

RESEARCH ARTICLE

Open Access



Statistical optimization of tretinoin-loaded penetration-enhancer vesicles (PEV) for topical delivery

Neda Bavarsad^{1,2*}, Abbas Akhgari³, Somayeh Seifmanesh², Anayatollah Salimi^{1,2} and Annahita Rezaie⁴

Abstract

Background: The aim of this study was to develop and optimize deformable liposome for topical delivery of tretinoin.

Methods: Liposomal formulations were designed based on the full factorial design and prepared by fusion method. The influence of different ratio of soy phosphatidylcholine and transcucol (independent variables) on incorporation efficiency and drug release in 15 min and 24 h (responses) from liposomal formulations was evaluated. Liposomes were characterized for their vesicle size and Differential Scanning Calorimetry (DSC) was used to investigate changes in their thermal behavior. The penetration and retention of drug was determined using mouse skin. Also skin histology study was performed.

Results: Particle size of all formulations was smaller than 20 nm. Incorporation efficiency of liposomes was 79–93 %. Formulation F7 (25:5) showed maximum drug release. Optimum formulations were selected based on the contour plots resulted by statistical equations of drug release in 15 min and 24 h. Solubility properties of transcucol led to higher skin penetration for optimum formulations compared to tretinoin cream. There was no significant difference between the amount of drug retained in the skin by applying optimum formulations and cream. Histopathological investigation suggested optimum formulations could decrease the adverse effect of tretinoin in liposome compared to conventional cream.

Conclusion: According to the results of the study, it is concluded that deformable liposome containing transcucol may be successfully used for dermal delivery of tretinoin.

Keywords: Statistical optimization, Tretinoin, Liposomal formulation, Topical delivery

Background

Tretinoin or all-trans retinoic acid in topical form is commonly used for the treatment of various skin problems like acne, photoaging and psoriasis. Furthermore, it has other functions such as sebum production, collagen synthesis and regulating growth and differentiation of epithelial cells [1]. Reduction the size and the number of comedones are considered to be the main effect of tretinoin in the treatment of acne [2]. However, it has several negative points even in topical use such as very low

water solubility, skin irritation and high instability in the presence of air, light and heat. Furthermore, local irritation such as erythema, peeling and burning at the application site and increased susceptibility to sunlight are its side effects [3].

Liposomes are spherical-shaped carriers which has an internal aqueous portion surrounded by one or multiple concentric lipidic bilayers. Liposomes are used as carriers for both lipophilic and water soluble molecules. Hydrophilic substances are encapsulated in the interior aqueous portions whereas lipophilic substances are entrapped within lipid bilayers [4].

Incorporation of tretinoin in nanostructure systems such as liposomes may lead to decrease the adverse effects and protect this molecule against degradation [5–8].

* Correspondence: nbavarsad@ajums.ac.ir

¹Nanotechnology Research Center, Ahvaz Jundishapur University of Medical Sciences, Ahvaz, Iran

²Department of Pharmaceutics, School of Pharmacy, Ahvaz Jundishapur University of Medical Sciences, Ahvaz, Iran

Full list of author information is available at the end of the article



Liposomes were first used as topical therapy by Mezei and Gulasekharan in 1980. The introduction of liposomes as skin drug delivery systems, initially promoted for localized effects with minimal systemic delivery. Reduction of vesicle size improves drug deposition into deeper strata. Recent advances and alteration in the composition and structure of vesicles result in vesicles with tailored properties [9].

The first generations of elastic vesicles are deformable liposomes (Transfersomes®) which consist of phospholipids and surfactant as an edge activator. Surfactant destabilizes lipid bilayers and increases deformability of the vesicles. Ethosome is another type of elastic vesicle which is composed of phospholipid, ethanol and water [10]. Recently, a novel family of liposomes which named the Penetration Enhancer-containing Vesicles (PEVs) is described for enhanced (trans)-dermal drug delivery [11]. They mainly consist of phospholipids and penetration enhancers with hydrosoluble glycols such as diethylene glycol monoethyl ether or propylene glycol. Transient reduction of the stratum corneum barrier function and improvement in vesicular bilayer fluidity are two main functions of the PEVs [12].

The aim of this study was to prepare and evaluate deformable liposome using soy phosphatidylcholine and transcutool (Diethylene glycol monoethyl ether) for topical delivery of tretinoin. Also optimization of formulation was performed in order to obtain suitable dermal delivery system for tretinoin.

Methods

Materials

Tretinoin was purchased from Sepidaj (Iran). Soy phosphatidylcholine (phospholipon 85G®) was obtained from lipoid (Germany). Cholesterol and HEPES (4-(2-Hydroxyethyl) 1-piperazine ethanesulfonic acid) were purchased from Sigma (Germany). Diethylene glycol monoethyl ether (Transcutol®) received as a gift from Gattefosse (France). Propyl paraben, methyl paraben, propylene glycol and vitamin E were obtained from Merck (Germany). Tretinoin cream 0.05 % was purchased from the pharmacy and its base cream was provided as a gift by Iran Daru (Iran).

Animals

Female NMRI mice 7–9 weeks old were obtained from laboratory Animals Care and Breeding Center of Ahvaz Jundishapur University of Medical Sciences (Ahvaz, Iran). The experiments were conducted in full compliance with regulatory principles of ethics committee of Ahvaz Jundishapur University of Medical Sciences.

Experimental design

A full factorial 3² design was used for optimization procedure. The studied independent variables were amount of soy phosphatidylcholine (SPC) (X₁) and amount of transcutool (X₂) in formulation. Types and levels of the independent variables are listed in Table 1. Dependent variables (responses) were percent of incorporation efficiency (Y₁), percent of drug release in 15 min (Y₂) and percent of drug release in 24 h (Y₃). The resulted formulations are illustrated in Table 2.

Preparation of deformable liposomes

Deformable liposomes were prepared by the fusion method. Briefly, the lipid components consisted of SPC (15–20–25 % wt), transcutool (5–10–15 % wt), tretinoin) 0.05 % wt), cholesterol (2 % wt), propylene glycol (3 % wt), vitamin E (0.3 % wt), methyl paraben (0.1 % wt), and propyl paraben (0.02 % wt); these components were melted at about 75 °C. HEPES buffer (10 mM, pH 5) was heated separately and was added up to 100 % to the previously heated melted lipids, and the mixture was homogenized with a homogenizer (Ultra-Turrax IKA T25) for 5 min at 12,000 rpm and allow it to cool down to room temperature [13].

Characterization of the liposomes

Particle size measurement

The particle size of the samples were measured in triplicate by laser light scattering (Scatterscope 1, Qudix, South Korea). Samples were diluted in HEPES buffer to a suitable concentration (0.2 g formulation in 1 ml HEPES buffer).

