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Eutectic-based liposome as a potential delivery system of paeonol

 Jieyu Wu, †^a Suqin Yang, †^b Tianxiang Yin *^a and Xiaoyong Wang ^a

Deep eutectic solvents (DESSs), as a new type of green solvent, show great advantages of easy preparation and no need of purification after synthesis and have great potential applications in various fields. Moreover, when combining active pharmaceutical ingredients (APIs) with DESSs, a new type of drug delivery system called a therapeutic deep eutectic solvent (THEDES) can be developed to increase the solubility of the APIs and/or their permeation ability as well as bioactivity. Herein, we have designed a new THEDES based on paeonol and lauric acid with a molar ratio of 1 : 1. The physicochemical properties as well as the bioactivity were investigated by a series of techniques. The THEDES showed better solubility in water and similar bioactivities as compared to the pure APIs. Besides, we further prepared a THEDES liposome, which possessed an average size of about 120 nm with an encapsulation efficiency of 84%. The obtained liposome showed good stability during the experimental period, which indicated a potential application for the delivery of paeonol. The results presented herein suggested that THEDES may be regarded as a new strategy for the effective delivery of APIs.

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1. Introduction

Deep eutectic solvents (DESSs), as a new type of solvent, can be simply obtained by gently heating a suitable mixture of a hydrogen-bond donor (HBD) component and a hydrogen-bond acceptor (HBA) component, which presents 100% atom economy and fits within green chemistry principles.^{1,2} They show great advantages of easy preparation and no need of purification after synthesis. Therefore, increasing interest in DESSs has been awakened due to their great potential applications in various fields, like extraction, catalysis, and gas absorption.^{3–11} Moreover, by incorporating active pharmaceutical ingredients (APIs) into DESSs, a new type of drug delivery system called a therapeutic deep eutectic solvent (THEDES) can be developed. In recent years, a series of THEDES have been reported to increase the solubility of APIs and/or their permeation ability as well as bioactivity, where the API was used as a counterpart of the DES or as a solute in the DES.^{12–22}

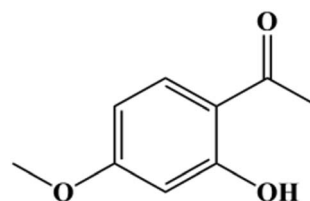
Paeonol(1-(2-hydroxy-4-methoxyphenyl) ethanone, Scheme 1), the main phenolic compound present in the traditional Chinese medicine herb *Paeonia suffruticosa*, shows great potential applications in pharmaceuticals due to its anti-inflammatory, antibacterial, and anti-tumor functions.^{23–25}

However, wide applications of paeonol are hindered by its poor solubility, *i.e.* 0.54 mg ml⁻¹,²⁶ and low oral bioavailability. Smart formulations and/or transdermal delivery with the help of penetration enhancers may be used to effectively deliver paeonol. Herein, we developed a novel eutectic system based on paeonol and lauric acid, and investigated the corresponding physicochemical properties and bioactivities. Liposomes have been extensively investigated as effective drug delivery systems due to their easily tunable physicochemical properties and good biocompatibility.^{27–29} Herein, a liposome based on the paeonol–lauric acid eutectic system was constructed to evaluate its potential application in paeonol delivery.

2. Experimental

2.1 Chemicals

Paeonol (≥98%), tyrosinase (≥500 U mg⁻¹), penicillin–streptomycin solution (100X), L-glutamine (≥99%), and cholesterol (≥99.5%) were purchased from Macklin. Lauric acid (≥99%)



Scheme 1 Chemical structure of paeonol.

^aSchool of Chemistry and Molecular Engineering, East China University of Science and Technology, Shanghai 200237, China. E-mail: yintx@ecust.edu.cn; Fax: +86 21 64250804; Tel: +86 21 64252012

^bThe Affiliated Huai'an Hospital of Xuzhou Medical University, The Second People's Hospital of Huai'an, No. 62, Huaihai Rd., Huai'an 223002, China

† These two authors contributed equally to this work.



