water. In the process of dilution what happens is that the system undergoes a transition where inversion from W/O to O/W emulsification occurs. At this inversion point, the interfacial tension of the oil-water interface becomes greatly reduced, therefore, nano-droplets having high specific surface area is obtained without significant energy or pressure required<sup>[69, 70]</sup>.

## **Nanoemulsions through transdermal**

### **Route: Basic considerations**

**The skin:** The skin is the most extensive multilayered organ of the body having a surface expanse of 1.7-2.0 m<sup>2</sup><sup>[71]</sup>. The skin not only covers the body surface but also serves as a shield for the internal organs against direct exposure to the external environment. Amongst its numerous functions are: immunity against microorganisms, electrolytic balance, protection against and from physical injuries, temperature regulation ability UV radiation and harmful chemical substances<sup>[72]</sup>. Additionally, the skin serves as a route of absorption for drugs as well as an avenue for expression of drug efficacy<sup>[72-74]</sup>.

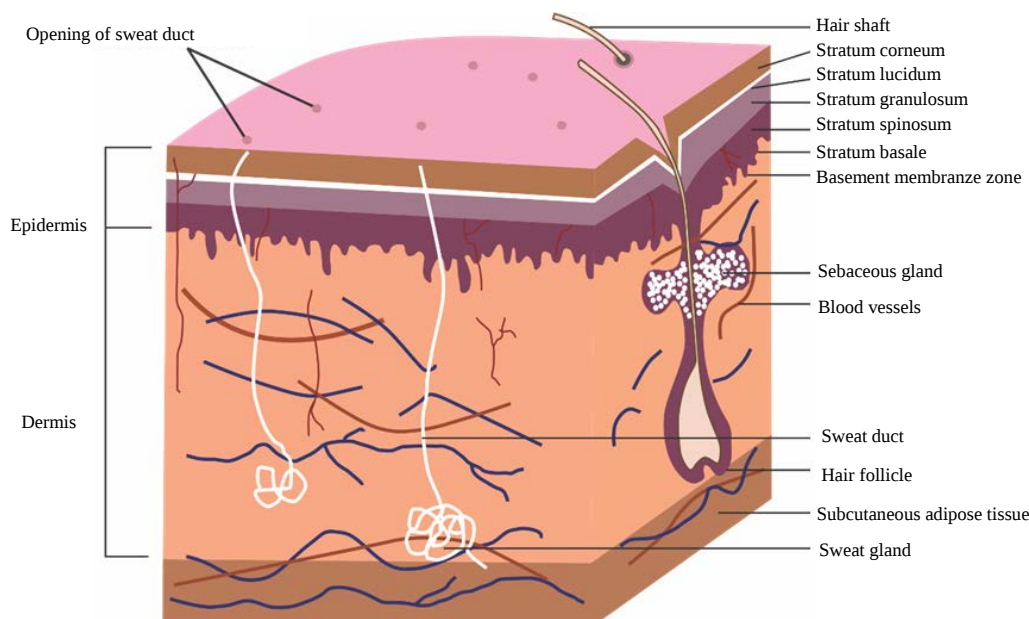


Fig. 2: The skin structure representing three layers microscopically, namely: epidermis, dermis and subcutaneous layers

As presented in Fig. 2, the skin represents three layers microscopically, namely: epidermis, dermis and subcutaneous layer, i.e., the outer, middle and the inner layer, respectively<sup>[58]</sup>. The Epidermis which is composed of several cell layers, can be subdivided into Viable Epidermis (VE) and non-viable epidermis. The non-viable epidermis is also known as Stratum Corneum (SC) where the SC represents the outermost layer of the epidermis while the viable epidermis represents the beneath layer<sup>[75,76]</sup>. The Stratum Corneum (SC) consists of keratin filled corneocytes that are surrounded by lipophilic matrix which comprises of esters, ceramides, cholesterol and fatty acids<sup>[71]</sup>. Corneocytes are flat dead cells that contain a huge amount of water including keratin filaments. The Corneocytes further have interconnections with some special juncture called Corneodesmosomes<sup>[75]</sup>. The complexity of the SC can therefore be summarized as described by Menon *et al.*<sup>[77]</sup> as 'brick and mortar' structure with the corneocytes representing the bricks stuffed in a mortar (the surrounding lipophilic matrix), hence, the barrier of this layer for transdermal drug permeation, especially, hydrophilic drugs<sup>[77]</sup>.

The dermis represents the thick layer of the skin with 500-3000  $\mu\text{m}$  depth. This layer contains lymph vessels, blood vessels, sweat glands and nerve endings. It is the middle layer of the skin<sup>[78]</sup>. The subcutaneous fatty layer forms the most inner layer of the skin. It lies below the dermis and it contains adipose tissue (fat). It is responsible for providing support such as nutrition, temperature regulation and physical protection<sup>[78]</sup>.