Incorporation efficiency

Incorporation efficiency of liposomes was determined indirectly. Certain amounts of liposomal dispersions were centrifuged (VS-35SMTI, Korea) at 20,000 rpm for 25 min at 25 °C. The supernatant was collected and analyzed at 362 nm using UV spectrophotometer (Biowave II, Biochrom, England) [14]. The percent of incorporation efficiency of drug was calculated by the following formula:

Table 1 Independent variables: Types and Levels

Variables	Levels		
X ₁ : amount of SPC (% wt)	15	20	25
X ₂ : amount of transcutool (% wt)	5	10	15

Table 2 Composition of experimental formulations (runs)

Run	X ₁ : SPC (% wt)	X ₂ : transcutool (% wt)
1	15	5
2	15	10
3	15	15
4	20	5
5	20	10
6	20	15
7	25	5
8	25	10
9	25	15

$$EE\% = \left[\frac{\left(\begin{array}{l} \text{amount of initial drug} \\ - \text{amount of free drug in supernatant} \end{array} \right)}{\text{amount of initial drug}} \right] \times 100$$

In vitro drug release

Drug release studies were performed using dialysis membrane method. Dialysis membranes were soaked before use in distilled water for 20 h. 1 g of formulation was placed in a dialysis membrane and both ends were closed. The membrane was float in a beaker containing 150 ml phosphate buffer (pH 7.4) and methanol (2:1 v/v); and stirred at 200 rpm at 37 °C. 1 ml of receiver medium was removed at 0.25, 0.5, 0.75, 1, 1.5, 2, 4, 6, 8 and 24 h and same volume of the fresh medium was replaced. The collected samples were analyzed for their tretinoin content. The derived concentration values were corrected by using the equation (1):

$$Mt(n) = Vr \times Cn + Vs \times \sum Cm \quad (1)$$

Where Mt(n) is the current cumulative mass of drug transported across the membrane at time t, n is the number (times) of sampling, Cn is the current concentration in the receiver medium, $\sum Cm$ is the summed total of the previously measured concentrations, Vr is the volume of the receiver medium, and Vs corresponds to the volume of the sample removed for analysis [15, 16].

Viscosity

Viscosity of selected formulations were measured by Brookfield viscometer (DV II + Pro, US) at 10 rpm and 25 °C using spindle 64.

In vitro skin penetration and retention

In vitro skin penetration studies were performed using jacketed Franz cells with a receiver medium of 25 ml

Phosphate buffer (pH 7.4) and methanol (2:1 v/v) at 37 °C and surface area of 4.84 cm². The dorsal skin of mouse was shaved with electric clippers one day before the experiment. A suitable size of full-thickness skin of mouse was cut and clamped between the donor and receiver compartment of Franz cell with the stratum corneum side facing upward. The skin samples were initially left in the Franz cells for 30 min in order to facilitate hydration. Subsequently, 1 g of the optimum formulations and tretinoin cream were placed onto the skin surface. 1 ml of receiver solution was removed at 0.25, 0.5, 0.75, 1, 1.5, 2, 4, 6 and 8 h and same volume of the fresh medium was replaced. The collected samples were analyzed for their tretinoin content. The derived concentration values were corrected according to equation (1).

For the determination of the amount of the drug retained in the skin, at the end of the experiment, the amount of the formulation remaining on the surface of the skin was collected and assayed for tretinoin. The amount of tretinoin retained in the skin was then calculated by subtracting the sum of the amount of tretinoin that remained on the surface and the amount of tretinoin that was released (penetrated through the skin) from the whole applied amount [13].

The cumulative amount of drug permeated was plotted against time. The steady-state permeation rate (J_{ss}) was calculated by divided the slop of the linear portion of the plot on the exposed surface area of the skin. Lag time was determined from the x-intercept of the linear portion of the plot [17].

Histological evaluation

For skin histological study the dorsal side of the mouse was shaved with electric clipper. The skin was cut and mounted between donor and receiver compartment of the jacketed Franz cell whit stratum corneum side facing upward. 1 g of optimum formulations and tretinoin cream were placed on the skin. After 38 h the excessive formulation and cream were removed and the skin cleaned with cotton soaked in a phosphate buffer solution (pH 7.4). The treated skins were fixed in 10 % formalin solution and embedded in paraffin wax. Then samples were cut in 5 μ m thick sections by using microtome and conventionally stained with haematoxylin and eosin (H&E). Finally the samples were examined by light microscope (Olympus, BH-2, Japan) [18].

Differential scanning calorimetry (DSC)

DSC thermograms of SPC, cholesterol, transcutool and tretinoin were recorded on a differential scanning calorimeter (Mettler Toledo, DSC-1, Switzerland). Thermograms of both blank and tretinoin loaded liposomes were recorded individually. Certain amount of sample

was placed in aluminum pan and scanned from 20 to 200 °C by scanning rate of 10 °C/min.

Stability study

Optimum formulations were stored at refrigerate temperature (4 °C) for 3 months. After 1 and 3 months, the particle size and drug incorporation efficiency of the formulations were measured. The results were compared with the initial size and drug incorporation efficiency of formulations [19].

Statistical analysis

All experiments were repeated three times and expressed as the mean \pm standard deviation. One way analysis of variance (ANOVA) followed by multiple comparisons Tukey test was used to substantiate statistical differences between groups. Results with $P < 0.05$ were considered to be significant.

The effects of independent variables (X) on the dependent variables (Y) were modeled using a polynomial second order equation as followed:

$$Y = c + b_1X_1 + b_2X_2 + b_3X_1^2 + b_4X_2^2 + b_5X_1X_2$$

The modeling was performed using SPSS (Version 16.0) with a backward, stepwise linear regression technique and significant expressions ($P < 0.05$) were selected for the final equations. Response surface plots and contour plots resulting from equations obtained by Statgraphics version Centurion XVI.

Results and discussion

In an attempt to obtain suitable formulation containing tretinoin for dermal delivery, deformable liposomes prepared with SPC and transcutool.

Transcutool (diethylene glycol monoethylether) is a non-toxic, biocompatible with skin, penetration enhancer which is miscible in polar and non polar solvents [20]. Effect of transcutool on the lipid organizational structure of human stratum corneum was evaluated by

Table 3 Particle size (nm) and Incorporation efficiency (%) of liposomal formulations (mean \pm SD, $n = 3$)

Formulation	Average size	Incorporation (%)
1	8.82 \pm 0.95	85.51 \pm 0.15
2	16.96 \pm 3.69	80.88 \pm 0.19
3	10.94 \pm 1.84	79.11 \pm 0.19
4	15.09 \pm 4.62	89.55 \pm 0.19
5	9.59 \pm 2.99	91.33 \pm 0.57
6	12.8 \pm 1.65	89.88 \pm 0.19
7	10.21 \pm 1.85	93.22 \pm 0.69
8	7.70 \pm 2.30	93.88 \pm 0.19
9	7.12 \pm 0.74	89.99 \pm 0.33

Moghadam et al. [21]. Their results of x-ray scattering study showed transcutool caused a slight disordering effect in the stratum corneum membrane and increasing its fluidity.

In this study the fusion method was used to prepare the topical liposomal formulations. This method is simple, efficient, and reproducible. The method is free of organic solvents like chloroform; and yields homogeneous liposomes with high incorporation efficiencies. Furthermore, liposomes prepared by fusion method showed enough viscosity that they could be applied directly on the skin without the need for the liposomal formulation to be mixed with other bases [9].