Table 1 Compositions of reaction solutions in the tyrosinase inhibitory assay

No.	Volume (ml)				Tyrosinase
	PBS	DMSO	Drug ^a	L-DOPA	
1	1.6	0.2	0	1	0.2
2	1.8	0.2	0	1	0
3	1.6	0	0.2	1	0.2
4	1.8	0	0.2	1	0

^a Both pure ingredients (paeonol and lauric acid) and THEDES were dissolved in DMSO.

Table 2 Orthogonal experiment factors and levels of preparation of the liposomes

Level	Factors			
	A (mg ml ⁻¹)	B (w/w)	C (w/w)	D (°C)
1	5	10 : 1	1 : 10	40
2	8	8 : 1	1 : 8	50
3	10	5 : 1	1 : 5	60

was supplied by Alfa Aesar. Phosphate buffered saline (PBS, pH = 6.8, 0.01 M) was purchased from Biorigin. L-Dopa ($\geq 98\%$) was supplied by Duly Biotech. Co. Ltd (Nanjing, China). Lecithin (from soybean, $>90\%$) was purchased from Aladdin. DMSO (Greagent, $\geq 99\%$) and dichloromethane (Greagent, 99.5%) were from Tansoole (Titan Co. Shanghai, China). CCK-8 was from Invigentech. RPMI 1640 cell culture medium and fetal bovine serum (FBS) were supplied by Biological Industries.

2.2 Preparation and characterization of THEDES

2.2.1 Construction of the paeonol-lauric acid eutectic solvent. Mixtures of paeonol and lauric acid at different mole fractions were prepared and heated up to 50 °C. Thereafter, the samples were cooled slowly to 37 °C with the temperature being controlled by a water bath. The sample that still remained in the

liquid state was chosen as the eutectic system (denoted as THEDES) for the following investigations.

Differential scanning calorimetry experiments were carried out for paeonol, lauric acid and THEDES (mole ratio of paeonol and lauric acid being 1 : 1). The thermograms were recorded from 0 °C to 80 °C with a scanning rate of 5 °C min⁻¹.

2.2.2 Characterization of THEDES

2.2.2.1 ¹H NMR. A suitable amount of THEDES was dissolved in CDCl₃, and the ¹H NMR spectrum was recorded by a Bruker Avance 400 spectrometer.

2.2.2.2 Viscosity and solubility measurement of THEDES. The viscosity of THEDES at 37 °C was determined by an autoviscosimeter (NDJ-5S, Hengping Co. Shanghai, China).

2.2.2.3 Solubility. The method used to determine THEDES solubility was as follows: slightly excess amount of THEDES was added to 10 ml water. The mixture was then vigorously stirred at 37 °C for 48 h, and was further centrifuged at 8000 rpm for 45 min. Thereafter, the supernate was carefully sucked and analyzed by a UV-vis spectrometer with the help of a calibration curve. The calibration curves of paeonol and THEDES were determined in aqueous solution by the UV-vis spectrometer (UV-2450, Shimadzu), which were $A = 0.0848C/(\text{mg ml}^{-1})$ for paeonol and $A = 0.0399C/(\text{mg ml}^{-1})$ for THEDES, where A and C refer to the absorbance and concentration, respectively. The process was repeated thrice and the average value was recorded.

2.2.3 Tyrosinase inhibitory assay. L-DOPA was used as a substrate and the absorption intensity change of its oxidation product at 475 nm was monitored to determine the tyrosinase inhibition efficiency. The reaction was carried out in a PBS solution containing 0.5 mM L-DOPA and tyrosinase solution (500 U ml⁻¹) at 37 °C. The substances were added in the order shown in Table 1 (from left to right). The mixture was incubated for 15 min before the addition of tyrosinase.