#### **Position of the skin in transdermal delivery:**

Transdermal drug delivery systems refer to the delivery of drug across the skin in order to accomplish local and systemic therapeutic effect<sup>[79]</sup>. Transdermal drug delivery is an attractive alternative to conventional administration of drugs owing to its numerous advantages<sup>[80]</sup> including convenient administration route of the drug, ease of accessibility, non-invasive nature and increase in patient compliance<sup>[81]</sup>. This mode of drug administration increasingly becomes a research focus in pharmaceutical preparation. Its numerous potential and features also includes the reduction in the fluctuation of drug concentration in the blood, easy drug detection, reduced chances of overdose, provision of steady plasma levels and escape from the gastrointestinal environment: pH and enzymatic reactions<sup>[82]</sup>. Most importantly is its ability to shelve the first pass effect, increases the therapeutic efficacy of drug and increases their long time stability because there will be little or no drug interference<sup>[83]</sup>. Despite its long list of advantages, drug absorption through the skin seems to be a bit difficult<sup>[84]</sup>. Drug permeation through the skin can be a little bit challenging. Its transport through the skin to systemic circulation can be divided into three steps:

- Penetration: release of drug from the vehicle/dosage form and passage into external layer of the skin
- Permeation: passage of the drug from one layer of the skin to the other
- Resorption: penetration of the drug into systemic circulation through the blood and lymph vessels within the dermis

The transepidermal (across SC) or the transappendegeal are the two potential pathways for drug permeation through the skin<sup>[71]</sup>. The SC route can take place as intra or intercellular penetration where the intracellular route passes through the corneocytes allowing hydrophilic substance to be transported in and out of cell membrane. On the other hand, intercellular transport allows transportation of lipophilic drugs around corneocytes in the lipid rich extracellular regions. For the transappendegeal route, transportation of substances are granted through the sebaceous glands, across hair follicles and sweat glands<sup>[9, 84]</sup>.

Animal model are usually explored in assessing penetration properties of drug candidates for human consumption. This is because ethical as well as practical issues have made specimens associated to human skin to be very less available. Hence, alternative models, especially models of animal skin have been developed<sup>[85]</sup>.

**Nanoemulsions and transdermal drug delivery: means of operation:** As earlier discussed, delivery of drugs through the skin experiences high resistance against diffusion into blood pool because upper layer of the skin are the most restricted layer, thereby making diffusion across the SC a rate limiting step for drugs<sup>[41]</sup>. To overcome this difficulty, nanoformulations seem to be the required choice. Amongst the most favorable nanoformulations for improvement of transdermal permeation and delivery of drugs is nanoemulsion technique<sup>[86]</sup>. Dermally applied nanoemulsion penetrates the stratum corneum and firmly exists in the whole horny layer altering both lipid and polar pathways<sup>[87]</sup>. Their small particle size brings better result on 'drug retention', 'drug specificity' and 'drug targeting' with these three making an ideal transdermal drug delivery<sup>[88]</sup>.

Nanoemulsion, a unique aspect of nanoformulations are isotropic, low viscous mixture with extremely small droplet size comprising of clear or translucent oil globules that are dispersed in an aqueous phase with stability offered through an interfacial surfactant or co-surfactant film<sup>[17]</sup>. Since, the droplet or particle size of nanoemulsions exists below 25% of the wavelength of visible light, they tend to come out clear or transparent<sup>[89]</sup>. Due to these unique particle size, large surface area and low surface tension, nanoemulsions show great wettability which make them to maintain a close contact with the skin. They also give shorter transdermal time and better percutaneous absorption<sup>[74]</sup>.

Bakshi *et al.*<sup>[90]</sup>, reported the topical delivery of nanoemulsion-based formulation containing heparinoid, an anticoagulant derived from heparin. Its potential in delivering a transdermal therapy towards the treatment of superficial thrombophlebitis was studied. Results

indicated that the developed O/W nanoemulsion transported greater amount of heparinoid ( $91.25 \pm 25.75$  and  $62.29 \pm 5.66 \mu\text{g cm}^{-2}$ ), respectively after 72 h<sup>[90]</sup>.

Aqil (2015) also developed a clove oil based nanoemulsion for transdermal supply of olmesartan, a lipophilic and an anti-hypertensive drug. The optimized formulation revealed a small droplet size of  $53.11 \pm 3.13$  nm with an enhanced bioavailability of Olmesartan proven by an improved penetration through the rat skin<sup>[36]</sup>.

Transdermal delivery of Diacerein that was formulated with glucosamine sulphate laden oil in water nanoemulsion as the homing carrier was reported by Chattopadhyay and Datta<sup>[91]</sup>. The nanoemulsion system was reported to contain an oleic acid oil phase with Tween 20 and PEG 400 as the surfactants. The result revealed a powerful permeation ability and good stability, thereby identifying nanoemulsion as a potential carrier to achieve an enhanced topical transport of lipophilic diacerein<sup>[91]</sup>.

The success of nanoemulsion in promoting drug penetration transdermally as well as achieving therapeutic success can be traced to its channel of operation. The first agent that can be considered is the nanoemulsion composition: the oil and the surface active agents<sup>[92]</sup>. According to Shakeel *et al.*<sup>[92]</sup>, increased solubility of the drug in the oil phase, surfactants and co-surfactants efficiently improves the extent at which the nanoemulsion system maintains the drug in the solubilized form. This leads to a better permeability for improved permeability and successful delivery. Also, since nanoemulsion contains both lipids and aqueous phase, it is expected that if the drug gets dissolved in the lipid domain of the nanoemulsion, direct penetration of the lipid layer of the SC can take place, causing destabilization of its bilayer structure<sup>[92]</sup>.

