

Etoposide Amorphous Nanopowder for Improved Oral Bioavailability: Formulation Development, Optimization, in vitro and in vivo Evaluation

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Introduction: Etoposide refers to a derivative of podophyllotoxin, which plays an important role in the treatment of cancer due to its prominent anti-tumor effect. As a BCS IV drug, etoposide exhibits insufficient aqueous solubility and permeability, thereby limiting its oral absorption. To enhance the oral bioavailability of etoposide, this study developed an amorphous nanopowder.

Methods: Based on preliminary screening and experimental design, the stabilizer and preparation process of etoposide nanosuspension were explored. Subsequently, using a Box–Behnken design, the effects of independent factors (ultrasonication time, ratio of two phases and stabilizer concentration) on response variables (particle size and polydispersity index) were studied, and then the formulation was optimized. Finally, nanosuspension was further freeze dried with 1% of mannitol resulting in the formation of etoposide amorphous nanopowder.

Results: The optimized etoposide nanopowder showed as spherical particles with an average particle size and polydispersity index of 211.7 ± 10.4 nm and 0.125 ± 0.028 . X-ray powder diffraction and differential scanning calorimetry confirmed the ETO in the nanopowder was amorphous. Compared with coarse powder and physical mixture, etoposide nanopowder achieved significantly enhanced saturated solubility and dissolution in various pH environments. The C_{\max} and AUC_{0-t} of etoposide nanopowder after oral administration in rats were respectively 2.21 and 2.13 times higher than the crude etoposide suspension. Additionally, the T_{\max} value of nanopowder was 0.25 h, compared with 0.5 h of reference group.

Discussion: In the present study, the optimized amorphous nanopowder could significantly facilitate the dissolution and oral absorption of etoposide and might act as an effective delivery method to enhance its oral bioavailability.

Keywords: etoposide, amorphous nanopowder, oral absorption, poor aqueous solubility, bioavailability enhancement

Introduction

Etoposide (ETO) is a semisynthetic podophyllotoxin derivative, with the molecular formula of $C_{29}H_{32}O_{13}$ (Figure 1). It is capable of interfering with mammalian DNA topoisomerase II in the late S or G2 phase of the cell cycle, and combines with enzymes and DNA to fabricate a drug-enzyme-DNA stable reversible complex, which can irreparably impair DNA, eventually causing apoptosis.^{1,2} For its prominent anti-tumor effect, it has been applied for treating various tumors, such as

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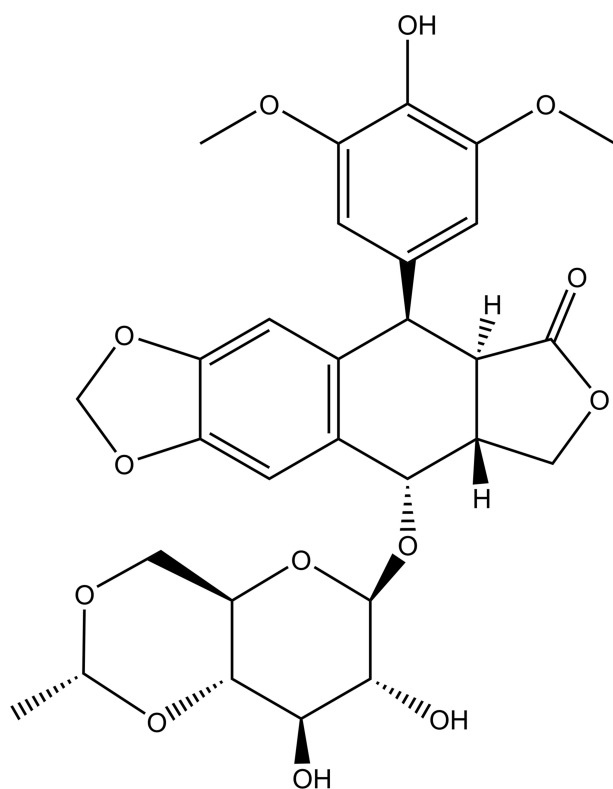


Figure 1 The chemical structure of ETO.

leukemia, lymphoma, small cell lung cancer and oropharyngeal cancer.^{3,4} However, as a BCS IV drug, etoposide exhibits insufficient aqueous solubility and permeability,^{5,6} resulting in many formulations of etoposide being prepared as injections.⁷⁻⁹ However, injections have many disadvantages, such as the need to add a large amount of surfactants and organic solvents,¹⁰⁻¹² which may cause allergic reactions in patients.¹³ Therefore, the development of oral preparations of etoposide has become an attractive research field. To facilitate the oral absorption of ETO, several strategies have been adopted, including nanostructured lipid carriers and self-emulsifying delivery systems. Some of them exhibit high *in vivo* performance.^{14,15}

Drug nanoparticles composed of drugs and stabilizers are considered one of the most promising nanotechnologies to promote insoluble drugs to exhibit enhanced solubility and oral bioavailability.¹⁶⁻¹⁸ With the decrease in particle size (PS) and the increase in specific surface area, drug nanoparticles can significantly elevate the saturation solubility and dissolution rate of drugs.¹⁹ Moreover, under the high drug loading capacity of drug nanoparticles, patients can have lower dosage and better compliance in oral administration.²⁰ The role of drug nanoparticles in improving solubility and oral bioavailability has been

confirmed in various literatures.²¹⁻²³ Drug nanoparticles may act as a preferable nanotechnology to facilitate the solubility and oral absorption of ETO, but this has rarely been studied. In this work, an ETO nanosuspension was prepared using a sonoprecipitation–high pressure homogenization method; subsequently, the formulation was optimized using a Box–Behnken design (BBD) of response surface method; lastly, it was transformed into ETO nanopowder by freeze-drying. The optimized ETO nanopowder was systematically analyzed in terms of morphology, particle size and physicochemical properties. Finally, *in vitro* dissolution and *in vivo* pharmacokinetics of optimized ETO nanopowder were assessed.

Materials and Methods

Materials

The following materials were provided by or purchased from the sources in parentheses: etoposide (Shanghai Xiandai Pharma Ltd, Shanghai, China); sodium dodecyl sulfate (SDS) and sodium deoxycholate (Tianjin Yong da Chemical Reagent Co. Ltd, Tianjin, China); D-alpha-Tocopheryl polyethylene glycol 1000 succinate (TPGS), poloxamer 188 (Pluronic F68), and poloxamer 407 (Pluronic F127) (BASF, Ludwigshafen, Germany); polyvinyl alcohol 217 (PVA 217) (Kuraray Co. Ltd, Japan, Osaka); polyethylene glycol 2000 (PEG-2000) and polyvinylpyrrolidone K30 (PVP K30) (Tianjin Bodi Chemical Co. Ltd, Tianjin, China); sodium carboxyl methyl cellulose (CMC-Na) (Nanjing Zhongnuo Bioengineering Co. Ltd, Nanjing, China); dimethyl sulfoxide (DMSO), methanol, acetone and acetonitrile of chromatographic grade (Concord Technology Co. Ltd, Tianjin, China). All reagents and chemicals used were of analytical or chromatographic grade.