The change of the absorbance A of DOPA chrome was measured at 475 nm using the UV-vis spectrometer. The tyrosinase inhibitory activity was then calculated by:

$$\text{Inhibitory Activity}\% = \frac{[(A_1 - A_2) - (A_3 - A_4)]}{(A_1 - A_2)} \times 100\% \quad (1)$$

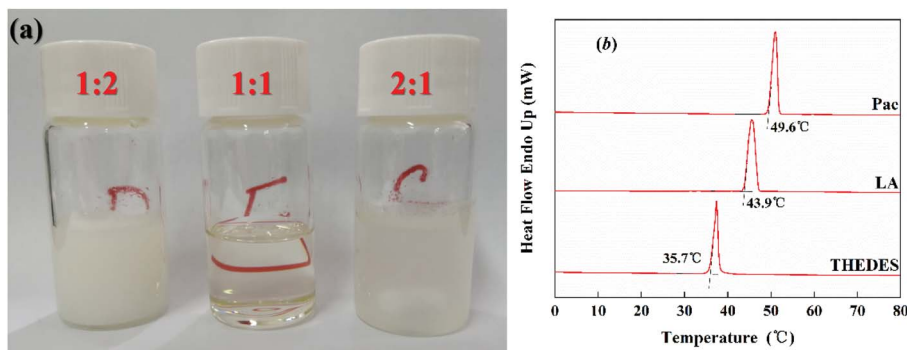


Fig. 1 (a) Images of the paeonol and lauric acid mixtures at different mole ratios (paeonol : lauric acid); (b) thermograms for the pure components paeonol (pae) and lauric acid (LA), and THEDES.



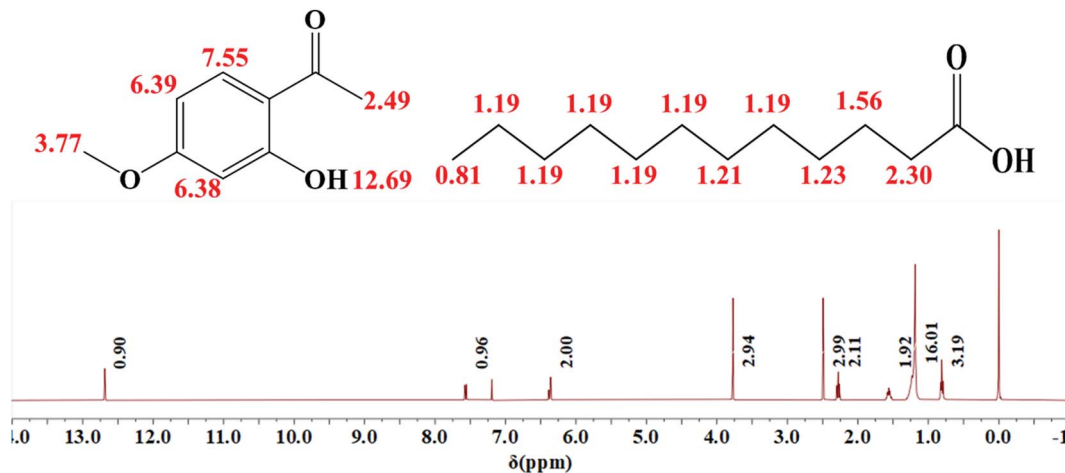


Fig. 2 ^1H NMR spectrum of THEDES.

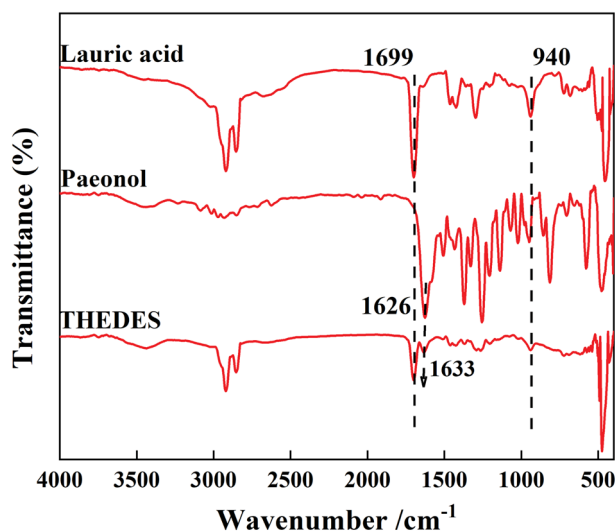


Fig. 3 FT-IR spectra of lauric acid, paeonol and THEDES.