This interaction leads to a high increase in the drug permeability through the transdermal lipid pathway. In contrast, the hydrophilic aspect of nanoemulsion can largely serve as a source of hydration to the SC resulting in transdermal uptake of drugs. Possibility of this can be explained that when the aqueous fluid of nanoemulsions migrates into the polar pathway, the interlamellar volume of the stratum corneum lipid bilayer increases, thereby resulting in disarrangement of its interfacial structure<sup>[93]</sup>. The oil component of nanoemulsion finds compatibility with the sebum, an oily secretion from the sebaceous gland found in the follicular openings. These openings are alternate entry point that opens up chances of leading and trapping drugs with therapeutic capabilities to clinically active skin lesions<sup>[17]</sup>.

The presence of surfactants in nanoemulsions increases skin permeation by altering the cellular integrity of the skin in several ways making the stratum corneum become hydrated<sup>[94]</sup>. Both oil and surfactants interact

readily with different layers of the skin and assist in the distribution of loaded moiety to deeper part of the skin<sup>[95]</sup>.

Moustafa *et al.*<sup>[96]</sup> reported his study on transdermal cumin oil nanoemulsions for effective and increased systemic antioxidant and hepato-protective properties. Surfactant/co-surfactant used in this study was tween 20/ethanol. Oleic acid /capryol were employed as the oil phase. From the results, phenolic content of Cumin oil showed increased encapsulation efficiency, including remarkable phenol permeation across the rat skin was recorded. From the different surfactant mix prepared, the formulation containing cumin/oleic acid as the oil phase Tween 20/ethanol (2:1) as surfactant/cosurfactant and distilled water as the aqueous phase showed the highest favorable formula.

This was proven from the *in vitro* and *in vivo* antioxidant effect and a great hepato-protective potential after 7 days of single transdermal application<sup>[96]</sup>.

Elmateeshy *et al.*<sup>[97]</sup> developed a novel nanoemulgel for enhanced transdermal permeability of Terbinafine (TB), an antifungal drug with inadequate water solubility. The formulation which comprises Peceol (oil phase) and Tween 80/propanol (Surfactant/co-surfactant mix) were prepared in different weight ratio which ranges from 1:9-9:1. From the pseudo ternary phase diagram, required formulae were obtained, followed by formulation and characterization of nanoemulsions. The thermodynamic stability studies including the *in vitro* drug release studies were also carried out. The nanoemulsion formulae containing 10 or 15% w/w oil, 45% w/w  $S_{mix}$  (1:2/1:3) and 45-40% w/w aqueous phase) were selected as the optimum formulae and incorporated into Carbopol 940 gel bases to form three groups of TB nanoemulsion formulated emulgel formulae (F1-F3). These were investigated for *ex vivo* drug permeation as well as *in vivo* antifungal activity. Results indicated that all the prepared nanoemulsion based gel formulae reveals significant ( $p < 0.05$ ) improvement when compared to the commercially prepared emulgel<sup>[97]</sup>.

Rachmawati<sup>[98]</sup> also studied a curcumin nanoemulsion for transdermal evaluation. The nanoemulsion prepared by self nanoemulsification method includes glyceryl monooleate as oil phase, cremophor RH-40 and PEG 400 as surfactant/co-surfactant. *In vitro* permeation of curcumin was studied using casted (shed) snake skin of a python reticulatus and a modified vertical diffusion cell. Evaluation of nanoemulsions was carried out using particle size analyzer, Polydispersity Index (PDI), zeta potential, physical stability and Raman spectra. The overall study revealed improved curcumin permeability; it also protected curcumin from chemical degradation<sup>[98]</sup>.

Another factor that contributes to permeability effect of nanoemulsion through skin is the little particle size and

high specific surface area of the nanoemulsion<sup>[99]</sup>. Transdermal delivery is made possible when the size of the nanoemulsion droplet is low. Permeability of nanoemulsions through the skin is inversely proportional to the magnitude of the nanoemulsion droplet. A nanoemulsion with droplet size above 150 nm may find difficulty in penetrating the skin efficiently, limiting its application topically while a nanoemulsion droplet below 60 nm ensures great possibilities in transdermal delivery<sup>[100]</sup>.

Other important factors could be associated to the special and unique structure of nanoemulsions which gives them the capability of solubilizing both lipophilic and hydrophilic drugs. When this occurs, a greater chance of deeper drug penetration is allowed since a high permeable concentration gradient will take place both in and out of the skin<sup>[101]</sup>. Also, nanoemulsion ability to act as tiny reservoirs for drugs gives them inner storage ability, thereby prolonging drug absorption. This is because diffusion of drug is constantly maintained because of the forceful pull from the external phase to the inner phase<sup>[100]</sup>.