Methods

Preparation of Amorphous Nanopowder

ETO amorphous nanopowder was prepared using a sonoprecipitation–high pressure homogenization method. Briefly, the crude etoposide was completely dissolved in DMSO. In the presence of sonication (SCIENTZ-IID ultrasonic processor, Scientz Biotechnology Co., Ltd, Ningbo, China), the ETO solution was rapidly added to the aqueous solution containing the stabilizer (Pluronic 127, PVP K30, CMC-Na, sodium deoxycholate, SDS, PVA, TPGS, Pluronic F68 or PEG-2000) to obtain a starting suspension. The starting suspension was then placed in a high-pressure

homogenizer (HPH, Pharmaceutical ultra-high-pressure homogenizer of AH100D, ATS Engineering Inc., Shanghai, China) at 4°C. After that, the nanosuspension was concentrated by centrifugation of 15,000 rpm/min for 15 min using a high-speed refrigerated centrifuge (Anhui Zhongke Zhongjia Scientific Instrument Co. Ltd, Anhui, China). Finally, the supernatant was removed and the precipitate was redistributed in an aqueous solution containing the stabilizer.

Further, the nanosuspension and cryoprotectant were lyophilized together (Labconco, USA) for 24 hours. The lyophilized ETO nanopowder was evaluated for appearance, re-dispersibility and Sf (particle size after lyophilization)/Si (initial particle size) ratio.

Formulation Optimization of ETO Nanopowder

Through the preliminary experiments of different process parameters, X_1 (ultrasonication time), X_2 (ratio of two phases) and X_3 (stabilizer concentration) were selected as the critical process parameters affecting PS (Y_1) and polydispersity index (PDI, Y_2) of ETO nanopowder. Then the prescription was further optimized by BBD. This study was a three-factor central composite design (CCD) using 17 batches. The ultimate goal of optimization was to obtain ETO nanopowder with minimum PS and PDI. The experiment was designed and analyzed by Design-Expert software (Version 8.0.6.1, Stat-Ease Inc., Minneapolis, MN). The variables and their levels are listed in Table 1.

Particle Size and Size Distribution Analysis

ETO nanopowder was re-dispersed in water and evaluated for PS and PDI using dynamic light scattering (ZEN 3600, Zetasizer nano ZS 90, Malvern, UK).²⁴ The re-dispersed ETO nanopowder was diluted 5-fold with distilled water

before measurement. Each sample was measured in triplicate at room temperature.

Morphology Observation

The morphology of the ETO nanopowder was analyzed with a transmission electronic microscope (TEM, JEM 2010; JEOL, Japan) operated at 120 kV.²⁵ The ETO nanopowder was diluted with distilled water before measurement. One drop of the re-dispersed ETO nanopowder was placed on the surface of a copper grid, and the excess liquid was drained on filter paper. The copper grid was strained in 2% phosphotungstic acid for 120 s.

Differential Scanning Calorimetry (DSC)

Differential scanning calorimetry analysis was carried out (DSC-3; METTLER TOLEDO, Switzerland). The samples were accurately weighed (2–5 mg) and placed in a standard aluminum pan, sealed with a lid and perforated. The samples were scanned at a rate of 10°C/min over a range of 40–300°C under 50 mL/min nitrogen flow rate.²⁶

Powder X-Ray Diffraction (PXRD)

The powder X-ray diffraction patterns of the ETO nanopowder were analyzed by D/Max-2400 X-ray fluorescence spectrometer (Rigaku, Osaka, Japan).²⁷ Data were collected with Cu-K α radiation and a voltage of 40 kV, and the scanning speed was 4°/min from 3° to 50°.

Determination of Saturation Solubility

The saturation solubility of coarse powder (PS = 13.33 \pm 3.44 μ m), physical mixture and nanopowder of ETO in different media (distilled water, pH 1.2 solution or pH 6.8 PBS) were analysed by a shaking-flask method.²⁸ Briefly, an excess amount of sample powder was dispersed into 0.5 mL different media, and then shaken for 72 h at 100 rpm and 37°C in the HCR-D210 shaking-air bath (Henan Instruments Co., Ltd, Henan, China). These samples were centrifuged at 15,000 rpm for 15 min, and the supernatant was immediately filtered through a 0.22 μ m syringe filter. Finally, the content of etoposide in samples was detected by high performance liquid chromatography (HPLC) analysis in triplicate. Drug analysis was conducted on a Kromasil 100–5 Phenyl column (250 \times 4.6 mm i.d., 5 μ m, Zhonghuida Technologies, China). The HPLC mobile phase consisted of acetonitrile and 0.02 M sodium acetate (pH 4.0, 30:70, v/v) at a flow rate 1.0 mL/min, column temperature was

Table 1 Independent Factors and Responses in the Box–Behnken Design

Independent Variables	Level		
	Low (-1)	Middle (0)	High (+1)
X_1 , Ultrasonication Time (min)	5.0	7.5	10.0
X_2 , Volume Ratios (organic phase/aqueous phase, v/v)	0.05	0.08	0.10
X_3 , Stabilizer Concentration (% w/v)	0.10	0.17	0.25
Dependent Variables	Goal		
Y_1 , PS (nm)	Minimize		
Y_2 , PDI	Minimize		

maintained at 40°C, and injection volume was 10 µL. The detection wavelength was set at 245 nm.²⁹

In vitro Dissolution Test

The dissolution profiles of coarse powder (PS = 13.33 ± 3.44 µm), physical mixture and nanopowder of ETO in different media (pH 1.2 solution, pH 6.8 PBS or pH 7.4 PBS) were tested with a dissolution apparatus (RC806D). The medium was stirred at 100 rpm and the temperature was set at 37°C. The accurately weighed powder (containing 5 mg etoposide) was added into 200 mL different media. Aliquots of 5 mL were withdrawn at 1, 2, 3, 5, 10, 15, 30, 45 and 60 min and replaced with an equal volume of fresh media. The samples were immediately filtered through a 0.22 µm syringe filter and quantified by HPLC in triplicate.²⁹

Pharmacokinetics Study

Twelve SD male rats weighting 180–220 g were purchased from the Institute of Laboratory Animal Science, Shenyang Pharmaceutical University (Shenyang, China). This study was reviewed and approved by the Laboratory Animal Welfare Ethics Committee of Shenyang Pharmaceutical University (Animal ethical number SYPU - IACUC - C2019 - 7 - 10 - 203) and all the operations were conducted in accordance with the “Guidelines of Shenyang Pharmaceutical University for the Care and Use of Experimental Animals.” Rats were fasted in a specific pathogen-free (SPF) animal house for 12–16 h with free access to water. The rats were randomly divided into two groups (n=6) and orally administered with the suspension of coarse etoposide powder (PS = 13.33 ± 3.44 µm) and ETO nanopowder respectively, each equivalent to 180 mg/kg ETO. Equivalent volume of blood samples (0.5 mL) were obtained from orbital veins and transferred into heparinized tubes at 0.25, 0.5, 1, 2, 4, 6, 8, 10, 12 and 24 h after oral administration. Then, the blood sample was centrifuged at 6000 rpm/min for 10 min.

