Preparation and Optimization of PEG-PLGA Loaded with Vincristine Sulfate and its In vitro Release

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Keywords: PEG-PLGA; Vincristine sulfate; Modified double emulsion method; In vitro drug release

Introduction

Vincristine sulphate (VCR), a dimeric indole alkaloid isolated from the periwinkle plant Catheranthus roseus [1], is one of the most common chemotherapeutic drugs used for the treatment of several forms of malignancies [2,3]. It has been found to be active against the acute lymphoblastic leukemia, Hodgkin’s disease, non-Hodgkin’s lymphoma, neuroblastoma, breast cancer, etc [4,5]. Its antineoplastic effects are to interfere with microtubule assembly and induce tubulin selfassociation into coiled spiral aggregates during mitosis, thus inhibit division of cells [6,7]. Unfortunately, many tumor cells are not sensitive to VCR because of efflux from the tumor cells mediated by P-glycoprotein (Pgp), multidrug resistance-associated protein 1 (MRP1), MRP2 and MRP3 [8,9]. Meanwhile, VCR also has severe nervous system toxicity when administered systematically [10]. To improve the effectiveness and decrease the side effects of VCR, it is a potential approach to deliver VCR by targeting delivery system [11].

Recently biodegradable polymeric NPs for drug delivery have shown significant therapeutic potential [12-14]. Biodegradable polymers such as poly (dl-lacticacid), poly (dl-lactic-co-glycolic acid) and their co-polymers diblocked or multiblocked with PEG have been commonly used to form core–shell structured NPs to encapsulate a variety of therapeutic compounds. These NPs have a number of appealing features: their hydrophobic core is capable of carrying highly insoluble drugs with high loading capacity, while their hydrophilic shell provides steric protection and functional groups for surface modification [15].

In this study, biodegradable polymeric NPs made from polyethylene glycol-modified poly (dl-lactide-co-glycolide) (PEG-PLGA-NPs), which have been extensively used as drug delivery systems for a variety of drugs were investigated for encapsulating VCR. Nanoparticles prepared from PEG-PLGA-NPs are being extensively investigated due to their controlled release, biodegradable and biocompatibility [16,17]. Several methods have been reported for the preparation of biodegradable polymer nanoparticles, such as solvent evaporation [18], nanoprecipitation [19] and salting-out [20]. Among these methods, the modified double emulsion solvent evaporation method is the most common used for the encapsulation of hydrophilic drug. VCR-loaded PEG-PLGA-NPs were prepared by the modified double emulsion (W/O/W) method, and the main experimental factors influencing the characteristics of nanoparticles were investigated and the preparation was optimized. The results showed that the physicochemical characteristics of VCR-loaded PEG-PLGA-NPs were affected by the polymer concentration, the ratio of internal water phase to oil phase, external water phase to oil phase and ultrasound time for the second time. VCR-loaded PEG-PLGA-NPs, with average particle size of 135.9nm, zeta potential of -12.83mV, encapsulation efficiency of 68.2% and drug loading of 8.34%, were prepared under optimal conditions. The release experiments in vitro showed the VCR release from PEG-PLGA-NPs exhibited consequently sustained release for more than 13d, which was in accordance with Higuchi equation.

Abstract

The aim of this study was to prepare the PEG-PLGA loaded with vincristine sulfate nanoparticles (VCR-loaded PEG-PLGA-NPs) and investigate its in vitro release properties. VCR-loaded PEG-PLGA-NPs were prepared by the modified double emulsion (W/O/W) method, and the main experimental factors influencing the characteristics of the nanoparticles were investigated and the preparation was optimized. The results showed that the physicochemical characteristics of VCR-loaded PEG-PLGA-NPs were affected by the polymer concentration, the ratio of internal water phase to oil phase, external water phase to oil phase and ultrasound time for the second time. VCR-loaded PEG-PLGA-NPs, with average particle size of 135.9nm, zeta potential of -12.83mV, encapsulation efficiency of 68.2% and drug loading of 8.34%, were prepared under optimal conditions. The release experiments in vitro showed the VCR release from PEG-PLGA-NPs exhibited consequently sustained release for more than 13d, which was in accordance with Higuchi equation.
water phase using probe sonication for a while to form the W1/O/W2 emulsion. The W1/O/W2 emulsion was gently stirred at room temperature in a fume hood until the evaporation of the organic solvent was complete.