## CONCLUSION

Nowadays, alternative route of drug delivery is highly invested as parenteral route seems challenging. Transdermal route of drug delivery is now embraced strongly as it has confidently reflected positivity and uniqueness. To resolve challenges of drug bioavailability, improved solubility, increased penetrability and enhanced therapeutic effectiveness, nanoemulsions have been considered. The identity of nanoemulsions has advantageously play outstanding part in the improved transdermal delivery of drugs. In this review, the components and transdermal mechanism of nanoemulsions is summarized with the aim to improve understanding of the role of nanoemulsions in transdermal drug delivery systems. This review encourages further research in the field drug delivery for improved transdermal treatment.

## REFERENCES

01. Shakeel, F. and W. Ramadan, 2010. Transdermal delivery of anticancer drug caffeine from water-in-oil nanoemulsions. *Colloids Surfaces B Biointerfaces*, 75: 356-362.
02. Rao, R., S. Mahant, L. Chhabra and S. Nanda, 2014. Transdermal innovations in diabetes management. *Current Diabetes Rev.*, 10: 343-359.
03. Ahad, A., F.I. Al-Jenoobi, A.M. Al-Mohizea, M. Aqil and K. Kohli, 2013. Transdermal delivery of calcium channel blockers for hypertension. *Expert Opin. Drug Deliv.*, 10: 1137-1153.

04. Ahad, A., A.M. Al-Mohizea, F.I. Al-Jenoobi and M. Aqil, 2016. Transdermal delivery of Angiotensin II Receptor Blockers (ARBs), Angiotensin-Converting Enzyme Inhibitors (ACEIs) and others for management of hypertension. *Drug Delivery*, 23: 579-590.
05. Desai, P., R.R. Patlolla and M. Singh, 2010. Interaction of nanoparticles and cell-penetrating peptides with skin for transdermal drug delivery. *Mol. Membr. Biol.*, 27: 247-259.
06. Paudel, K.S., M. Milewski, C.L. Swadley, N.K. Brogden, P. Ghosh and A.L. Stinchcomb, 2010. Challenges and opportunities in dermal/transdermal delivery. *Ther. Delivery*, 1: 109-131.
07. Sutradhar, K.B. and M.L. Amin, 2013. Nanoemulsions: Increasing possibilities in drug delivery. *Eur. J. Nanomed.*, 5: 97-110.
08. Rai, V.K., N. Mishra, K.S. Yadav and N.P. Yadav, 2018. Nanoemulsion as pharmaceutical carrier for dermal and transdermal drug delivery: Formulation development, stability issues, basic considerations and applications. *J. Controlled Release*, 270: 203-225.
09. Alkilani, A.Z., M.T. McCrudden and R.F. Donnelly, 2015. Transdermal drug delivery: Innovative pharmaceutical developments based on disruption of the barrier properties of the stratum corneum. *Pharmaceutics*, 7: 438-470.
10. Barakat, N., E. Fouad and A. Elmedany, 2011. Formulation design of indomethacin-loaded nanoemulsion for transdermal delivery. *Pharm Anal. Acta S.*, 2: 1-8.
11. Kogan, A. and N. Garti, 2006. Microemulsions as transdermal drug delivery vehicles. *Adv. Colloid Interface Sci.*, 123: 369-385.
12. Bruno, B.J., G.D. Miller and C.S. Lim, 2013. Basics and recent advances in peptide and protein drug delivery. *Ther. Delivery*, 4: 1443-1467.
13. Yamada, M., H. Tayeb, H. Wang, N. Dang and Y.H. Mohammed *et al.*, 2018. Using elongated microparticles to enhance tailorable nanoemulsion delivery in excised human skin and volunteers. *J. Controlled Release*, 288: 264-276.
