·基础与临床研究 ·

PEG-CL/LA 膜负载甲硝唑药物的体外缓释及抗菌性能研究

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【摘 要】目的 研究以聚乙二醇 - 聚己内酯 - 丙交酯 (PEG-CL/LA) 为聚合物载体的甲硝唑缓释膜的体外抗菌和药物释放性能,旨在研制一种新型的具有抗菌作用的药物缓释膜。方法 用紫外分光光度法测定 5wt%、8wt% 及 10wt% 甲硝唑药膜中药物的体外释放度;选用牙周可疑致病菌具核梭杆菌 (Fn) 和致龋菌变异链球菌 (Sm),采用抑菌圈法研究 3 种药膜对厌氧菌和兼性厌氧菌的抗菌性能。结果 (1) 药膜在载药量为 5wt%~10wt% 时具有较好的缓释性能,药物释放时间随着甲硝唑含量的增加而缩短,3 种比例的载药膜的药物释放时间均可达 7 天以上,其中,5wt% 药膜可达 10 天。(2) 随着药物含量增加,抗菌作用逐渐增强,5wt% 载药膜对 Sm 和 Fn 的抑菌圈直径分别为 (8.00±0.16) mm 和 (8.07±0.09) mm; 8wt% 载药膜为 (10.13±0.19) mm 和 (11.13±0.09) mm; 10wt% 载药膜为 (12.07±0.09) mm 和 (19.33±0.94) mm。该药膜对具核梭杆菌的抑制作用强于对变异链球菌的抑制作用。结论 PEG-CL/LA 载药膜对牙周致病菌的抑制作用效果明显,且具有良好的药物缓释性能,是一种有望用来辅助治疗牙周疾病的药物膜。

【关键词】 甲硝唑聚乙二醇 - 己内酯 - 丙交酯聚合物 药物缓释 抗菌性

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In vitro study of drug release properties and antibacterial activity of medical PEG-CL/LA membranes loaded metronidazole

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[Abstract] Objective To evaluate drug release properties and antibacterial effect of metronidazole sustained-release membranes made of polyethylene glycol-caprolactone-lactide(PEG-CL-LA)with various drug loadings (5wt%, 8wt% and 10wt%). Methods 1. In vitro drug release properties were examined by ultraviolet spectrophotometry. 2. Antibacterial effect on Streptococcus mutans and Fusobacterium nucleatum was observed on solid culture medium. Results 1. Membranes tested here released drug slowly. Time required for drug release was longer than 7 days, and was decreased by increasing drug loading. 2. Membranes had strong inhibitory effect on both S. mutans and F. nucleatum, and antibacterial activity was enhanced with higher drug loading. Membranes loaded with drug at 10wt% showed larger inhibition zone with a diameter of (12.07 \pm 0.09) mm and (19.33 \pm 0.94) mm

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