The plasma was separated and stored at –80°C before analysis. Etoposide concentration in rat plasma was determined by UPLC/MS/MS. The UPLC/MS/MS method is as follows: Chromatographic separation was carried out with an ACQUITY™ UPLC system (Waters Corp., Milford, MA, USA). The separation was conducted on an ACQUITY UPLC™ BEH C₁₈ column (50 mm × 2.1 mm i.d., 1.7 µm, Waters Corp., Milford, MA, USA). The mobile phase consisted of acetonitrile (A) and 0.1%

formic acid in water (B), and was delivered at a flow rate of 0.2 mL/min. The column was kept at 35°C, and the autosampler was maintained at 4°C. The linear gradient elution program was (1) A decreased from 70% to 30% during the first 1.0 min; (2) A was held at 30% for 1.0 min; (3) A was reset to the initial composition in 0.2 min; (4) A was held at 70% for 0.8 min. The injection volume was 5 µL using the partial loop mode. Mass spectrometric detection was carried out with a Waters ACQUITY™ TQD triple quadrupole tandem mass spectrometer (Waters Corp., Manchester, UK) in positive ESI mode. The optimized source/gas parameter was capillary 3.1 kV; cone voltage 3.0 V; radio frequency 0.3 V; source temperature 100°C; desolvation temperature 400°C. Nitrogen was used as the desolvation gas (450 L/h) and cone gas (50 L/h). For collision-induced dissociation, argon was used as the collision gas at a flow rate of 0.2 mL/min. Quantitation was performed using multiple reaction monitoring of the fragmentation transitions of m/z 589.05→228.90 amu for ETO, and m/z 656.89→382.91 amu for the internal standard, with a scan time of 0.02 s per transition.^{30,31} Pharmacokinetic data were collected by Masslynx™ NT4.1 software (Waters Corp., Milford, MA, USA) and processed with the QuanLynx™ program (Waters Corp, Milford, MA, USA).³¹ Pharmacokinetic parameters were analyzed with DAS 2.0 software (Mathematical Pharmacology Professional Committee of China, Shanghai, China).⁴ All descriptive parameters are expressed as mean ± SD.

Results and Discussion

Formulation Development and Optimization

Selection of Organic Solvent, Stabilizers and Process Parameters

In this section, the results and scientific explanations of the formulation selection of ETO nanopowder are presented. The content of the preliminary experiment aimed to delve into the influence of various experimental parameters on PS and PDI, such as the type of organic phase, ratio of two phases, ultrasonication time, the type of stabilizer, stabilizer concentration and HPH operation parameters. Then, the critical process parameters were selected to optimize the formulation of ETO nanopowder by BBD.

The sonoprecipitation–high pressure homogenization method is a combination of top-down and bottom-up methods.^{32,33} In the bottom-up method, the nucleation is

critical to harvest uniform drug nanoparticles in a narrow size distribution. The supersaturation induced by the anti-solvent is vital for the control of the nucleation process. By selecting a suitable antisolvent to achieve high supersaturation, more rapid nucleation can be induced, thereby resulting in crystals of similar size.¹⁵ In this study, the effects of different organic solvents on PS and PDI of ETO nanosuspension were compared (Figure 2A). As revealed

from the results, the precipitation system with acetone and acetonitrile led to the formation of large particulate aggregates, which was not conducive to subsequent processing with a high-pressure homogenizer. However, the suspension with DMSO as antisolvent did not produce this phenomenon. Moreover, the solubility of ETO in DMSO was significantly higher than that in other solvents. Therefore, choosing DMSO as antisolvent can reduce the volume of