14. Ahad, A., M. Aqil, K. Kohli, H. Chaudhary, Y. Sultana, M. Mujeeb and S. Talegaonkar, 2009. Chemical penetration enhancers: A patent review. *Expert Opin. Therap. Patents*, 19: 969-988.
15. Ghalandarlaki, N., A.M. Alizadeh and S. Ashkani-Esfahani, 2014. Nanotechnology-applied curcumin for different diseases therapy. *BioMed Res. Int.*, Vol. 2014, 10.1155/2014/394264
16. Shakeel, F., S. Shafiq, N. Haq, F.K. Alanazi and I.A. Alsarra, 2012. Nanoemulsions as potential vehicles for transdermal and dermal delivery of hydrophobic compounds: An overview. *Expert Opin. Drug Delivery*, 9: 953-974.
17. Singh, Y., J.G. Meher, K. Raval, F.A. Khan, M. Chaurasia, N.K. Jain and M.K. Chourasia, 2017. Nanoemulsion: Concepts, development and applications in drug delivery. *J. Controlled Release*, 252: 28-49.
18. Ravi T. and P. Thiagarajan, 2011. Nanoemulsions for drug delivery through different routes *Biotechnol.*, 2: 1-13.
19. Jaiswal, M., R. Dudhe and P.K. Sharma, 2015. Nanoemulsion: An advanced mode of drug delivery system. *3 Biotechnol.*, 5: 123-127.
20. Sood, S., K. Jain and K. Gowthamarajan, 2014. Optimization of curcumin nanoemulsion for intranasal delivery using design of experiment and its toxicity assessment. *Colloids Surf. B. Biointerfaces*, 113: 330-337.
21. Kumar, D., J. Ali and S. Baboota, 2016. Omega 3 fatty acid-enriched nanoemulsion of thiocolchicoside for transdermal delivery: Formulation, characterization and absorption studies. *Drug Delivery*, 23: 591-600.
22. Gao, F., Z. Zhang, H. Bu, Y. Huang and Z. Gao *et al.*, 2011. Nanoemulsion improves the oral absorption of candesartan cilexetil in rats: Performance and mechanism. *Journal of Controlled Release* 149: 168-174.
23. Javadzadeh, Y. and L.A. Bahari, 2017. Therapeutic Nanostructures for Dermal and Transdermal Drug Delivery. In: *Nano-and Microscale Drug Delivery Systems: Design and Fabrication*, Grumezescu, A.M. (Ed.), Elsevier, Amsterdam, Netherlands, pp: 131-146.
24. Sigward, E., N. Mignet, P. Rat, M. Dutot and S. Muhamed *et al.*, 2013. Formulation and cytotoxicity evaluation of new self-emulsifying multiple W/O/W nanoemulsions. *Int. J. Nanomed.*, 8: 611-625.
25. Patil, P.A. and B.R. Bhutkar, 2016. Biomedical application of nanoemulsion. *Int. J. Res. Methodol.*, 1: 37-58.
26. Pengon, S., C. Limmatvapirat and S. Limmatvapirat, 2009. Preparation and evaluation of antimicrobial nanoemulsion containing herbal extracts. *Drug Metab. Rev.*, 41: 85-85.
27. Gupta, A., H.B. Eral, T.A. Hatton and P.S. Doyle, 2016. Nanoemulsions: Formation, properties and applications. *Soft Matter.*, 12: 2826-2841.
28. Azeem, A., M. Rizwan, F.J. Ahmad, Z. Iqbal, R.K. Khar, M. Aqil and S. Talegaonkar, 2009. Nanoemulsion components screening and selection: A technical note. *Aaps Pharmscitech*, 10: 69-76.
29. Vecchione, R., V. Quagliarillo, D. Calabria, V. Calcagno, E. De Luca, R.V. Iaffaioli and P.A. Netti, 2016. Curcumin bioavailability from oil in water nano-emulsions: *In vitro* and *in vivo* study on the dimensional, compositional and interactional dependence. *J. Controlled Release*, 233: 88-100.