The tyrosinase inhibitory assay was performed thrice and the average values were reported.

2.2.4 Cytotoxicity. The CCK-8 assay was used to evaluate cell viability to indicate the safety of THEDES. The L02 cell was cultured in RPMI 1640 supplemented with 10% FBS, 200 U ml^{-1} penicillin, 100 $\mu\text{g ml}^{-1}$ streptomycin and 0.3 g ml^{-1} L-glutamine under a humidified atmosphere of 95% CO_2 at 37 $^\circ\text{C}$. The cells were seeded at a density of approximately 10^4 cells per ml in a 96-well plate for 48 h. Thereafter, they were incubated with THEDES, paeonol, or lauric acid at a series of concentrations. The cytotoxicity was tested by the CCK-8 method, where the optical density was measured at 450 nm using an automatic microplate reader (SPARK 10M, TECAN). The cell viability was displayed as a percentage compared to the untreated cells. Three independent experiments were carried out, and the average values were taken and reported with standard deviation.

2.3 Preparation and characterization of the liposome

2.3.1 Preparation of the liposome. The liposome was prepared by a traditional thin film hydration method. Briefly, lecithin, cholesterol, and THEDES were dissolved in 20 ml dichloromethane with the help of ultrasound. Thereafter, the organic solvent was removed using a vacuum rotary evaporator. The thin lipid film formed on the wall of the flask was further hydrated with double distilled water for 1 h, and the mixture was further ultra-sounded for 5 min to give the liposome solution.

2.3.2 Determination of encapsulation efficiency (EE). The ultrafiltration centrifugation method was used to determine the encapsulation efficiency. Ethanol was added to a certain amount of liposome solution to break down the liposomes. The absorbance of the obtained solution at 274 nm was determined by a UV-vis spectrometer and denoted as A_0 . On the other hand, 0.4 ml liposome solution was added to an ultrafiltration centrifuge tube with a molecular weight cutoff of 10 KDa, which was centrifuged at a working speed of 8000 rpm for 40 min. The lower part of the solution was carefully taken, whose absorbance at 274 nm was measured by a UV-vis spectrometer and denoted as A_1 . Thus, the encapsulation efficiency was calculated by:

$$\text{EE} = \frac{A_0 - A_1}{A_0} \times 100\% \quad (2)$$

2.3.3 Optimization of the formula of the liposome. Four main factors that have shown significant impacts on the encapsulation efficiency, *i.e.* lecithin concentration (A), lecithin-to-cholesterol ratio (B), drug-to-lecithin ratio (C) and hydration temperature (D), were chosen to design a four-factor three-level orthogonal test with encapsulation efficiency as the evaluation index. The orthogonal test factors and levels of the preparation process of the liposomes are shown in Table 2.

2.3.4 Characterization of the liposome. The size distribution and zeta potential of the liposome were determined using a Malvern Zetasizer Nano ZS instrument (Southborough, MA)