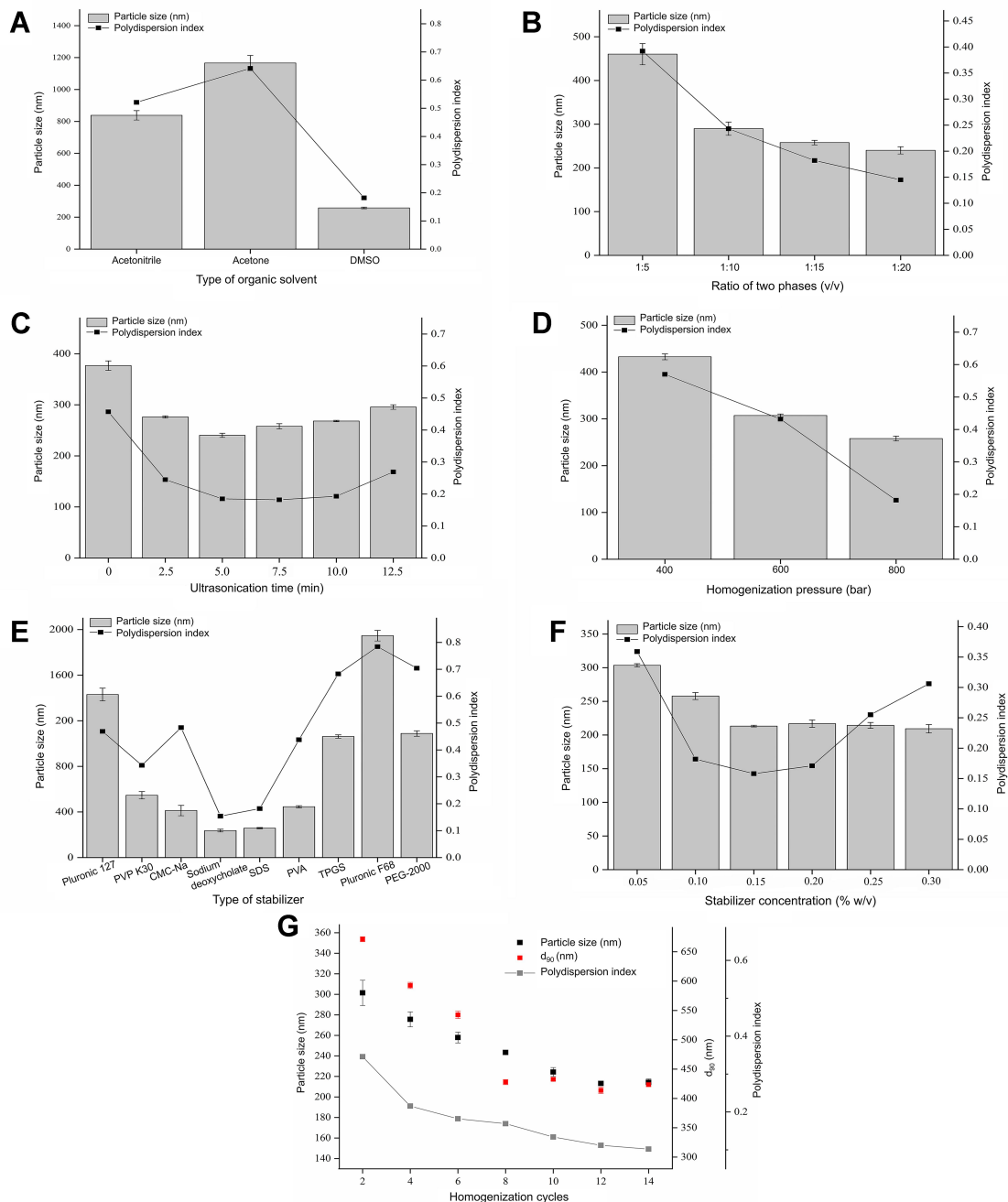


Figure 2 The effect of various parameters, (A) type of organic solvent, (B) ratio of two phases, (C) ultrasonication time, (D) homogenization pressure, (E) type of stabilizer, (F) stabilizer concentration, (G) homogenization cycles in each prescription, on mean PS and PDI of ETO nanopowder.

the organic solvent in the suspension, which is conducive to reduce the PS and PDI of the nanosuspension (Figure 2B). In existing studies, ultrasonication was able to enhance nucleation by producing acoustic cavitation in solution and subsequently shortening the induction period, so as to obtain a suspension with uniform particle size distribution.^{34,35} Accordingly, ultrasonication time is also an important factor affecting PS and PDI of a nanosuspension. As shown in Figure 2C, with the increase in ultrasonication time, PS and PDI of the nanosuspension decreased first and then increased. The reason might be that with the increase in ultrasonication time, the input energy increases, generating heat and thus leading to the aggregation of particles.

High-pressure homogenization refers to a top-down method capable of effectively reducing PS and PDI of nanoparticles.^{36,37} The effect of different homogenization pressure and cycle times on the PS and PDI of the nanosuspension was explored in this study. As demonstrated from the results, with the increase in homogenization pressure and cycle times of high-pressure homogenization, the energy input increased, and the PS and PDI of ETO nanosuspension declined (Figure 2D and G). Moreover, it is noteworthy that with the increase in homogenization cycle times, the decrease in d_{90} was greater than that of the average particle size, indicating that high-pressure homogenization could effectively reduce the large particles in the system and promote the system to be more uniform.

The selection of the stabilizer is the most important step in the preparation of a nanosuspension. The PS and PDI of a nanosuspension can be altered by adding different stabilizers in the crystallization milieu.^{38,39} As suggested in Figure 2E, under identical preparation conditions, the use of charged stabilizers (SDS or sodium deoxycholate) can effectively reduce PS and PDI of ETO nanosuspension. However, the addition of polymers such as Pluronic 127, PVP K30, CMC-Na, PVA 217, TPGS, Pluronic F68 or PEG-2000 could not reduce PS and PDI of ETO nanosuspension. It was therefore demonstrated that electrical repulsion is dominant in the formation of the ETO nanosuspension system. However, though sodium deoxycholate could give excellent PS and PDI to a nanosuspension, it is unstable in gastric acid. Accordingly, SDS was chosen as the stabilizer of ETO nanosuspension in this study. Subsequently, the effect of stabilizer concentration on PS and PDI of the nanosuspension was determined. It was demonstrated that with an increase in SDS concentration,

PS of ETO nanosuspension decreased. However, when a higher stabilizer concentration was added to the formula, the nanosuspension exhibited smaller PS as expected, whereas PDI increased (Figure 2F). Similar results were identified in previous studies.⁴⁰ An explanation for these results is that when the concentration of stabilizer is low, the stabilizer should cover the whole hydrophobic surface of nanoparticles to mask their hydrophobic region, minimizing the free stabilizer in the system. However, when a higher concentration of stabilizer is added, the free stabilizer in the system increases. Under the concentration of free stabilizer higher than the critical micelle concentration, the mentioned free stabilizers tend to form micelles, which may be the main reason for the uneven particle size distribution of the system.⁴⁰

Formulation Optimization

The design matrix of the 17 experiments was designed by BBD with three factors and three levels, as shown in Table 2. These experiments were carried out randomly to reduce systematic error caused by extraneous factors. Then, the responses and relevant data of the 17 experiments were simultaneously fitted to linear, two-factor (2F) and quadratic models by Design Expert software. As shown in Table 3, quadratic models were the best fitted model for PS and PDI, with coefficient of multiple

Table 2 Arrangement and Response Variables of Box–Behnken Design

Formulations	Factors			Response Variables	
	X ₁ (min)	X ₂ (v/v)	X ₃ (% w/v)	Y ₁ (nm)	Y ₂
1	7.5	0.08	0.17	221.4	0.120
2	5.0	0.10	0.17	279.8	0.183
3	10.0	0.08	0.10	357.9	0.324
4	10.0	0.05	0.17	283.6	0.186
5	5.0	0.08	0.10	258.4	0.158
6	7.5	0.08	0.17	220.8	0.123
7	10.0	0.08	0.25	249.7	0.207
8	5.0	0.05	0.17	253.2	0.136
9	7.5	0.10	0.10	308.9	0.294
10	7.5	0.05	0.10	268.9	0.268
11	7.5	0.08	0.17	221.1	0.121
12	7.5	0.05	0.25	200.4	0.147
13	10.0	0.10	0.17	314.8	0.311
14	5.0	0.08	0.25	273.2	0.239
15	7.5	0.10	0.25	253.8	0.237
16	7.5	0.08	0.17	214.8	0.125
17	7.5	0.08	0.17	219.7	0.131