30. Fofaria, N.M., H.S.S. Qhattal, X. Liu and S.K. Srivastava, 2016. Nanoemulsion formulations for anti-cancer agent piplartine-Characterization, toxicological, pharmacokinetics and efficacy studies. *Int. J. Pharm.*, 498: 12-22.
31. Chouhan, S., K. Sharma and S. Guleria, 2017. Antimicrobial activity of some essential oils-present status and future perspectives. *Medicines*, Vol. 4, No. 3. 10.3390/medicines4030058
32. Gundel, S.D.S., M.C. Velho, M.K. Diefenthaler, F.R. Favarin and P.M. Copetti *et al.*, 2018. Basil oil-nanoemulsions: Development, cytotoxicity and evaluation of antioxidant and antimicrobial potential. *J. Drug Delivery Sci. Technol.*, 46: 378-383.
33. Bajarski, L., L.R. Michels, L.M. Colome, E.A. Bender and R.J. Freddo *et al.*, 2016. The use of Brazilian vegetable oils in nanoemulsions: An update on preparation and biological applications. *Braz. J. Pharm. Sci.*, 52: 347-363.
34. Ge, L., X. He, Y. Zhang, Y. Zhang and F. Chai *et al.*, 2018. A dabigatran etexilate phospholipid complex nanoemulsion system for further oral bioavailability by reducing drug-leakage in the gastrointestinal tract. *Nanomed. Nanotechnol. Biol. Med.*, 14: 1455-1464.
35. Tubtimsri, S., C. Limmatvapirat, S. Limsirichaikul, P. Akkaramongkolporn, Y. Inoue and S. Limmatvapirat, 2018. Fabrication and characterization of spearmint oil loaded nanoemulsions as cytotoxic agents against oral cancer cell. *Asian J. Pharm. Sci.*, 13: 425-437.
36. Aqil M., M. Kamran, A. Ahad and S.S. Imam, 2016. Development of clove oil based nanoemulsion of olmesartan for transdermal delivery: box-behnken design optimization and pharmacokinetic evaluation. *J. Mol. Liq.*, 214: 238-248.
37. Barradas, T.N., J.P. Senna, S.A. Cardoso, S. Nicoli and C. Padula *et al.*, 2017. Hydrogel-thickened nanoemulsions based on essential oils for topical delivery of psoralen: Permeation and stability studies. *Eur. J. Pharm. Biopharm.*, 116: 38-50.
38. Helgeson, M.E., 2016. Colloidal behavior of nanoemulsions: Interactions, structure and rheology. *Curr. Opin. Colloid Interface Sci.*, 25: 39-50.
39. Wulff-Perez, M., A. Martin-Rodriguez, M.J. Galvez-Ruiz and J. de Vicente, 2013. The effect of polymeric surfactants on the rheological properties of nanoemulsions. *Colloid Polymer Sci.*, 291: 709-716.
40. Anton, N. and T.F. Vandamme, 2011. Nanoemulsions and micro-emulsions: Clarifications of the critical differences. *Pharmaceut. Res.*, 28: 978-985.
41. Patel, M.R., R.B. Patel and S.D. Thakore, 2018. Nanoemulsion in Drug Delivery. In: *Applications of Nanocomposite Materials in Drug Delivery*, Inamuddin, M.A. Abdullah and A. Mohammad (Eds.), Woodhead Publishing, Cambridge, UK., pp: 667-700.
42. Koroleva, M.Y. and E.V. Yurtov, 2012. Nanoemulsions: The properties, methods of preparation and promising applications. *Russ. Chem. Rev.*, 81: 21-43.
43. Kawakami, K., T. Yoshikawa, T. Hayashi, Y. Nishihara and K. Masuda, 2002. Microemulsion formulation for enhanced absorption of poorly soluble drugs: II. *In vivo* study. *J. Controlled Release*, 81: 75-82.
44. Ali, M.S., M.S. Alam, N. Alam and M.R. Siddiqui, 2014. Preparation, characterization and stability study of dutasteride loaded nanoemulsion for treatment of benign prostatic hypertrophy. *Iranian J. Pharm. Res. (IJPR.)*, 13: 1125-1140.
45. Li, S.J., Y.F. Liao and Q. Du, 2018. Research and application of quercetin-loaded nano drug delivery system. *China J. Chin. Materia Med.*, 43: 1978-1984.
46. Isailovic, T.M., M.N. Todosijevic, S.M. Dordevic and S.D. Savic, 2017. Natural Surfactants-Based Micro/Nanoemulsion Systems for NSAIDs-Practical Formulation Approach, Physicochemical and Biopharmaceutical Characteristics/Performances. In: *Microsized and Nanosized Carriers for Nonsteroidal Anti-Inflammatory Drugs*, Calija, B. (Ed.). Academic Press, Cambridge, Massachusetts, pp: 179-217.
47. Zhang, Y., Z. Shang, C. Gao, M. Du, S. Xu, H. Song and T. Liu, 2014. Nanoemulsion for solubilization, stabilization and *in vitro* release of pterostilbene for oral delivery. *Aaps Pharm.*, 15: 1000-1008.
48. Yin, J., C. Xiang, P. Wang, Y. Yin and Y. Hou, 2017. Biocompatible nanoemulsions based on hemp oil and less surfactants for oral delivery of baicalein with enhanced bioavailability. *Int. J. Nanomed.*, 12: 2923-2931.
49. McClements, D.J. and S.M. Jafari, 2018. General Aspects of Nanoemulsions and Their Formulation. In: *Nanoemulsions*, Jafari, S.M. and D.J. McClements (Eds.). Academic Press, Cambridge, Massachusetts, pp: 3-20.
50. Shams, N. and M.A. Sahari, 2016. Nanoemulsions: Preparation, structure, functional properties and their antimicrobial effects. *Applied Food Biotechnol.*, 3: 138-149.
51. Tayeb, H.H. and F. Sainsbury, 2018. Nanoemulsions in drug delivery: Formulation to medical application. *Nanomedicine*, 13: 2507-2525.
52. Singh, T.G.D., J. Manish, I.S. Sandhu and M. Chitkara, 2016. Nanobiomaterials: Applications in Biomedicine and Biotechnology. In: *Fabrication and Self-Assembly of Nanobiomaterials*, Grumezescu, A.M. (Ed.). Elsevier Science B.V., Amsterdam, pp: 409-421.
53. Debnath, S., K.V. Satayanarayana and V.K. Gampa, 2011. Nanoemulsion-a method to improve the solubility of lipophilic drugs. *Pharmanest*, 2: 72-82.