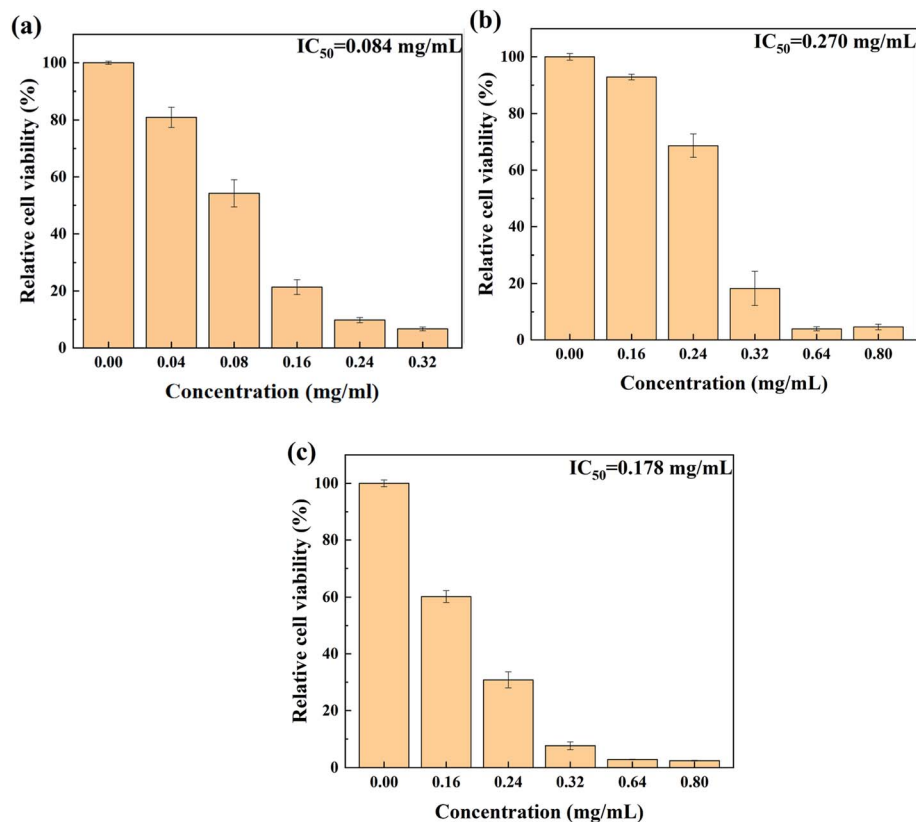


Fig. 4 Cytotoxicity of THEDES (a), paeonol (b) and lauric acid (c) using L02 cells.

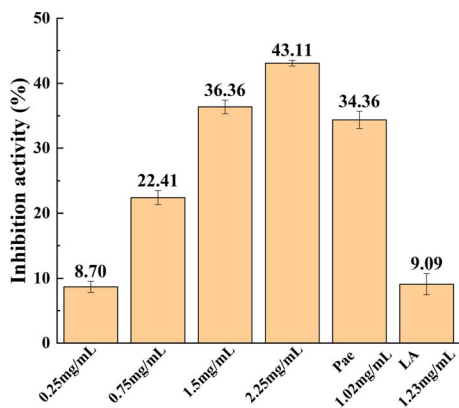


Fig. 5 Inhibition activities of THEDES at various concentrations and those of the pure components lauric acid (LA) and paeonol (pae).

with a backscattering detector (173.8°) and a laser with 633 nm wavelength. Measurements were taken in batch mode at 298.15 K using a plastic cuvette with a path length of 1 cm.

Negative stained transmission electron microscopy (NS-TEM) measurements were performed to show the morphology of the liposome. A drop of liposome solution was placed on a carbon-coated copper grid (300 meshes), and the excess was removed by filter paper. After that, a drop of 1% phosphotungstic acid solution was dropped on the copper grid as the

staining agent, and the excess liquid was removed after 15 s. The image of the sample after drying at room temperature was obtained using a JEM-1400 electron microscope at a working voltage of 80 kV.

3. Results and discussions

3.1 Preparation and physicochemical properties of THEDES

The images of paeonol and lauric acid at different mole ratios were obtained at 37°C and some of these samples are displayed in Fig. 1a. The eutectic system occurs when the mole ratio of paeonol and lauric acid is 1 : 1. Furthermore, the thermograms of the pure components and THEDES were collected and are displayed in Fig. 1b. The melting temperatures for paeonol and lauric acid as well as the eutectic temperature were determined to be 49.6°C , 43.9°C , and 35.7°C , respectively.