Table 3 Results of Statistical Models for Responses

Models		r^2	Adjusted r^2	Predicted r^2
Linear	Y_1	0.4000	0.2615	0.0097
	Y_2	0.3458	0.1948	-0.0736
2F	Y_1	0.5363	0.2581	-0.2776
	Y_2	0.4968	0.1949	-0.3553
Quadratic	Y_1	0.9887	0.9743	0.8353
	Y_2	0.9605	0.9097	0.3810

determinations (r^2) of 0.9887 and 0.9605, respectively. Accordingly, the best formulation of ETO nanopowder was calculated and predicted by the quadratic equations. The polynomial regression equations of the quadratic model are as follows:

$$\begin{aligned} \text{PS} = & 219.56 + 17.68X_1 + 18.90X_2 - 27.13X_3 \\ & + 1.15X_1X_2 - 30.75X_1X_3 + 3.35X_2X_3 + 45.05X_1^2 \\ & + 18.25X_2^2 + 20.19X_3^2 (P < 0.0001) \end{aligned} \quad (1)$$

$$\begin{aligned} \text{PDI} = & 0.12 + 0.039X_1 + 0.036X_2 - 0.027X_3 \\ & + 0.019X_1X_2 - 0.050X_1X_3 + 0.016X_2X_3 \\ & + 0.038X_1^2 + 0.042X_2^2 + 0.070X_3^2 (P = 0.0004) \end{aligned} \quad (2)$$

In an analysis of variance (ANOVA, Table 4), it was found that ultrasonication time (X_1), ratio of two phases (X_2) and

stabilizer concentration (X_3) were the main factors affecting the particle size of ETO nanopowder. And the influence of the ratio of two phases (X_2) and stabilizer concentration (X_3) was the most significant. The second-order interaction of ultrasonication time and stabilizer concentration (X_1X_3) also had a significant effect on the particle size of ETO nanopowder. For the PDI of ETO nanopowder, in addition to X_1 , X_2 and X_3 , the second-order interaction of ultrasonication time and stabilizer concentration (X_1X_3) was also statistically significant on the PDI of the developed formulations. For a more intuitive assessment of the factor effects on the CQAs, the three-dimensional response surface plots are shown in Figure 2. With the increase in ultrasonication time, PS and PDI of ETO suspension first ascended and then increased (Figure 3A). In Figure 3B and C, the higher the concentration of SDS, the smaller the particle size of ETO nanosuspension, but the PDI first decreased and then increased. These results were consistent with the previous preliminary experimental results. In Figure 3A and C, both PS and PDI decreased with the decrease of the proportion of organic phase to aqueous phase. Therefore, reducing the volume of organic phase not only helps obtain excellent PS and PDI nanosuspension, but also solves the problem of drug precipitation being difficult to recover after centrifugation of nanosuspension due to more organic solvents.

Table 4 Analysis of Variance (ANOVA) in the Quadratic Model for Responses (Y_1 and Y_2)

Source	PS (Y_1)				PDI (Y_2)			
	Sum of Squares	Degree of Freedom	F-value	P-value (Prob > F)	Sum of Squares	Degree of Freedom	F-value	P-value (Prob > F)
Model	27,794.50	9	68.32	<0.0001	0.078	9	18.90	0.0004
X_1	2499.24	1	55.29	0.0001	0.012	1	26.37	0.0013
X_2	2857.68	1	63.21	<0.0001	0.010	1	22.47	0.0021
X_3	5886.12	1	130.21	<0.0001	5.725×10^{-3}	1	12.40	0.0097
X_1X_2	5.29	1	0.12	0.7423	1.521×10^{-3}	1	3.30	0.1123
X_1X_3	3782.25	1	83.67	<0.0001	9.801×10^{-3}	1	21.24	0.0025
X_2X_3	44.89	1	0.99	0.3522	1.024×10^{-3}	1	2.22	0.1799
X_1^2	8543.38	1	188.99	<0.0001	6.000×10^{-3}	1	13.00	0.0087
X_2^2	1401.60	1	31.00	0.0008	7.516×10^{-3}	1	16.29	0.0050
X_3^2	1717.21	1	37.99	0.0005	0.021	1	45.03	0.0003
Residual	316.44	7			3.231×10^{-3}	7		
Lack of Fit	286.47	3	12.74	0.0163	3.155×10^{-3}	3	55.34	0.0010
Pure Error	29.97	4			7.600×10^{-5}	4		
Cor Total	28,110.94	16			0.082	16		