54. Reza, K.H., 2011. Nanoemulsion as a novel transdermal drug delivery system. *Int. J. Pharm. Sci. Res.*, 2: 1938-1946.
55. Sharma, N., S. Mishra, S. Sharma, R.D. Deshpande and R.K. Sharma, 2013. Preparation and optimization of nanoemulsions for targeting drug delivery. *Int. J. Drug Dev. Res.*, 5: 37-48.
56. Floury, J., A. Desrumaux, M.A. Axelos and J. Legrand, 2003. Effect of high pressure homogenisation on methylcellulose as food emulsifier. *J. Food Eng.*, 58: 227-238.
57. Mason, T.G., J.N. Wilking, K. Meleson, C.B. Chang and S.M. Graves, 2006. Nanoemulsions: Formation, structure and physical properties. *J. Phys. Condens. Matt.*, 18: 635-643.
58. Jain, S., N. Patel, M.K. Shah, P. Khatri and N. Vora, 2017. Recent advances in lipid-based vesicles and particulate carriers for topical and transdermal application. *J. Pharm. Sci.*, 106: 423-445.
59. Yukuyama, M.N., D.D.M. Ghisleni, T. Pinto and N.A. Bou Chacra, 2016. Nanoemulsion: Process selection and application in cosmetics-a review. *Int. J. Cosmetic Sci.*, 38: 13-24.
60. Quintanilla-Carvajal, M.X., B.H. Camacho-Diaz, L.S. Meraz-Torres, J.J. Chanona-Perez, L. Alamilla-Beltran, A. Jimenez-Aparicio and G.F. Gutierrez-Lopez, 2010. Nanoencapsulation: A new trend in food engineering processing. *Food Eng. Rev.*, 2: 39-50.
61. Maali, A. and M.H. Mosavian, 2013. Preparation and application of nanoemulsions in the last decade (2000-2010). *J. Dispersion Sci. Technol.*, 34: 92-105.
62. Kale, S.N. and S.L. Deore, 2017. Emulsion micro emulsion and nano emulsion: A review. *Syst. Rev. Pharm.*, 8: 39-47.
63. Komaiko, J.S. and D.J. McClements, 2016. Formation of food grade nanoemulsions using low energy preparation methods: A review of available methods. *Compr. Rev. Food Sci. Food Safety*, 15: 331-352.
64. Forgiarini, A.M., J. Esquena, C. Gonza'lez and C. Solans, 2001. Formation and Stability of Nanoemulsions in Mixed Non Ionic Surfactants System. In: *Trends in Colloids and Interface Science*, Koutsoukos, P. (Ed.). Springer, Berlin, Germany, pp: 184-189.
65. Solc, I., A. Maestro, C.M. Pey, C. Gonzalez, C. Solans and J.M. Gutierrez, 2006. Nano-emulsions preparation by low energy methods in an ionic surfactant system. *Colloids Surf. A. Physicochem. Eng. Aspects*, 288: 138-143.
66. Date, A.A., N. Desai, R. Dixit and M. Nagarsenker, 2010. Self-nanoemulsifying drug delivery systems: Formulation insights, applications and advances. *Nanomedicine*, 5: 1595-1616.
67. Lovelyn, C. and A.A. Attama, 2011. Current state of nanoemulsions in drug delivery. *J. Biomater. Nanobiotechnol.*, 2: 626-639.
68. Sharma, S., S. Kumar and R. Gupta, 2012. A review on transdermal drug delivery. *IJAPBC.*, 1: 100-110.
69. Forgiarini, A., J. Esquena, C. Gonzalez and C. Solans, 2000. Studies of the Relation between Phase Behavior and Emulsification Methods with Nanoemulsion Formation. In: *Trends in Colloid and Interface Science XIV*, Buckin, V. (Eds.). Springer, Berlin, Germany, pp: 36-39.
70. Izquierdo, P., J. Esquena, T.F. Tadros, C. Dederen, M.J. Garcia, N. Azemar and C. Solans, 2002. Formation and stability of nano-emulsions prepared using the phase inversion temperature method. *Langmuir*, 18: 26-30.
71. Czajkowska-Kosnik, A., M. Szekalska and K. Winnicka, 2019. Nanostructured lipid carriers: A potential use for skin drug delivery systems. *Pharmacological Rep.*, 71: 156-166.
72. Yasin, Z.A., F. Ibrahim, N.N. Rashid, M.F. Razif and R. Yusof, 2017. The importance of some plant extracts as skin anti-aging resources: A review. *Curr. Pharm. Biotechnol.*, 18: 864-876.
73. Ventrelli, L., M.L. Strambini and G. Barillaro, 2015. Microneedles for transdermal biosensing: Current picture and future direction. *Adv. Healthcare Mater.*, 4: 2606-2640.
74. Zhou, X., Y. Hao, L. Yuan, S. Pradhan, K. Shrestha and O. Pradhan *et al.*, 2018. Nano-formulations for transdermal drug delivery: A review. *Chin. Chem. Lett.*, 29: 1713-1724.
75. Bouwstra, J.A., H.E.J. Hofland, F. Spies, G.S. Gooris and H.E. Junginger, 1992. Changes in the Structure of the Human Stratum Corneum Induced by Liposomes. In: *Liposome Dermatics*, Braun-Falco, O., H.C. Korting and H.I. Maibach (Eds.), Springer, Berlin, Germany, pp: 121-136.
76. El Maghraby, G.M., B.W. Barry and A.C. Williams, 2008. Liposomes and skin: From drug delivery to model membranes. *Eur. J. Pharm. Sci.*, 34: 203-222.
77. Menon, G.K., G.W. Cleary and M.E. Lane, 2012. The structure and function of the stratum corneum. *Int. J. Pharm.*, 435: 3-9.
78. Chen, X., 2018. Current and future technological advances in transdermal gene delivery. *Adv. Drug Delivery Rev.*, 127: 85-105.
79. Hao, Y., W. Li, X. Zhou, F. Yang and Z. Qian, 2017. Microneedles-based transdermal drug delivery systems: A review. *J. Biomed. Nanotechnol.*, 13: 1581-1597.
80. Schoellhammer, C.M., D. Blankschtein and R. Langer, 2014. Skin permeabilization for transdermal drug delivery: Recent advances and future prospects. *Expert Opin. Drug Delivery*, 11: 393-407.
81. Barry, B.W., 1987. Mode of action of penetration enhancers in human skin. *J. Controlled Release*, 6: 85-97.