Characterization of the liposomes

Particle size

The mean particle size of the liposomal formulations was shown in Table 3. ANOVA analysis showed statistical significant differences between F2 and F1, F8, F9 ($P < 0.05$) and also, F4 and F9 ($P < 0.05$). Different concentrations of phospholipid and transcutool showed no effect on particle size of liposomal formulations.

Incorporation efficiency

The incorporation efficiency of formulations was in range of 79 to 93 % (Table 3). F8 showed maximum

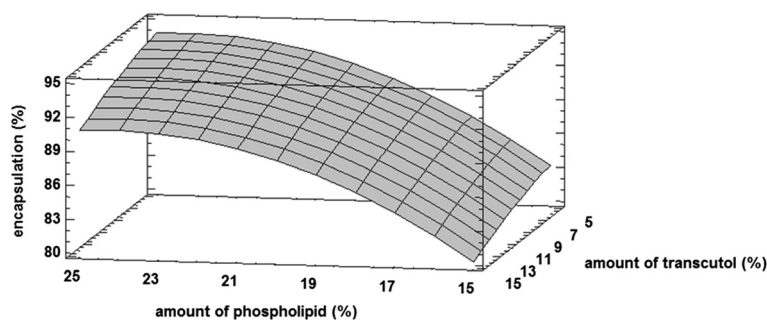


Fig. 1 Response surface plot for Y_1 response (percent of incorporation efficiency)

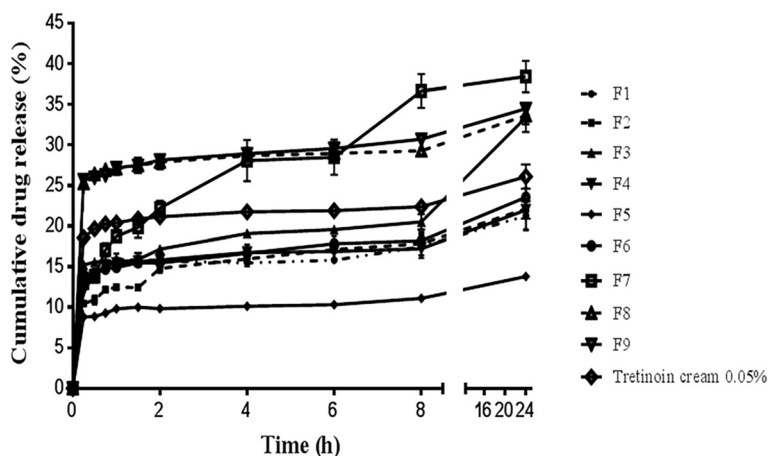


Fig. 2 Release profiles of tretinoin from various liposomal formulations and tretinoin cream

percent of incorporation. There were no statistical significant differences among F4, F6, F9 ($P > 0.05$) and also, between F7 and F8 ($P > 0.05$).

Mathematical relationships were generated between the responses and independent variables using the statistical package SPSS. The equations of the responses represent the quantitative effect of independent variables (X_1 and X_2) upon the responses (Y_1 , Y_2 and Y_3). Coefficients with more than one factor represent the interaction between factors while coefficients with second order terms indicate the quadratic nature of the phenomena.

The equation of the Y_1 (percent of incorporation efficiency) is given below:

$$Y_1 = 20.560 + 6.101 X_1 - 0.126 X_1^2 - 0.016 X_2^2 \quad (2)$$

Three-dimensional response surface plot for Y_1 is shown in Fig. 1. According to the Fig. 1, incorporation efficiency was increased by increasing the amount of phospholipid, while it was slightly decreased with higher amounts of transcucol. The lipophilic nature of tretinoin may lead to incorporation of this drug between the lipid

Table 4 Analysis of variance (ANOVA) of dependent variables (percent of incorporation efficiency (Y_1) and percent of drug release in 15 min (Y_2) and 24 h (Y_3))

Source of variation	Sum of squares	Degree of freedom	Mean square	F ratio	P-value
Y_1					
Regression	604.968	3	201.656	87.409	0.000
Residuals	53.062	23	2.307		
Total	658.030	26			
$R^2 = 0.919$					
Y_2					
Regression	716.867	4	179.217	23.780	0.000
Residuals	165.800	22	7.536		
Total	882.666	26			
$R^2 = 0.812$					
Y_3					
Regression	1569.312	4	392.328	87.047	0.000
Residuals	99.156	22	4.507		
Total	1668.468	26			
$R^2 = 0.941$					

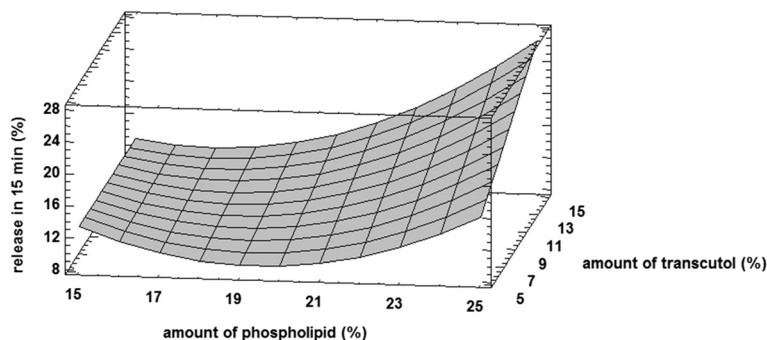


Fig. 3 Response surface plot for Y_2 Response (percent of drug release in 15 min)

bilayers. This can also explain the high incorporation efficiency of the tretinoin in liposomal formulations [22].

In vitro drug release

The release profile of tretinoin from various liposomal formulations and conventional tretinoin cream are shown in Fig. 2. In the first 2 h, drug release from F8 and F9 were significantly ($P < 0.05$) higher than other formulations, at the 4th hour no statistical significant differences were observed among F7, F8 and F9 ($P > 0.05$) and after 8 and 24 h, F7 showed maximum drug release compared with other formulations. The release pattern of tretinoin cream and all liposomal formulations except F7, showed burst release in initial times and then had a reduced rate of release [23].

The target of optimization was to obtain formulations with slow release in first 15 min and maximum release at 24 h. Therefore constraints for the Y_2 (percent of drug release in 15 min) and Y_3 (percent of drug release in 24 h) were:

$$Y_2 < 15 \% \text{ and } Y_3 > 30 \%$$

The equations of the responses Y_2 and Y_3 are given below:

$$Y_2 = 101.07 - 9.207 X_1 - 1.666 X_2 + 0.111 X_1 X_2 + 0.223 X_1^2 \tag{3}$$

$$Y_3 = 149.821 - 14.491 X_1 - 0.177 X_1 X_2 + 0.431 X_1^2 + 0.194 X_2^2 \tag{4}$$

Analysis of variance (ANOVA) (Table 4) demonstrated that the mathematical models generated were statistically significant and valid for each considered response.

