

论文

抗支原体喹诺酮的合成及其构效关系

杨玉社;嵇汝运;陈凯先;叶辉;武济民

中国科学院上海药物研究所, 上海 200031

摘要:

目的: 开发新型抗支原体药物。方法与结果: 设计合成了一系列新型左旋氧氟沙星类似物(18~24), 测试其体外抗支原体活性(MIC值), 并讨论了他们的构效关系。结论: 所合成的化合物有较好的抗支原体活性。哌嗪环或高哌嗪环4位氮原子有吸电子基团时, 可能有利于提高喹诺酮的抗支原体活性。

关键词: 左旋氧氟沙星 氟喹诺酮 合成 抗菌活性

STUDIES ON SYNTHESIS AND STRUCTURE-ACTIVITY RELATIONSHIPS OF ANTIMYCOPLASMA QUINOLONES

Yang Yushe; Ji Ruyun; Chen Kaixian; Ye Hui and Wu Jimin

Abstract:

AIM: To develop new antimycoplasma drugs. METHODS and RESULTS: A series of new analogues of (S)-(-)-ofloxacin with antimycoplasma activities were prepared. Compounds 18~24 were new compounds. Their in vitro susceptibilities to mycoplasma were tested. The influences on structure-activity relationships were also discussed. CONCLUSION: The synthesized compounds have good activities against mycoplasma. The electron-withdrawing groups on the 4-position of piperazine or homopiperazine may be favorable for antimycoplasma activity.

Keywords: mycoplasma (S)-(-)-ofloxacin synthesis antimycoplasma quinolone

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作者简介:

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