

中文标题 🔻 检索 跨刊检索

Home 注册 订阅 英文版

中国中药杂志 **China Journal of Chinese Materia Medica**

免疫亲和柱净化-在线柱后光化学衍生-HPLC-FLD 同时测定甘草中黄曲霉毒素 $\mathbf{B_1,B_2,G_1,G_2}$ 和赭曲霉毒素A的含量

投稿时间: 2011-03-04 责任编辑: 马超一 <u>点此下载全文</u>

引用本文,韦日伟杨小驱-仇峰-杨美华-翠洁萍-兔疫亲和柱净化-在线柱后光化学衍生-HPLC-FLD 同时测定甘草中黄曲霉毒素 $\mathbf{B}_1\mathbf{B}_2\mathbf{G}_1\mathbf{G}_2$ 和越曲霉毒素A的含量[J].中国中药杂志.2011.36(17):2342.

DOI: 10.4268/cjcmm20111708

摘要点击次数:706

全文下载次数: 247











作者 中文 名	作者英 文名	单位中文名	单位英文名	E-Mail
<u>事日</u> <u>伟</u>	WEI Riwei	中国医学科学院 中国协和 医科大学 药用植物研究所、 北京 100193 广西中医学院,广西 南宁 530001	Institute of Medicinal Plant Development, Chinese Academy of Medical Sciences, Peking Union Medical College, Beijing 100193, China Guangxi Traditional Chinese Medical University, Nanning 530001, China	
<u>杨小</u> 丽	YANG Xiaoli	中国医学科学院 中国协和 医科大学 药用植物研究所、 北京 100193 江苏大学 药学院,江苏 镇 江 212013	Institute of Medicinal Plant Development, Chinese Academy of Medical Sciences, Peking Union Medical College, Beijing 100193, China School of Pharmacy, Jiangsu University, Zhenjiang 212013, China	
仇峰	QIU Feng	中国医学科学院 中国协和 医科大学 药用植物研究所, 北京 100193	Institute of Medicinal Plant Development, Chinese Academy of Medical Sciences, Peking Union Medical College, Beijing 100193, China	
<u>杨美</u> 生	YANG Meihua	中国医学科学院 中国协和 医科大学 药用植物研究所, 北京 100193	Institute of Medicinal Plant Development, Chinese Academy of Medical Sciences, Peking Union Medical College, Beijing 100193, China	yangmeihua15@hotmail.com
<u>單洁</u> <u>李</u>	QIN Jieping	广西中医学院,广西 南宁 530001	Guangxi Traditional Chinese Medical University, Nanning 530001, China	

基金項目:国家"重大新药创制"科技重大专项(2009ZX09502-025);国家中医药管理局中医药行业科研专项(200807042)

中文摘要:目的:建立同时检测甘草中黄曲霉毒素B₁,B₂,G₁,G₂和赭曲霉毒素A的免疫亲和柱净化-在线柱后光化学衍生- HPLC- FLD 测定的方法。 方法: 拌品经甲醇-水(80:20)超声漫取后,用兔疫亲和肚净化和富集;以甲醇和0.5% 乙酸溶液为流动相进行梯度完 脱.通过柱后光化学衍生,荧光检测器测定。 结果:黄曲霉毒素G₂G₁B₂B₁和蘇曲霉毒素A的检测限分别为0.02,0.06,0.015,0.03,0.25 $\mu g \cdot k g^{-1}$,平均加样回收率为76.0%~103%,RSD低于13%。 结论 :该方法快速简便、准确,可用于甘草中同时测定黄曲霉毒素 B_1,B_2,G 1,G,和赭曲霉毒素A的含量。

中文关键词:<u>真菌毒素 甘草 免疫亲和柱 光化学衍生</u> <u>HPLC</u> <u>荧光检测</u>

 $Simultaneous\ determination\ of\ aflatoxin\ B_1,B_2,G_1,G_2\ and\ ochratoxin\ A\ in\ \textit{Glycyrrhiza\ uralensis}\ by$ HPLC-FLD after immunoaffinity column with online post-column photochemical derivatization

Abstract:Objective: To develop a method for the simultaneous determination of aflatoxin B₁, B₂, G₁, G₂ and ochratoxin A in Glycyrrhiza uralensis by HPLC-FLD after immunoaffinity column with online post-column photochemical derivatization. Method: Sample was extracted with MeOH: $\rm H_{2}O~(80~\odot~20)$ and cleaned up by immunoaffinity column. The toxins were separated by reversed-phase HPLC and the mobile phase was consisted of methanol and 0.5% acetic acid solution with gradient elution. The determination was carried out by fluorescence detector after photochemical derivatization. Result: The detection limits of aflatoxin G_2 , G_1 , B_2 , B_1 and ochratoxin A were 0.02, 0.06, 0.015, 0.03 and 0.25 $\mu g \cdot k g^{-1}$, respectively. The recoveries of analytes were from 76.0% to 103% and the relative standard deviations (RSDs) were below 13%. Conclusion: The method is a simple, accurate and can be used to determine the contents of aflatoxin B₁, B₂, G₁, G₂ and ochratoxin A in G. uralensis simultaneously.

keywords:mycotoxin Glycyrrhiza uralensis immunoaffinity column photochemical derivatization HPLC fluorescence detector

查看全文 查看/发表评论 下载PDF阅读器

版权所有 © 2008 《中国中药杂志》编辑部 京ICP备11006657号-4 您是本站第7615384位访问者 今日一共访问5273次 当前在线人数:582 北京市东直门内南小街16号 邮编:100700技术支持:北京勤云科技发展有限公司 linezinglilli.