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和厚朴酚自微乳化给药系统的研究(PDF) 分享到:

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Title: Development of self-microemulsifying drug delivery system for honokiol

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关键词: 和厚朴酚; 自微乳化给药系统; 伪三元相图

Keywords: honokiol; self-microemulsifying drug delivery system; pseudo-ternary phase diagram

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摘要: 目的 筛选并优化了和厚朴酚自微乳化给药系统(honokiol-SMEDDS)的处方。方法 利用溶解度, 辅料配伍实验及伪三元相图筛选空白处方, 以粒径和体外释放评价为指标选取最佳处方。结果 优化后的honokiol-SMEDDS处方油相为辛酸甘油酯(MCT), 表面活性剂为聚氧乙烯氢化蓖麻油(Cremophor RH40), 助表面活性剂为二乙二醇单乙基醚(Transcutol P), 油相比例20%, 表面活性剂与助表面活性剂的比值 K_m 为2, 载药量为10%, 平均粒径为28 nm。结论 成功制备了honokiol-SMEDDS, 显著提高了和厚朴酚的溶解度。

Abstract: Objective To screen and optimize the formulation of self-microemulsifying drug delivery system (SMEDDS) for honokiol. Methods The formulation of honokiol-SMEDDS was screened by solubility experiment, compatibility test for excipients and pseudo-ternary phase diagrams. And the best formulation was determined by the particle size and *in vitro* release evaluation. Results In the best formulation of honokiol-SMEDDS, octyl and decyl glycerate (MCT), polyoxyethylene hydrogenated castor oil (Cremophor RH40) and diethylene glycol monoethyl ether (Transcutol P) were used as oil phase, surfactant and co-surfactant, respectively. The percentage of oil phase was 20%, K_m of surfactant to cosurfactant was 2, the drug loading was 10%, and the average particle size was 28 nm. Conclusion The honokiol-SMEDDS is prepared successfully, and the solubility of honokiol is improved significantly.

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