Three-dimensional response surface plot for Y_2 and Y_3 are shown in Figs. 3 and 4. As shown in Fig. 3, drug release was decreased in 15 min by increase in the concentrations of phospholipids up to 20 %. Meanwhile, increase in the amount of phospholipid from 20 to 25 %, higher the value of drug release. Also it was shown that formulations with higher amounts of transcucol in their compositions released more elements of drug at 15 min.

According to Fig. 4, liposomes containing about 17–19 % phospholipid exhibited the lowest drug release among the formulations. However, it was increased by either lower or higher phospholipid concentrations so that drug release at 24 h from formulations composed of 25 % phospholipid (F7, F8 and F9) was maximums.

The phase transition temperature (T_m) of the lipids affected the release of liposome [24]. Phospholipids at their T_m changed from solid to liquid state. Thus the

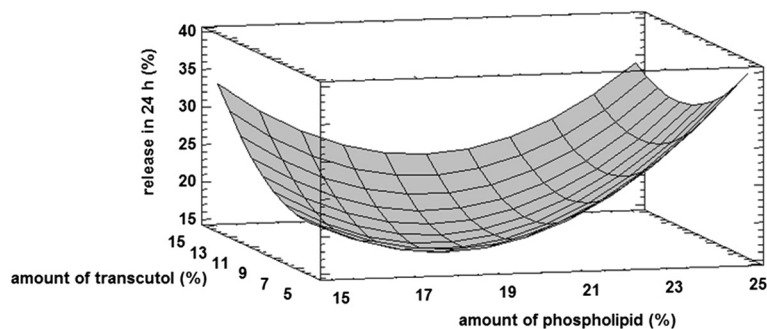
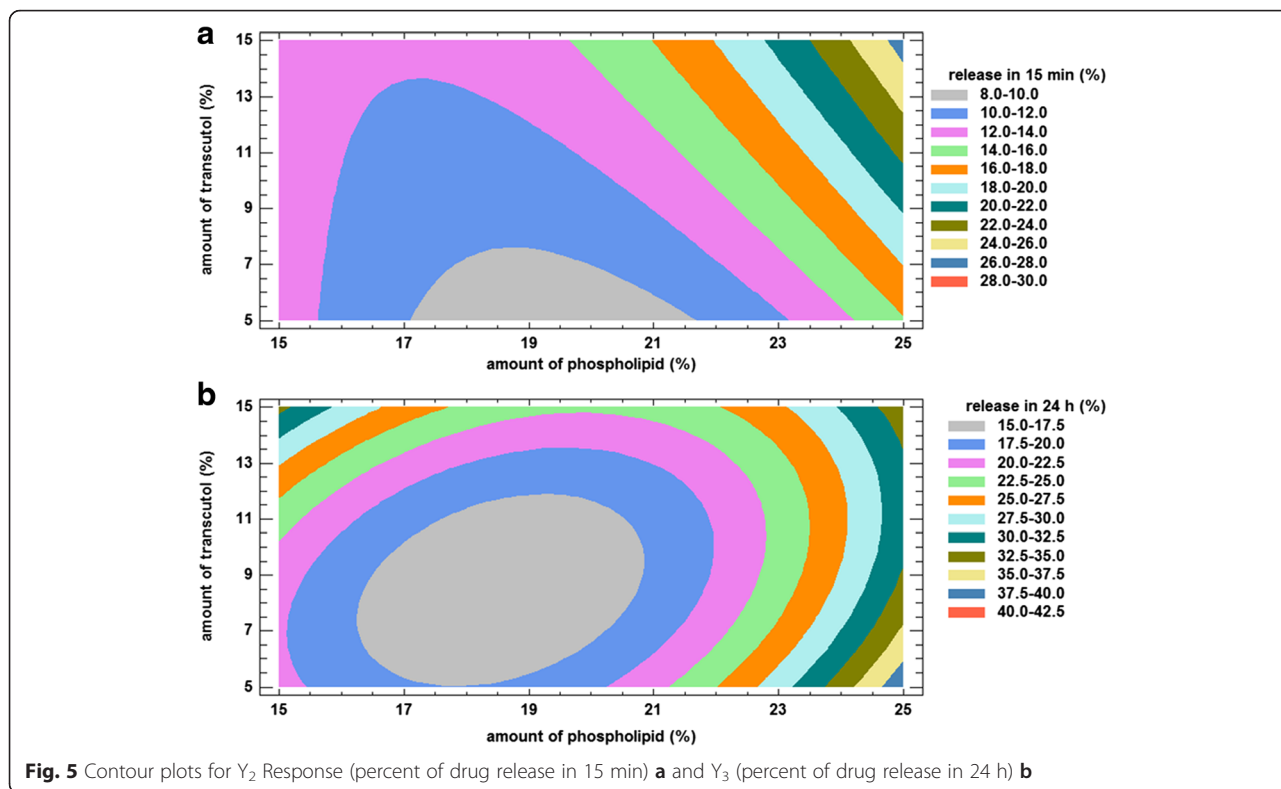


Fig. 4 Response surface plot for Y_3 Response (percent of drug release in 24 h)



permeability of liposomal membrane is increased and encapsulated drug released [25]. Chen et al. [26] prepared stealth liposome with different phosphatidylcholine and found liposome composed SPC in the presence of rat plasma showed maximum drug release which due to lower T_m of SPC.

According to contour plots of responses Y_2 and Y_3 (Fig. 5), optimized formulations were selected. The defined desirable areas of responses Y_2 and Y_3 were superimposed and the region of interest was found. The ratio of SPC: transcutol to obtained optimum formulations were 15.5:14.5, 24:7 and 25:5.

As a validation method for the process, liposomal formulations were prepared at the predicted levels of the independent variables and evaluated for percent of drug release in 15 min and 24 h. Observed and predicted responses for optimum formulations were then compared and the results were shown in Table 5.

Viscosity of F (25:5) and F (15.5:14.5) were 354.53 ± 4.51 and 266.94 ± 3.17 poise, respectively. Formulation (25:5) with the highest amount of phospholipid and lowest amount of transcutol had more viscosity compared with F (15.5:14.5). Thus Formulation (25:5) showed slower release in initial times. Fetih et al. [27] developed and evaluated liposomal gels of celecoxib and concluded that inverse relationship presented between viscosity of liposomal gels with drug diffusion rate and percent of drug released.

According to the results presented in Table 5, observed responses were close to predicted ones which confirmed that the factorial design was valid for predicting the optimum formulation.