The produced nanoparticles were collected by centrifugation (16000 r/min) at 4°C for 1 hour and washed with distilled water, thrice, to remove the emulsifier. The obtained nanoparticle suspension was freeze-dried and kept in a desiccator.

Particle size distribution and zeta potential

Particle size distribution and zeta potential were determined by Zetasizer 2000. Each sample of nanoparticles preparation was analyzed in duplicate with 30 readings for per nanoparticle sample suspended in distilled water. Figure 1 shows the particle size distribution of the VCR-PEG-PLGA NPs.

Determination of the drug loading capacity of PEG-PLGA-NPs

The obtained nanoparticle suspension was centrifuged (16000 r/min) at 4°C for 1 hour and then the supernatant was detected using UV at 297 nm to calculate the drug entrapment efficiency and loading capacity.

In vitro release

50 mg of VCR-loaded PEG-PLGA-NPs dispersed in 5 mL of PBS (pH = 7.4) were enclosed in dialysis bags and incubated in 500 mL of PBS (pH 7.4), agitated using the shaker at 120 r/min at 37°C. At predetermined time intervals, 5 ml samples were withdrawn and centrifuged for 1 h. The supernatant was assayed and replaced by 5 ml PBS (pH 7.4) immediately after withdrawn. Each nanoparticles batch was analyzed in triplicate.

Results and Discussion

Single factor

The factors that may influence the particle size and drug loading capacity of the VCR-loaded PEG-PLGA-NPs were investigated in the study. The prescription factors include: (1) PEG-PLGA concentration (mg/ml); (2) internal water phase/oil phase (ml/ml); (3) external water phase/oil phase (ml/ml); (4) the method to evaporation the organic phase. The results are showed in Table 1 to Table 8. The tables show that there are some factors that remarkably influenced the particle size and drug loading of the PEG-PLGA-NPs. The best conditions of the prescription were: (1) copolymer: 10 mg/ml; (2) internal water phase/oil phase: 3:10; (3) ultrasound power: (4) the method to evaporation the organic phase.
orthogonal experimental design method was adopted and L9 (34) table orthogonal test according to the results of the single factor study. The physicochemical characteristics of VCR-loaded PEG-PLGA-NPs were commonly prepared by the modified double emulsion method. The shown in Figure 2

Orthogonal test
The best preparation conditions were optimized through the orthogonal test according to the results of the single factor study. The orthogonal experimental design method was adopted and L9 (34) table was chosen for the experiment as the Table 9 showed. The results of visual analysis and variance analysis are shown in Table 10 and Table 11.

In view of the visual analysis, we can see the factors influence the entrapment efficiency listed in a decreasing order as follows: A> C> B > D according to the R value. In the variance analysis we can see that A has s significant different. So the maximum entrapment efficiency of VCR-PEG-PLGA-NPs was obtained when A2B3C1D2 taken (the copolymer concentration: 15mg/ml, the internal water phase/oil phase: 7/20, external water phase/oil phase: 6:1, the ultrasound time for the second time: 1.5min.).

In vitro release
The result of in vitro release of VCR from PEG-PLGA-NPs is shown in Figure 2 After an obvious burst release of about 32% in the first 2 hours, a smooth release of VCR can maintain for about 13 days, its cumulative release was up to 81%. Figure 2. shows that the drug release behavior of PEG-PLGA-NPs was fitted into a Higuchi equation Q=3.4670t+40.346, R²=0.9582.

Conclusions
In the present investigation, VCR-loaded PEG-PLGA-NPs were commonly prepared by the modified double emulsion method. The physicochemical characteristics of VCR-loaded PEG-PLGA-NPs were affected by polymer concentration, internal water phase/oil phase, external water phase/oil phase and ultrasound time for the second time. VCR-loaded PEG-PLGA-NPs with particle size of 135.9nm, zeta potential of -12.83mV, encapsulation efficiency of 68.2%, and drug loading of 8.34% were prepared under optimal conditions. PEG-PLGA-NPs exhibited a sustained in vitro release that lasted for more than 13 days, which fit the Higuchi equation. This could be valuable for realizing the potential of these drug carriers. The bio-distribution, cellular toxicity, pharmacokinetics, pharmacodynamics of VCR-loaded PEG-PLGA-NPs in cells and animal models are currently under investigation in this laboratory.

Acknowledgements
We thank to National Hi-tech Project of China for financial support (2007AA021603, 2007AA021901).

References