82. Marwah, H., T. Garg, A.K. Goyal and G. Rath, 2016. Permeation enhancers strategy in transdermal drug delivery. *Drug Deliv.*, 23: 564-578.
83. Singh, K., N. Arora and T. Garg, 2013. RFID: A trustable security tool in pharmaceutical industry. *Am. J. Pharm. Tech. Res.*, 2: 113-127.
84. Shahzad, Y., R. Louw, M. Gerber and J. Du Plessis, 2015. Breaching the skin barrier through temperature modulations. *J. Controlled Release*, 202: 1-13.
85. Abd, E., S.A. Yousef, M.N. Pastore, K. Telaprolu and Y.H. Mohammed *et al.*, 2016. Skin models for the testing of transdermal drugs. *Clin. Pharmacol. Adv. Appl.*, 8: 163-176.
86. Osborne, D.W., A.J.I. Ward and K.J. O'Neill, 1991. Microemulsions as topical drug delivery vehicles: *In vitro* transdermal studies of a model hydrophilic drug. *J. Pharm. Pharmacol.*, 43: 451-454.
87. Thacharodi, D. and K.P. Rao, 1994. Transdermal absorption of nifedipine from microemulsions of lipophilic skin penetration enhancers. *Int. J. Pharm.*, 111: 235-240.
88. Liu, J., Q.W. Chen and K. Wu, 2017. On-surface construction of low-dimensional nanostructures with terminal alkynes: Linking strategies and controlling methodologies. *Chin. Chem. Lett.*, 28: 1631-1639.
89. Balamohan, P., C.H. Anjali and A. Ravindran, 2013. Nanoemulsion: Synthesis, characterization and its applications. *J. Bionanosci.*, 7: 323-333.
90. Bakshi, P., Y. Jiang, T. Nakata, J. Akaki, N. Matsuoka and A.K. Banga, 2018. Formulation development and characterization of nanoemulsion-based formulation for topical delivery of heparinoid. *J. Pharm. Sci.*, 107: 2883-2890.
91. Chattopadhyay, H. and S. Datta, 2018. Transdermal delivery of Diacerein with homing carrier Glucosamine sulphate laden in oil-in-water nanoemulsion. *Mater. Today Proc.*, 5: 9690-9697.
92. Shakeel, F., S. Baboota, A. Ahuja, J. Ali and S. Shafiq, 2008. Skin permeation mechanism and bioavailability enhancement of celecoxib from transdermally applied nanoemulsion. *J. Nanobiotechnol.*, Vol. 6. 10.1186/1477-3155-6-8
93. Baboota, S., F. Shakeel, A. Ahuja, J. Ali and S. Shafiq, 2007. Design, development and evaluation of novel nanoemulsion formulations for transdermal potential of celecoxib. *Acta Pharm.*, 57: 315-332.
94. Schneider, M., F. Stracke, S. Hansen and U.F. Schaefer, 2009. Nanoparticles and their interactions with the dermal barrier. *Dermato-Endocrinol.*, 1: 197-206.
95. Lboutounne, H., Y.C. Guillaume, L. Michel, S. Makki, P.H. Humbert and J. Millet, 2004. Study and development of encapsulated forms of 4, 5', 8 trimethylpsoralen for topical drug delivery. *Drug Dev. Res.*, 61: 86-94.
96. Mostafa, D.M., A.A. Kassem, M.H. Asfour, S.Y. Al Okbi, D.A. Mohamed and T.E.S. Hamed, 2015. Transdermal cumin essential oil nanoemulsions with potent antioxidant and hepatoprotective activities: *In-vitro* and *in-vivo* evaluation. *J. Mol. Liq.*, 212: 6-15.
97. Elmataeeshy, M.E., M.S. Sokar, M. Bahey-El-Din and D.S. Shaker, 2018. Enhanced transdermal permeability of Terbinafine through novel nanoemulgel formulation; Development *in vitro* and *in vivo* characterization. *Future J. Pharm. Sci.*, 4: 18-28.
98. Rachmawati, H., D.K. Budiputra and R. Mauludin, 2015. Curcumin nanoemulsion for transdermal application: Formulation and evaluation. *Drug Dev. Ind. Pharm.*, 41: 560-566.
99. Zhou, H., Y. Yue, G. Liu, Y. Li and J. Zhang *et al.*, 2010. Preparation and characterization of a lecithin nanoemulsion as a topical delivery system. *Nanoscale Res. Lett.*, 5: 224-230.
100. Khurana, S., N.K. Jain and P.M.S. Bedi, 2013. Nanoemulsion based gel for transdermal delivery of meloxicam: Physico-chemical, mechanistic investigation. *Life Sci.*, 92: 383-392.
101. Lawrence, M.J. and G.D. Rees, 2000. Microemulsion-based media as novel drug delivery systems. *Adv. Drug Delivery Rev.*, 45: 89-121.