THEDES was verified by a ^1H NMR experiment, as displayed in Fig. 2. It can be estimated from Fig. 2 that the ratio of the peak area between $\delta = 12.69$ (–OH in paeonol) and $\delta = 0.8$ (m, – CH_3 in Lauric acid) is about 3; that is, the mole ratio of paeonol to lauric acid is nearly 1 : 1 in THEDES. The interaction between the –COOH group in lauric acid and the OH group in paeonol can be speculated from the FT-IR spectra, shown in Fig. 3, where the C=O stretching peak at 1626 cm^{-1} of paeonol moves towards a higher wavenumber from 1626 cm^{-1} to 1633 cm^{-1} . Moreover, the ratio of the absorbance between the –OH bending peak at 940 cm^{-1} of lauric acid and the C=O stretching peak at



Table 3 Results of the orthogonal test

No.	Factors				EE (%)
	A	B	C	D	
1	1	1	1	1	75.8
2	1	2	2	2	78.1
3	1	3	3	3	72.3
4	2	1	2	3	75.1
5	2	2	3	1	74.1
6	2	3	1	2	76.4
7	3	1	3	2	80.8
8	3	2	1	3	81.3
9	3	3	2	1	76.1
K_1	75.4	77.2	77.8	75.3	
K_2	75.2	77.8	76.5	78.4	
K_3	79.4	74.9	75.7	76.2	
R	4.21	2.90	2.10	3.11	

1699 cm^{-1} of lauric acid declined obviously, which further indicated the interaction between lauric acid and paeonol.

The solubility of THEDES in water was measured to be $1.67 \pm 0.02 \text{ mg ml}^{-1}$ (the corresponding amount of paeonol is 0.76 mg ml^{-1}). The solubility of paeonol in water was also measured herein to be 0.59 mg ml^{-1} , which was similar to the reported value (0.54 mg ml^{-1}).²⁶ Up to 30% increase in solubility for paeonol was achieved by the formation of THEDES, which was better than the previously reported eutectic system formed by paeonol and menthol (molar ratio of 4 : 6).³⁰ The viscosity of THEDES at a temperature of 37°C was determined to be $6.31 \text{ mPa}\cdot\text{s}$, which displayed a relatively low viscosity as compared to other commonly investigated DES systems.

To evaluate the safety of THEDES for its application, the cytotoxicity of THEDES was determined and compared to those of paeonol and lauric acid, which are displayed in Fig. 4. The IC_{50} values for paeonol, lauric acid, and THEDES were calculated to be 0.084 mg ml^{-1} , 0.270 mg ml^{-1} , and 0.178 mg ml^{-1} , respectively. For THEDES, the molar ratio of paeonol to lauric acid is 1 : 1, which gives a mass ratio of 1 : 1.2. For 0.178 mg ml^{-1} THEDES (IC_{50}), the corresponding concentrations of paeonol and lauric acid were 0.08 mg ml^{-1} and 0.098 mg ml^{-1} , respectively. As compared to the relative cell viability at these

concentrations for pure paeonol and lauric acid, it can be found that THEDES presented slightly lower cytotoxicity.

Moreover, in order to show the bioactivity of THEDES, the inhibitory activity on tyrosinase of THEDES was evaluated and compared to those of the pure components (paeonol and lauric acid). The results of the inhibitory activity tests are shown in Fig. 5, which shows that the tyrosinase inhibitory activity of THEDES increases with concentration. Moreover, the inhibition activity of THEDES with a concentration of 2.25 mg ml^{-1} is similar to the sum of those of paeonol and lauric acid at the same concentrations as those in THEDES, which indicates that THEDES retains the bioactivity of the pure APIs.

3.2 THEDES-based liposome

The preparation process of the THEDES-based liposome was optimized by a $L_9(3^4)$ orthogonal test. The results are displayed in Table 3.