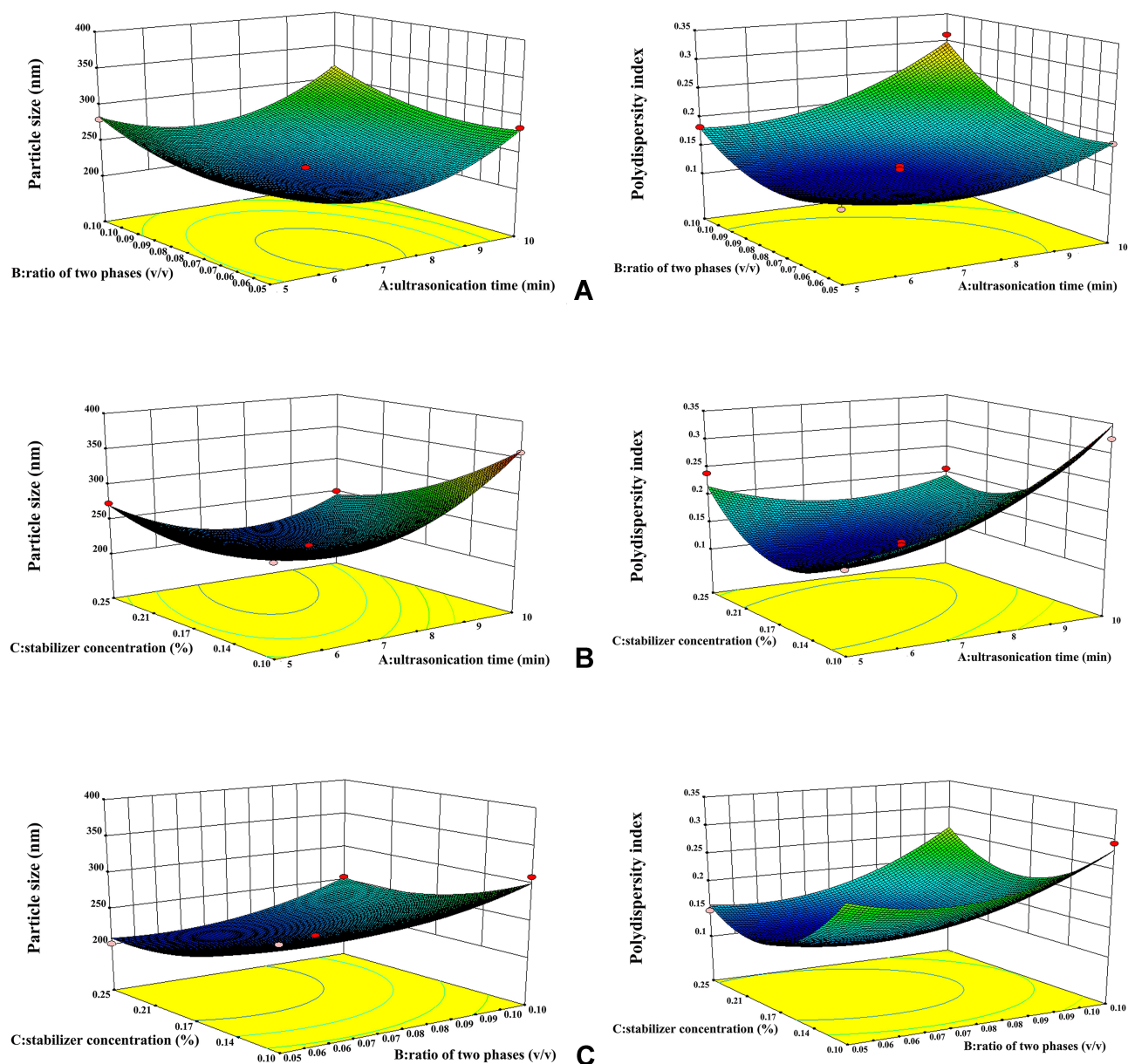


Figure 3 Response surface profiles showed the effects of ultrasonication time (X_1), ratio of two phases (X_2) and stabilizer concentration (X_3) on PS (Y_1) and PDI (Y_2) of ETO nanopowder. **(A)** The effect of second-order interaction of ultrasonication time and stabilizer concentration (X_1X_3) on the PS and PDI of ETO suspension. **(B)** The effect of second-order interaction of ultrasonication time and stabilizer concentration (X_1X_3) on the PS and PDI of ETO suspension. **(C)** The effect of second-order interaction of ultrasonication time and stabilizer concentration (X_2X_3) on the PS and PDI of ETO suspension.

According to the results of the BBD experiment, the optimal formulation is as follows: the ultrasonication time was 7.22 min, the content of stabilizer was 0.2% w/v and the volume ratio of water phase to organic phase was 0.06. The best prediction value was PS of 207.2 nm, PDI of 0.112. The actual particle size and PDI of the best prescription were 211.7 ± 10.4 nm and 0.125 ± 0.028 , which were consistent with the predicted values.

Lyophilization Study

Solidification is a very important step for the preparation of nanopowder. In this paper, we evaluated the type and concentration of cryoprotectant on the appearance, re-dispersibility, PS, PDI and Sf/Si ratio of ETO nanopowder. The results are shown in Table 5. The ETO nanopowder using 1% (w/v) and 2% (w/v) mannitol as the cryoprotectant had excellent performance in all aspects. In order to

Table 5 Effect of Different Cryoprotectants on Physical Characterization

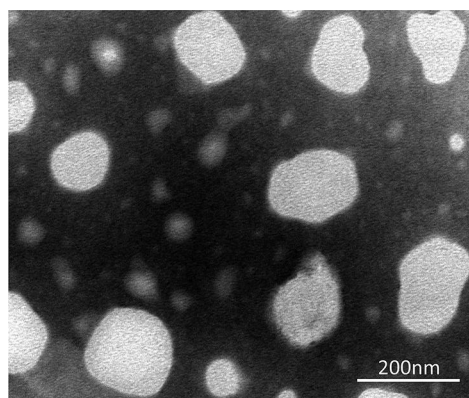
Cryoprotectant (% w/v)	Appearance	Redispersion	PS (nm)	PDI	Sf/Si Ratio
None	–	–	579.7 ± 20.3	0.840 ± 0.128	2.74
1% Mannitol	++	+	217.6 ± 14.3	0.147 ± 0.036	1.03
2% Mannitol	++	+	214.4 ± 10.2	0.150 ± 0.034	1.01
3% Mannitol	++	+	227.6 ± 11.3	0.157 ± 0.041	1.08
1% Maltose	–	+	279.7 ± 12.4	0.240 ± 0.026	1.32
1% Glucose	–	+	257.7 ± 18.5	0.181 ± 0.043	1.22
1% Lactose	–	+	262.3 ± 14.7	0.149 ± 0.032	1.24
1% Sucrose	–	+	266.2 ± 6.5	0.074 ± 0.002	1.26
1% PEG-2000	–	+	284.6 ± 7.2	0.127 ± 0.044	1.34
1% Mannitol + 1% Maltose	+	+	246.4 ± 11.5	0.139 ± 0.062	1.16
1% Mannitol + 1% Glucose	+	+	240.8 ± 17.5	0.138 ± 0.057	1.14
1% Mannitol + 1% Lactose	+	+	276.7 ± 13.5	0.125 ± 0.042	1.31
1% Mannitol + 1% Sucrose	+	+	244.7 ± 15.6	0.132 ± 0.007	1.16
1% Mannitol + 1% PEG-2000	+	+	260.6 ± 14.3	0.133 ± 0.012	1.23

Notes: Appearance: ++ indicated compacted cake; + indicates cake with slightly shrinkage; - indicates non-cake; Redispersion: + indicates easy to redisperse; - indicates hard to redisperse.

increase the drug loading, we selected 1% (w/v) mannitol as the cryoprotectant of ETO nanopowder.

Characterization of ETO Nanosuspension Morphology and Particle Size

The PS of the fresh ETO nanosuspension was 211.7 ± 10.4 nm, PDI was 0.125 ± 0.028 and zeta potential (ZP) was -46.2 mv. The PS of the lyophilized ETO nanopowder was 217.6 ± 14.3 nm and PDI was 0.147 ± 0.036 . After freeze-drying, the particle size of the re-dispersed nanopowder was slightly larger than that of the fresh nanosuspension, which indicated that a certain degree of particle aggregation occurred in the process of freeze-drying. The TEM image of re-dispersed nanopowder was spherical particles with a diameter range of about 200 nm (Figure 4).



Crystalline Characterization

The X-ray powder diffraction patterns of the ETO samples are illustrated in Figure 5A. The major characteristic peaks of ETO (at 2θ of 13.486° , 17.987° and 22.489°) were detected in diffractograms of the coarse powder and physical mixture of ETO (Figure 5A (B and C)), indicating that the crystalline state was maintained in the physical mixture. However, compared with the coarse powder and physical mixture, the ETO nanopowder showed a relatively flat curve without a strong peak (Figure 5A (A)), which confirmed that the ETO in the nanopowder was amorphous.