In vitro skin penetration and retention

The skin penetration profile of F (15.5:14.5), F (24:7), F (25:5) and conventional tretinoin cream are shown in

Table 5 Observed and predicted responses (percent of drug release in 15 min (Y_2) and 24 h (Y_3)) for optimum formulations

Formulation (SPC: transcutol)	Dependent variables	Observed response	Predicted response	Residual
F (15.5:14.5)	Y_2	14.88	12.72	2.61
	Y_3	30.41	29.76	0.65
F (24:7)	Y_2	14.88	15.53	-0.65
	Y_3	33.28	30.06	3.22
F (25:5)	Y_2	12.97	15.81	-2.84
	Y_3	38.43	39.64	-1.21

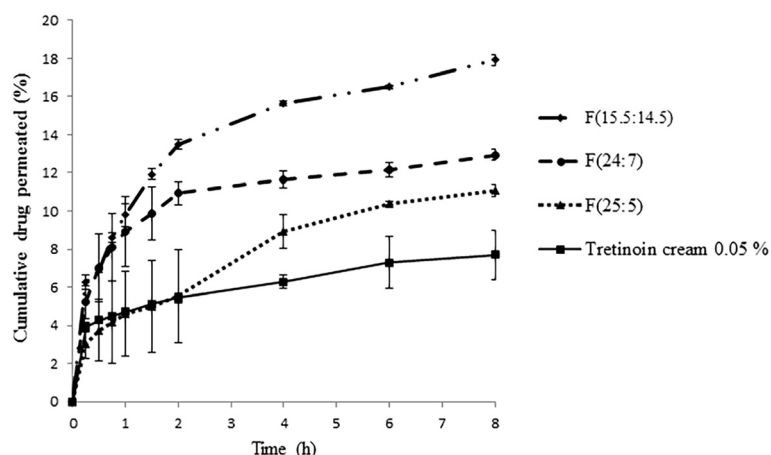


Fig. 6 In vitro permeation profiles of tretinoin from cream and optimum formulations through mice skin

Fig. 6. The penetration percent of optimum formulations was higher ($P < 0.05$) than tretinoin cream. F (15.5:14.5) showed maximum percent of penetration.

According to the Table 6, F (15.5:14.5) showed maximum flux. Formulations and cream penetrated through the skin without any lag time. The flux of F (15.5:14.5) and F (24:7) were higher than cream ($P < 0.05$), while no statistical significant differences found between F (25:5) and tretinoin cream ($P > 0.05$).

ANOVA analysis showed statistical significant differences between J_{ss} of optimum Formulations ($P < 0.05$). These results showed that by enhancement of transcutol concentration in the optimum formulations, percent of penetration and flux was increased which can be due to solubilizing properties of transcutol [28].

The percent of retained tretinoin and ratio of retained drug in the skin to penetrated drug for optimum formulations and tretinoin cream calculated and results showed in Table 6. There was no significant difference in the percent of retained tretinoin between optimum formulations and tretinoin cream ($P > 0.05$). ANOVA analysis showed that no significant difference among ratio of retention/penetration of optimum formulations ($P > 0.05$) but tretinoin cream showed higher ratio of retention/penetration than F(15.5:14.5) and F(24:7) ($P < 0.05$) that could be attributed to higher flux in formulations containing higher amount of transcutol.

Manconi et al. [29] evaluated PEVs with different penetration enhancers for dermal delivery of tretinoin. Their results showed PEVs improved cutaneous drug accumulation compared to control liposome. However, PEVs containing transcutol and labrasol had lower drug accumulated/drug permeated ratio because these vesicles increased both tretinoin deposition and flux.

Mura et al. [30] prepared PEVs with labrasol, transcutol and cineole for (trans) dermal delivery of minoxidil. They concluded that PEVs improved drug deposition into the skin when compared to the classic liposomes.

Caddeo et al. [31] developed PEVs with transcutol or propylene glycol, liposomes and ethosomes for delivered the diclofenac to the skin. Results showed PEVs containing transcutol led to higher drug accumulation in the skin compared to other carriers, diclofenac solution and voltaren. Also they found PEVs containing transcutol permeated the drug more than other vesicular formulations.

Histological evaluation

Histopathological investigation revealed that the thickness of epidermis in skin samples treated with F (15.5:14.5) (Fig. 7b) and F (25:5) (Fig. 7c) was same as untreated skin (control group) (Fig. 7a) and hyperkeratosis was increased in comparison with control group. In tretinoin cream treated skin, proliferations of keratinocytes were obvious and also hyperkeratosis was evident (Fig. 7d). Similar results obtained by Ascenso et al. [32].

Table 6 Permeation parameters, Retention (%) and ratio of retention/penetration of tretinoin from optimum formulations and cream (mean \pm SD, n = 3)

Formulation	J_{ss} ($\mu\text{g}/\text{cm}^2\cdot\text{h}$)	T_{lag} (h)	Retention (%)	Retention/Penetration
F(15.5:14.5)	4.71 ± 1	0	12.97 ± 2	0.72 ± 0.1
F(24:7)	3.29 ± 0.08	0	8.27 ± 1.91	0.63 ± 0.15
F(25:5)	1.05 ± 0.25	0	13.19 ± 0.71	1.18 ± 0.07
Tretinoin cream	0.88 ± 0.04	0	13.15 ± 2.77	1.72 ± 0.44

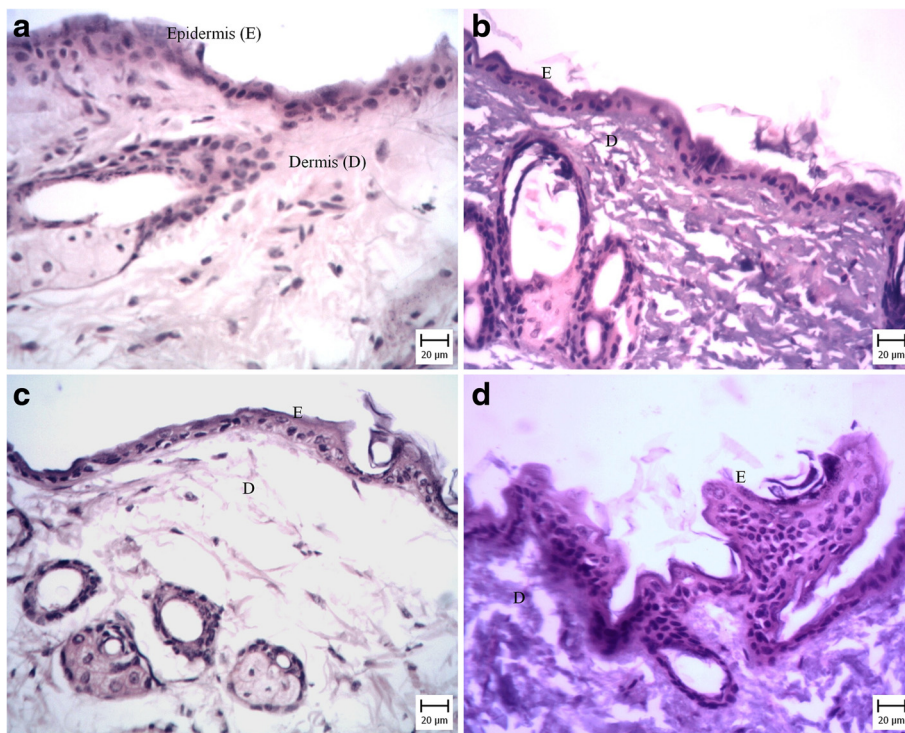


Fig. 7 Photomicrographs (20x) of hairless mice skin sections after haematoxylin and eosin staining: **a** untreated skin, **b** skin treated with F(15.5:14.5), **c** skin treated with F(25:5) and **d** skin treated with tretinoin cream 0.05 %

They evaluated in vivo skin irritation potential of tretinoin loaded ultradeformable vesicles in comparison to ketrel® and demonstrated ultradeformable vesicles caused lowest skin irritation. Their histological study showed in the skin treated with tretinoin loaded ultradeformable vesicle, hyperkeratosis was occurred and for ketrel® treated skin hyperplasia was observed.