From Table 3, it can be speculated that the order of the factors affecting the encapsulation efficiency of the THEDES liposome is as follows: lecithin concentration (A) > hydration temperature (D) > lecithin-to-cholesterol ratio (B) > drug-to-lecithin ratio (C). The optimized parameters for the preparation of the THEDES liposome were determined to be: lecithin concentration (10 mg ml^{-1}), hydration temperature (50°C), lecithin-to-cholesterol ratio (8 : 1), and drug-to-lecithin ratio (1 : 10). Under this condition, three different batches of liposome were prepared and the encapsulation efficiencies were measured to be 84.5%, 83.9%, and 84.1%, which showed good reproducibility. The liposome obtained in the optimized condition was thus used in the following studies.

The size distribution and morphology of the obtained liposome were determined by the methods of dynamic light scattering and TEM, respectively, which are displayed in Fig. 6a and b, respectively. An average size of $117.5 \pm 0.6 \text{ nm}$ with $\text{PDI} = 0.283$ was obtained by the dynamic light scattering measurement (Fig. 6a), which shows reasonable agreement with that observed by TEM (Fig. 6b).

The zeta potential was determined to be $-16.1 \pm 0.2 \text{ mV}$, which may result in good stability of the prepared liposome. This was further checked by a stability test, where the prepared liposome solution was stored at 4°C for three months. The size

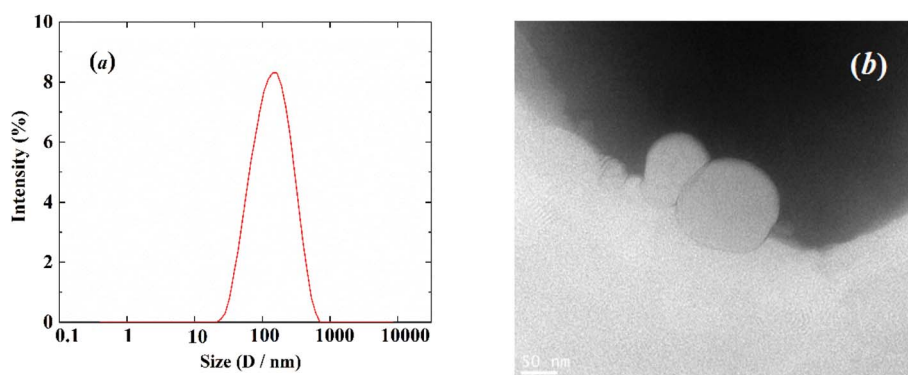


Fig. 6 (a) Size distribution of the liposome from dynamic light scattering measurement; (b) TEM image of the liposome.



of the liposome, zeta potential, and encapsulation efficiency (EE) were re-determined to be 124.2 ± 0.5 nm, -14.7 ± 0.3 mV and 81.7%, respectively; no obvious changes of these properties can be observed. These results indicated the good stability of the liposome and the great potential of this THEDES-based liposome as an efficient drug delivery system for paeonol.

4. Conclusion

In this work, we have designed a new THEDES based on paeonol and lauric acid with a molar ratio of 1 : 1. The physicochemical properties as well as the bioactivity were investigated by various methods. THEDES showed better solubility in water and similar inhibitory activity on tyrosinase as compared to the pure APIs. Moreover, THEDES displayed slightly weaker cytotoxicity than the pure components. We further prepared the THEDES liposome under optimized conditions of lecithin concentration (10 mg ml^{-1}), hydration temperature ($50 \text{ }^\circ\text{C}$), lecithin-to-cholesterol ratio (8 : 1), and drug-to-lecithin ratio (1 : 10). The obtained liposome possessed an average size of about 120 nm and an encapsulation efficiency of 84%. The liposome showed good stability during the experimental period of about 3 months. These results presented herein indicate the great potential of this THEDES-based liposome as an efficient drug delivery system for paeonol.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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