DSC patterns of coarse ETO powder, SDS, mannitol, the physical mixture and nanopowder are shown in Figure 5B. The thermogram of coarse ETO powder showed

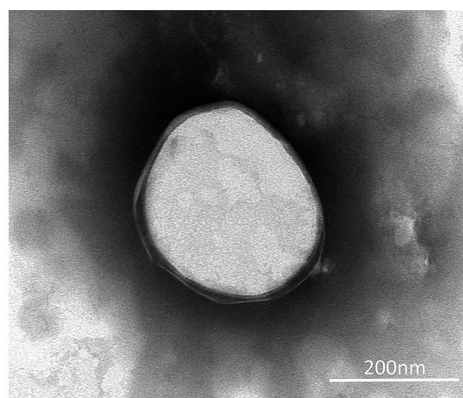


Figure 4 TEM images of ETO nanopowder. (Scale bar, 200 nm).

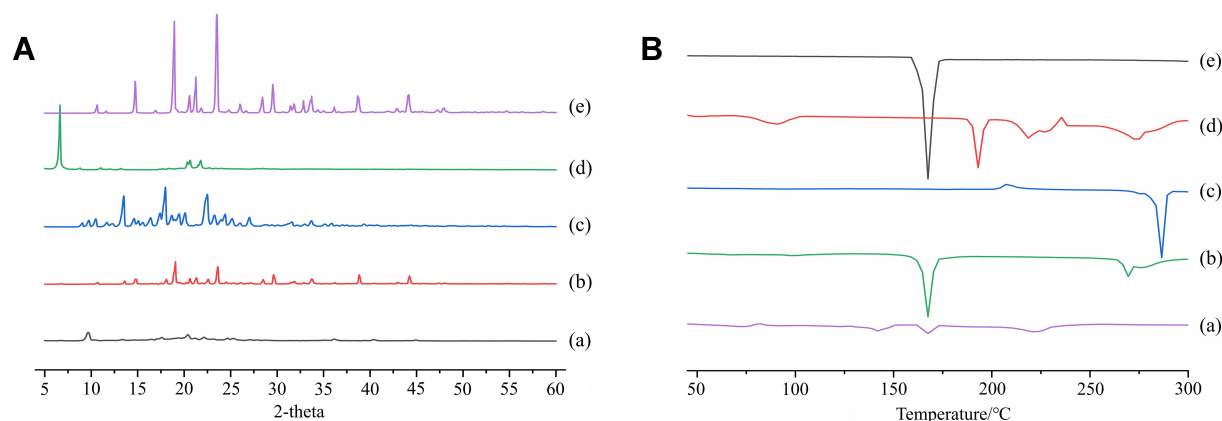


Figure 5 PXRD patterns (A) and DSC thermograms (B) of ETO nanopowder (a); physical mixture of ETO (b); coarse ETO (c); SDS (d); mannitol (e).

a characteristic peak at 286.5°C, which revealed the crystalline nature of the drug (Figure 5B (C)). But a broad peak at around 280°C was observed in the physical mixture (Figure 5B (B)), which may be caused by the overlapping of the melting endothermic peak of ETO and SDS. Mannitol showed a sharp endothermic peak at 168°C, which could be observed in the thermogram of physical mixture and nanopowder of ETO (Figure 5B (E)). However, the DSC curves of ETO nanopowder showed the ETO melting peak was absent (Figure 5B (A)), which indicates that ETO in the nanopowder was mainly amorphous. This result was consistent with that of XRPD. The melting peaks at 142°C and 221°C indicated that the ETO nanopowder had changed, which may be a result of the interaction between the drug and the excipients.

Determination of Saturation Solubility

The saturated solubility of coarse ETO powder, physical mixture and optimized ETO nanopowder in different media are listed in Table 6. In comparison to the coarse powder, the solubility of ETO were significantly increased by 2.31-fold in distilled water, 2.74-fold in pH 1.2 solution and 2.41-fold in pH 6.8 PBS for the nanopowder. The solubility of ETO in physical mixture was also improved,

but the solubility of nanopowder in distilled water, pH 1.2 solution and pH 6.8 PBS were still 1.90-, 2.36- and 1.92-fold higher than that of physical mixture. The Kelvin and Ostwald–Freundlich equations are applicable to illustrate the reason for this phenomenon. According to the Kelvin equation, the decrease of the particle size of nanopowder (from $13.33 \pm 3.44 \mu\text{m}$ to $217.6 \pm 14.32 \text{ nm}$) is helpful to the increase of the surface curvature, which leads to the increase of the dissolution pressure. This causes the equilibrium to move in the direction of dissolution, resulting in an increase in saturated solubility.⁴¹ The Ostwald–Freundlich equation more directly explains that the saturation solubility of drug increases with the decrease of the particle size.⁴² Moreover, the solubilizing effect of SDS is also one of the reasons for the increase of solubility of nanopowder and physical mixture.⁴³

In vitro Dissolution Test

The dissolution profile of coarse ETO powder, physical mixture and optimized ETO nanopowder in different media are shown in Figure 6. ETO nanopowder showed significantly improved dissolution performance in all media compared with coarse powder and physical mixture of ETO. The dissolution rate of ETO nanopowder in pH

Table 6 Saturation Solubility of the Coarse Powder, Physical Mixture and Nanopowder of ETO in Different Media

Media	Saturation Solubility of ETO ($\mu\text{g/mL}$)		
	Coarse ETO	Physical Mixture	Nanopowder
Distilled water	59.80 ± 4.02	$72.53 \pm 4.29^{\#}$	$137.97 \pm 13.33^{\#\#*}$
pH 1.2 solution	86.94 ± 3.79	101.12 ± 8.23	$238.54 \pm 5.98^{\#\#*}$
pH 6.8 PBS	71.43 ± 1.64	$89.61 \pm 3.84^{\#\#}$	$171.84 \pm 2.42^{\#\#*}$

Notes: [#] $P < 0.05$ with respect to coarse ETO; ^{##} $P < 0.01$ with respect to coarse ETO; ^{*} $P < 0.01$ with respect to the physical mixtures.