Raza et al. [1] developed lipid-based nanocarriers for dermal delivery of tretinoin and concluded these carriers were well-tolerated on mouse skin while in the skin

treated with marketed product inflammation was observed.

Castro et al. [33] prepared retinoic acid loaded solid lipid nanoparticles and found this carriers considerably reduced skin irritation in rabbit and rhino mouse models when compared to marketed cream. Furthermore their histological evaluation showed in skin treated with retinoic acid loaded solid lipid nanoparticles and marketed gel epidermal thickness was increased in comparison to placebo group.

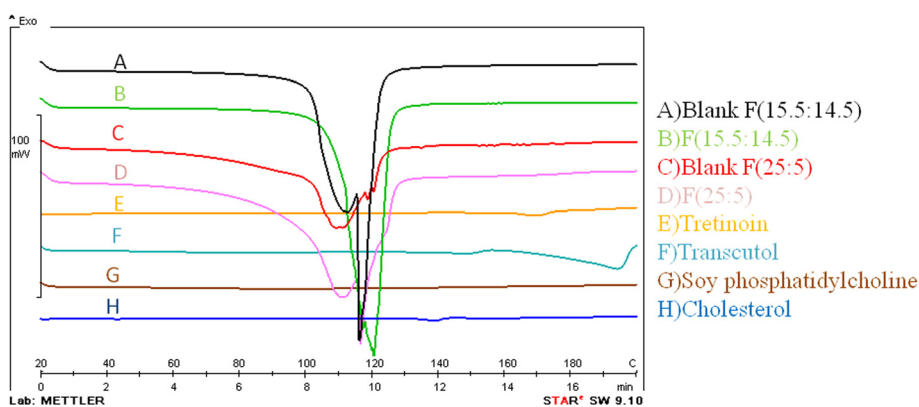


Fig. 8 DSC thermograms of **A** Blank F(15.5:14.5), **B** F(15.5:14.5), **C** Blank F(25:5), **D** F(25:5), **E** Tretinoin, **F** Transcutol, **G** Soy phosphatidylcholine, **H** Cholesterol

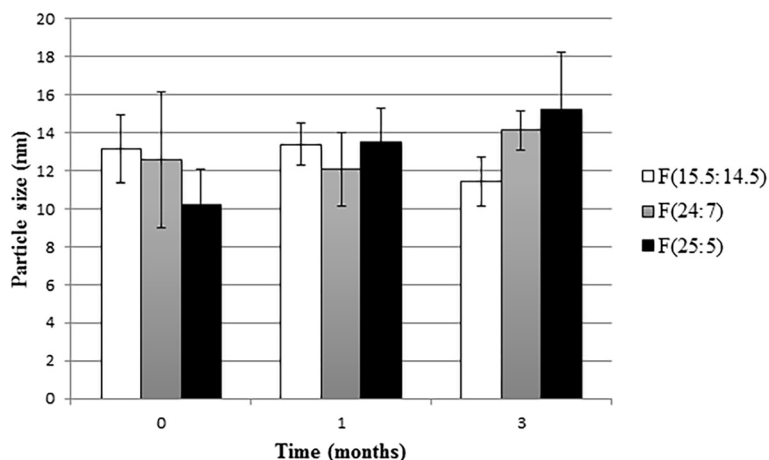


Fig. 9 Mean particle size (nm) of optimum formulations after 1 and 3 months storage at 4 °C

DSC

According to Fig. 8, DSC thermogram of tretinoin and cholesterol showed endothermic peak at 170 °C and 140 °C corresponding to their melting points, respectively [34, 35]. Transcutol thermogram showed endothermic peak about 195 °C, indicating its boiling point [36]. SPC thermogram showed a broad peak at 88 °C. In case of tretinoin loaded liposomes, the melting point of tretinoin was not observed which indicates that it is encapsulated in the liposome. The blank and tretinoin loaded liposome showed peak around 100–120 °C in their thermograms, which may be caused by evaporation of bounded water.

Stability study

The results of stability study for optimum formulations are shown in Figs. 9 and 10. Over the course of 3 months, the mean particle size of formulations did not change significantly ($P > 0.05$). This matter could be

resulted by the presence of transcutol in the formulation that may cause flexibility in vesicles [30] and cholesterol that has stabilizing effect against aggregation and fusion of the liposomes [37]. Similar results obtained by Mura et al. [38] that found particle size of PEVs with different amounts of transcutol remained constant during 3 months at 4 °C. While, particle size of liposomes without transcutol increased significantly. Incorporation efficiency of F (24:7) and F (25:5) were significantly decreased ($P < 0.05$) after 1 and 3 months while drug loading of F (15.5:14.5) only decreased about 6 % after 3 months ($P < 0.05$).

Chessa et al. [39] evaluated influence of PEVs with different types of penetration enhancer on (trans) dermal delivery of quercetin and concluded all PEVs promoted drug deposition in the skin.

Srisuk et al. [40] prepared deformable liposomes containing oleic acid for transepidermal delivery of

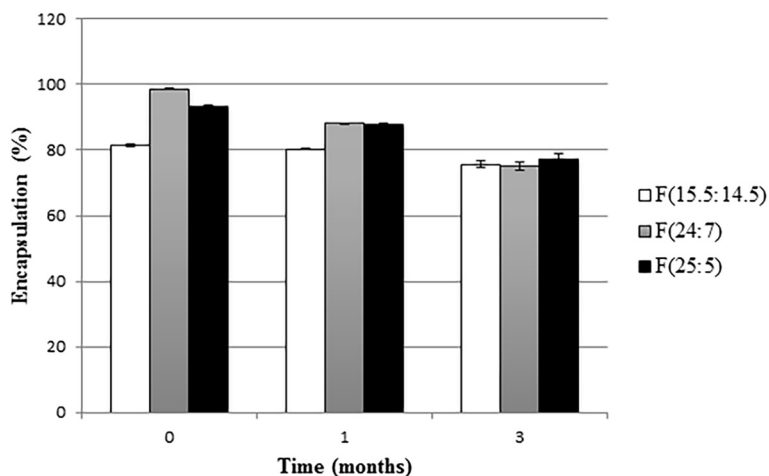


Fig. 10 Incorporation efficiency (%) of optimum formulations after 1 and 3 months storage at 4 °C

methotrexate. These vesicles showed the highest skin permeation, accumulation and flux compared to liposomes without oleic acid and methotrexate solution.

Charoenputtakun et al. [41] developed solid lipid nanoparticles, nanostructured lipid carriers and nanoemulsions containing limonene or cineol as penetration enhancer for dermal delivery of all-trans retinoic acid. They demonstrated solid lipid nanoparticles containing limonene had the highest skin permeation and flux in comparison to other formulations and all-trans retinoic acid suspension.

