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论文	扩展功能
3,4-二氢-海南新碱类似物的合成及抗溃疡作用	本文信息
刘瑞武;高由松;张守仁;梁晓天	Supporting info
中国医学科学院中国协和医科大学药物研究所,北京100050	▶ PDF <u>(699KB)</u> ▶ [HTML全文]
	▶参考文献
摘要:	服务与反馈
为考察3,4-二氢-海南新碱类似物C ₁ 取代基对化合物抗溃疡活性的影响,设计并合成了13对共26个新的3,4-二氢-海南新碱类似物,A,B互为非对映异构体,大部分化合物在大鼠冷应激溃疡模型中有一定的抗溃疡活性,其中IX ₃ A,IX ₇ A,IX ₈ A,IX ₁₂ A,IX ₁₂ B,IX ₁₃ A6个化合物表现较强活性,超过西咪替丁。对构效关系进行了初步分析。	▶把本文推荐给朋友 ▶加入我的书架 ▶加入引用管理器 ▶引用本文
关键词: 3.4-二氢-海南新碱类似物 抗溃疡活性 构效关系	▶ Email Alert ▶ 文章反馈
SYNTHESIS AND ANTIULCER ACTIVITY OF 3,4-DIHYDRO-HAINANENSINE ANALOGS	▶浏览反馈信息 本文关键词相关文章
Liu Ruiw; Gao Yousong; Zhang Shouren and Liang Xiaotian	▶3.4-二氢-海南新碱类似物 ▶抗溃疡活性 ▶构效关系
Abstract:	本文作者相关文章 ▶ 刘瑞武
In order to search for new compounds with new mode of action, high antiulcer activity and lower toxicity,	
26(13 pairs of diastereoisomer A and B) 3,4-dihydro-hananensine analogs were synthesized. All compounds were tested in M ₁ receptor combined assay and gastric ulcer induced by cold-immersion	▶张守仁
stress in rats. Most compounds showed antiulcer activity, among them IX_3A , IX_7A , IX_8A , $IX_{12}A$, $IX_{12}B$	▶梁晓天
and IX ₁₃ A exhibited more potent antiulcer activity than the control compound cimetidine. Meanwhile, the relationship between their structures and activity was discussed.	. asmea
relationship between their structures and activity was discussed.	Article by
	Article by Article by
Keywords: Antiulcer activity SAR studies 3,4-Dihydro-hainanensine analogs	Article by
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