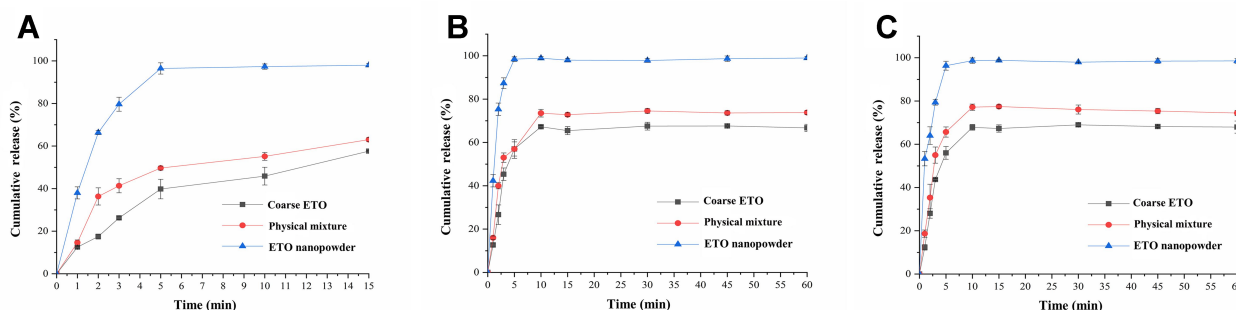


Figure 6 Dissolution profile of coarse powder, physical mixture and nanopowder of ETO in pH 1.2 solution (A); pH 6.8 PBS (B) and pH 7.4 PBS (C).

1.2 solution improved by 1.70-fold and 1.55-fold, with $98.01 \pm 0.82\%$ release in 15 min when compared with coarse powder ($57.60 \pm 0.29\%$) and physical mixture ($63.03 \pm 0.86\%$) of ETO, respectively. Whereas, the concentration of ETO of all samples in pH 1.2 solution began to decrease gradually after 15 minutes, which possibly was due to the stability of etoposide in strong acid solvent. Moreover, the dissolution rate of optimized ETO nanopowder was improved in pH 6.8 PBS ($98.97 \pm 0.78\%$ vs $66.73 \pm 1.68\%$, $73.82 \pm 0.86\%$) and pH 7.4 PBS ($98.53 \pm 1.11\%$ vs $67.95 \pm 2.79\%$, $74.44 \pm 1.23\%$) in 60 min, respectively. In addition, the dissolution velocities of ETO nanopowder in all different media increased significantly. Compared with coarse powder and physical mixture of ETO (<45.3% and <55%, respectively), the dissolution of ETO nanopowder in 3 min was more than 80%. These results suggested that the formation of nanopowder strongly improves the dissolution rate and degree of ETO. The increase of dissolution rate and degree of physical mixture was due to the presence of SDS.⁴³ However, the further dramatic increase of the dissolution rate and degree mainly depends on the decrease of the particle size (from $13.33 \pm 3.44 \mu\text{m}$ to $217.6 \pm 14.32 \text{ nm}$). According to the Noyes–Whitney equation, the decrease of the drug particle size will lead to an increase of the surface

area and a decrease of the diffusion layer thickness, thus increasing the dissolution rate.⁴¹

Pharmacokinetics Study

In order to further verify the advantages of nanopowder in improving the oral absorption of etoposide, the differences between the pharmacokinetic parameters of nanopowder and coarse powder of ETO in rats were compared. The key pharmacokinetic parameters are shown in Table 7, and the plasma drug concentration-time curve is shown in Figure 7. After oral administration, the C_{max} and AUC_{0-t} values of nanopowder were 2.21- and 2.13-fold higher than those of coarse ETO. This improvement of oral bioavailability of ETO could be related to the increase of saturation solubility and dissolution of drug via nanopowder formulation. Additionally, the T_{max} values were 0.25/0.5 h for nanopowder and coarse ETO, respectively. The shortened T_{max} of nanopowder could be related to its ability to accelerate the dissolution of drugs, leading to the rapid drug

Table 7 Pharmacokinetic Parameters of Oral ETO Coarse Powder and Nanopowder

Parameters	Coarse ETO	Nanopowder
C_{max} (ng/mL)	294.87 ± 10.97	$652.52 \pm 54.62^{**}$
AUC_{0-t} (ng [*] h/L)	1176.79 ± 55.85	$2503.79 \pm 222.64^{**}$
T_{max} (h)	0.50 ± 0.00	0.25 ± 0.00
$T_{1/2}$ (h)	14.01 ± 5.01	15.70 ± 5.08

Notes: * $P < 0.05$ with respect to coarse ETO (control group); ** $P < 0.01$ with respect to coarse ETO.

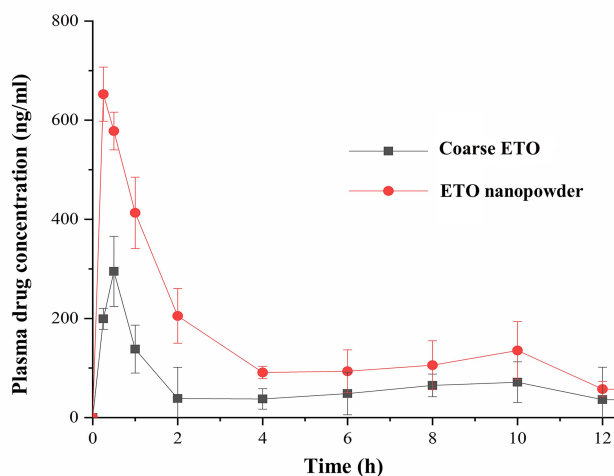


Figure 7 Plasma concentration-time curves of oral ETO coarse powder and the optimized nanopowder.

absorption.⁴⁴ These results suggested that the nanopowder has a good contribution in improving the oral absorption of etoposide.

Conclusion

In this study, etoposide amorphous nanopowder was successfully prepared using a sonoprecipitation–high pressure homogenization method and then optimized with BBD. The physicochemical properties of the optimized formulation were evaluated by TEM, DSC and PXRD. Compared with the coarse powder and physical mixture, the aqueous solubility and in vitro dissolution rate of ETO in nanopowder were significantly improved. Pharmacokinetic study demonstrated that the nanopowder formulation effectively facilitated drug absorption of ETO in rats after oral administration. All the findings revealed that amorphous nanopowder is a promising strategy for enhanced oral bioavailability of hydrophobic compounds, such as ETO.

Abbreviations

AUC, area under the curve; BBD, Box–Behnken design; CCD, central composite design; CMC-Na, sodium carboxyl methyl cellulose; DMSO, dimethyl sulfoxide; DSC, differential scanning calorimetry; ETO, etoposide; HPH, high-pressure homogenizer; PBS, phosphate buffered saline; PDI, polydispersity index; PEG-2000, polyethylene glycol 2000; PS, particle size; PVA 217, polyvinyl alcohol 217; PVP K30, polyvinylpyrrolidone K30; SDS, sodium dodecyl sulfate; SPF, specific pathogen-free; TPGS, D-alpha-Tocopheryl polyethylene glycol 1000 succinate; TEM, transmission electronic microscopy; ZP, zeta potential.

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Disclosure

The authors report no conflicts of interest for this work.

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