Conclusion

PEVs are novel class of liposomes which consist of phospholipid and penetration enhancer. In this work deformable liposome containing SPC and transcucol was employed for dermal delivery of tretinoin. Particle size of all formulations was smaller than 20 nm. Liposomes showed high incorporation efficiency which may be due to lipophilic nature of tretinoin. Formulations containing 25 % phospholipid exhibited the highest drug release at 24 h while the amount of transcucol did not significantly change drug release. Drug penetrated through the skin for optimum formulations was higher than cream which can be due to solubilizing properties of transcucol. Optimum formulations compared to tretinoin cream caused milder hyperkeratosis without hyperplasia. These results suggested that deformable liposomes could gradually released tretinoin and thus decreased its adverse effects such as erythema, peeling, burning and also increased patient compliance.

Abbreviations

DSC: Differential Scanning Calorimetry; Jss: Steady-state permeation rate; PEV: Penetration-Enhancer Vesicle; SPC: Soy Phosphatidylcholine.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

Conception and Design: NB, AA, AS, AR. Acquisition of data: SS, AR. Analysis and interpretation of data: AA, NB, SS, AR. Drafting manuscript: NB, AA, SS. All authors read and approved the final manuscript.

Acknowledgments

This paper is issued from Pharm D thesis of Somayeh Seifmanesh and financial support was provided by a grant (N-32) from Nanotechnology Research Center of Ahvaz Jundishapur University of Medical Sciences, Ahvaz, Iran. The authors are thankful to Iran Daru for providing us base cream of tretinoin as a gift.

Author details

¹Nanotechnology Research Center, Ahvaz Jundishapur University of Medical Sciences, Ahvaz, Iran. ²Department of Pharmaceutics, School of Pharmacy, Ahvaz Jundishapur University of Medical Sciences, Ahvaz, Iran. ³Targeted Drug Delivery Research Center, School of Pharmacy, Mashhad University of Medical Sciences, Mashhad, Iran. ⁴Department of Pathobiology, Faculty of Veterinary Medicine, Shahid Chamran University of Ahvaz, Ahvaz, Iran.

Received: 17 August 2015 Accepted: 5 February 2016

Published online: 29 February 2016

References

- Raza K, Singh B, Lohan S, Sharma G, Negi P, Yachha Y, Katara, O. P. Nano-lipoidal carriers of tretinoin with enhanced percutaneous absorption, photostability, biocompatibility and anti-psoriatic activity. *Int J Pharm.* 2013; 456:65–72. doi:10.1016/j.ijpharm.2013.08.019.
- Brisaert M, Gabriels M, Matthijs V, Plaizier-Vercammen J. Liposomes with tretinoin: a physical and chemical evaluation. *J Pharm Biomed Anal.* 2001;26: 909–17.
- Sinico C, Manconi M, Peppi M, Lai F, Valenti D, Fadda AM. Liposomes as carriers for dermal delivery of tretinoin: in vitro evaluation of drug permeation and vesicle-skin interaction. *J Control Release.* 2005;103:123–36. doi:10.1016/j.jconrel.2004.11.020.
- Laouini A, Jaafar Maalej C, Limayem Blouza I, Sfar S, Charcosset C, Fessi H. Preparation, Characterization and Applications of Liposomes: State of the Art. *J Colloid Sci Biotechnol.* 2012;1:147–68. doi:10.1166/jcsb.2012.1020.
- Shah KA, Date AA, Joshi MD, Patravale VB. Solid lipid nanoparticles (SLN) of tretinoin: potential in topical delivery. *Int J Pharm.* 2007;345:163–71. doi:10.1016/j.ijpharm.2007.05.061.
- loele G, Cione E, Risoli A, Genchi G, Ragno G. Accelerated photostability study of tretinoin and isotretinoin in liposome formulations. *Int J Pharm.* 2005;293:251–60. doi:10.1016/j.ijpharm.2005.01.012.
- Manconi M, Sinico C, Valenti D, Lai F, Fadda AM. Niosomes as carriers for tretinoin. III. A study into the in vitro cutaneous delivery of vesicle-incorporated tretinoin. *Int J Pharm.* 2006;311:11–9. doi:10.1016/j.ijpharm.2005.11.045.
- Anadolu RY, Sen T, Tarimci N, Birol A, Erdem C. Improved efficacy and tolerability of retinoic acid in acne vulgaris: a new topical formulation with cyclodextrin complex. *J Eur Acad Dermatol Venereol.* 2004;18:416–21. doi:10.1111/j.1468-3083.2004.00929.x.
- Bavarsad N, Fazly Bazzaz BS, Khamesipour A, Jaafari MR. Colloidal, in vitro and in vivo anti-leishmanial properties of transfersomes containing paromomycin sulfate in susceptible BALB/c mice. *Acta Trop.* 2012;124:33–41. doi:10.1016/j.actatropica.2012.06.004.
- Elsayed MM, Abdallah OY, Naggar VF, Khalafallah NM. Deformable liposomes and ethosomes: mechanism of enhanced skin delivery. *Int J Pharm.* 2006;322:60–6. doi:10.1016/j.ijpharm.2006.05.027.
- Manconi M, Caddeo C, Sinico C, Valenti D, Mostallino MC, Biggio G, et al. Ex vivo skin delivery of diclofenac by transcucol containing liposomes and suggested mechanism of vesicle-skin interaction. *Eur J Pharm Biopharm.* 2011;78:27–35. doi:10.1016/j.ejpb.2010.12.010.
- Romero EL, Morilla MJ. Highly deformable and highly fluid vesicles as potential drug delivery systems: theoretical and practical considerations. *Int J Nanomedicine.* 2013;8:3171–86. doi:10.2147/ijn.s33048.
- Jaafari MR, Bavarsad N, Bazzaz BS, Samiei A, Soroush D, Ghorbani S, et al. Effect of topical liposomes containing paromomycin sulfate in the course of Leishmania major infection in susceptible BALB/c mice. *Antimicrob Agents Chemother.* 2009;53:2259–65. doi:10.1128/aac.01319-08.
- Kulkamp-Guerreiro IC, Berlitz SJ, Contri RV, Alves LR, Henrique EG, Barreiros VR, et al. Influence of nanoincorporation on the sensory properties of cosmetic formulations containing lipoic acid. *Int J Cosmet Sci.* 2013;35:105–11. doi:10.1111/ics.12013.
- Jafari B, Rafie F, Davaran S. Preparation and characterization of a novel smart polymeric hydrogel for drug delivery of insulin. *BiolImpacts.* 2011;1:135–43. doi:10.5681/bi.2011.018.
- Khan GM, Frum Y, Sarheed O, Eccleston GM, Meidan VM. Assessment of drug permeability distributions in two different model skins. *Int J Pharm.* 2005;303:81–7. doi:10.1016/j.ijpharm.2005.07.005.
- Sinko PJ, Singh, Y. Martin's physical pharmacy and pharmaceutical sciences. 6ed. Philadelphia; Lippincott Williams & Wilkins; 2006.
- Han SB, Kwon SS, Jeong YM, Yu ER, Park SN. Physical characterization and in vitro skin permeation of solid lipid nanoparticles for transdermal delivery of quercetin. *Int J Cosmet Sci.* 2014;36(6):588–97. doi:10.1111/ics.12160.
- Panwar P, Pandey B, Lakhera PC, Singh KP. Preparation, characterization, and in vitro release study of albendazole-encapsulated nanosize liposomes. *Int J Nanomedicine.* 2010;5:101–8.
- Mura P, Fucci MT, Bramanti G, Corti P. Evaluation of transcucol as a clonazepam transdermal permeation enhancer from hydrophilic gel formulations. *Eur J Pharm Sci.* 2000;9:365–72.
- Moghadam SH, Saliaj E, Wettig SD, Dong C, Ivanova MV, Huzil JT, et al. Effect of chemical permeation enhancers on stratum corneum barrier lipid

- organizational structure and interferon alpha permeability. *Mol Pharm*. 2013; 10:2248–60. doi:10.1021/mp300441c.
22. Awad RS, Wahed WS, Bitar Y. Evaluation the impact of preparation conditions and formulation on the accelerated stability of tretinoin loaded liposomes prepared by heating method. *Int J Pharm Pharm Sci*. 2015;7:171–8.
 23. Tabbakhian M, Sharifian A, Shatalebi MA. Preparation and in vitro characterization of tretinoin-containing microspheres suited for dermatological preparations. *Res Pharm Sci*. 2008;3:31–40.
 24. Aygun A, Torrey K, Kumar A, Stephenson LD. Investigation of factors affecting controlled release from photosensitive DMPC and DSPC liposomes. *Appl Biochem Biotechnol*. 2012;167(4):743–57. doi:10.1007/s12010-012-9724-6.
 25. Chen J, Cheng D, Li J, Wang Y, Guo JX, Chen ZP, et al. Influence of lipid composition on the phase transition temperature of liposomes composed of both DPPC and HSPC. *Drug Dev Ind Pharm*. 2013;39(2):197–204. doi:10.3109/03639045.2012.668912.
 26. Chen J, Yan GJ, Hu RR, Gu QW, Chen ML, Gu W, et al. Improved pharmacokinetics and reduced toxicity of brucine after incorporation into stealth liposomes: role of phosphatidylcholine. *Int J Nanomedicine*. 2012;7: 3567–77. doi:10.2147/ijn.s32860.
 27. Fetih G, Fathalla D, El-Badry M. Liposomal gels for site-specific, sustained delivery of celecoxib: in vitro and in vivo evaluation. *Drug Dev Res*. 2014; 75(4):257–66. doi:10.1002/ddr.21179.
 28. Prasanthi D, Lakshmi PK. Effect of chemical enhancers in transdermal permeation of alfuzosin hydrochloride. *ISRN Pharmaceutics*. 2012;2012:1–8. doi:10.5402/2012/965280.
 29. Manconi M, Sinico C, Caddeo C, Vila AO, Valenti D, Fadda AM. Penetration enhancer containing vesicles as carriers for dermal delivery of tretinoin. *Int J Pharm*. 2011;412:37–46. doi:10.1016/j.ijpharm.2011.03.068.
 30. Mura S, Manconi M, Sinico C, Valenti D, Fadda AM. Penetration enhancer-containing vesicles (PEVs) as carriers for cutaneous delivery of minoxidil. *Int J Pharm*. 2009;380:72–9. doi:10.1016/j.ijpharm.2009.06.040.
 31. Caddeo C, Sales OD, Valenti D, Sauri AR, Fadda AM, Manconi M. Inhibition of skin inflammation in mice by diclofenac in vesicular carriers: liposomes, ethosomes and PEVs. *Int J Pharm*. 2013;443:128–36. doi:10.1016/j.ijpharm. 2012.12.041.
 32. Ascenso A, Salgado A, Euleterio C, Praca FG, Bentley MV, Marques HC, et al. In vitro and in vivo topical delivery studies of tretinoin-loaded ultra-deformable vesicles. *Eur J Pharm Biopharm*. 2014;88:48–55. doi:10.1016/ j.ejpb.2014.05.002.
 33. Castro GA, Oliveira CA, Mahecha GA, Ferreira LA. Comedolytic effect and reduced skin irritation of a new formulation of all-trans retinoic acid-loaded solid lipid nanoparticles for topical treatment of acne. *Arch Dermatol Res*. 2011;303:513–20. doi:10.1007/s00403-011-1130-3.
 34. Zidan AS, Spinks C, Fortunak J, Habib M, Khan MA. Near-infrared investigations of novel anti-HIV tenofovir liposomes. *AAPS J*. 2010;12:202–14. doi:10.1208/s12248-010-9177-1.
 35. Ascenso A, Guedes R, Bernardino R, Diogo H, Carvalho FA, Santos NC, et al. Complexation and full characterization of the tretinoin and dimethyl-beta-cyclodextrin complex. *AAPS PharmSciTech*. 2011;12:553–63. doi:10.1208/ s12249-011-9612-3.
 36. Sullivan Jr DW, Gad SC, Julien M. A review of the nonclinical safety of Transcutol, a highly purified form of diethylene glycol monoethyl ether (DEGEE) used as a pharmaceutical excipient. *Food Chem Toxicol*. 2014;72: 40–50. doi:10.1016/j.fct.2014.06.028.
 37. Elmeshad AN, Mortazavi SM, Mozafari MR. Formulation and characterization of nanoliposomal 5-fluorouracil for cancer nanotherapy. *J Liposome Res*. 2014;24:1–9. doi:10.3109/08982104.2013.810644.
 38. Mura S, Manconi M, Valenti D, Sinico C, Vila AO, Fadda AM. Transcutol containing vesicles for topical delivery of minoxidil. *J Drug Target*. 2011;19: 189–96. doi:10.3109/1061186x.2010.483516.
 39. Chessa M, Caddeo C, Valenti D, Manconi M, Sinico C, Fadda AM. Effect of Penetration Enhancer Containing Vesicles on the Percutaneous Delivery of Quercetin through New Born Pig Skin. *Pharmaceutics*. 2011;3:497–509. doi:10.3390/pharmaceutics3030497.
 40. Srisuk P, Thongnopnua P, Raktanonchai U, Kanokpanont S. Physico-chemical characteristics of methotrexate-entrapped oleic acid-containing deformable liposomes for in vitro transepidermal delivery targeting psoriasis treatment. *Int J Pharm*. 2012;427:426–34. doi:10.1016/j.ijpharm.2012.01.045.
 41. Charoenputtakun P, Pamornpathomkul B, Opanasopit P, Rojanarata T, Ngawhirunpat T. Terpene composited lipid nanoparticles for enhanced dermal delivery of all-trans-retinoic acids. *Biol Pharm Bull*. 2014;37:1139–48.

Submit your next manuscript to BioMed Central and we will help you at every step:

- We accept pre-submission inquiries
- Our selector tool helps you to find the most relevant journal
- We provide round the clock customer support
- Convenient online submission
- Thorough peer review
- Inclusion in PubMed and all major indexing services
- Maximum visibility for your research

Submit your manuscript at
www.biomedcentral.com/submit

