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ISSN 1672-3651

CN 32-1708/R

Chinese Journal of Natural Medicines

中国天然药物

ZHONGGUO TIANRAN YAOWU

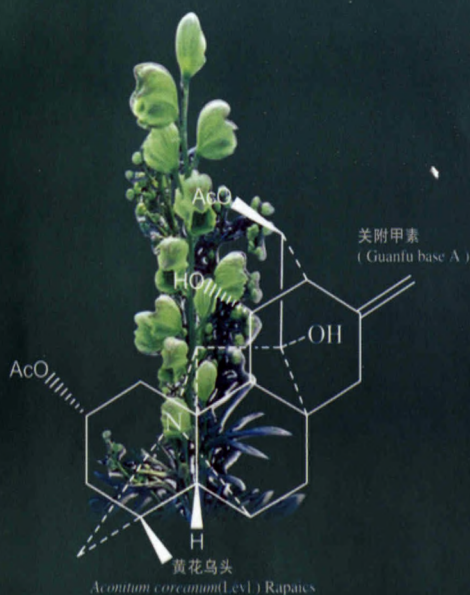
2012 3

第10卷 第3期

2012年 5月

Volume 10, Number 3

May, 2012



中国药科大学
China Pharmaceutical University 主办

中国药学会
Chinese Pharmaceutical Association

科学出版社 出版
Science Press



ELSEVIER

Chinese Journal of Natural Medicines

(Bimonthly, Founded in 2003)

Volume 10, Number 3

May 2012

ISSN 1672-3651 CN 32-1708/R

Directed by Ministry of Education PRC

Sponsored by China Pharmaceutical University

Chinese Pharmaceutical Association

Editors-in-Chief WU Xiao-Ming, SUN Han-Dong

Published by Science Press

Edited by Editorial Office of

Chinese Journal of Natural Medicines

Executive editors DING Zuo-Qi, ZHENG Xiao-Nan

Address 24 Tongjia Xiang, Nanjing, China

Post code 210009

Tel 86-25-83271565, 83271568

Fax 86-25-83271229, 83271568

E-Mail cjnm@cpu.edu.cn

Web Http://mc03.manuscriptcentral.com/cjnm

Http://www.cpcujnm.com/zgryw

Printed by Printing House of NUCT

Domestic Distributor Nanjing Post Office

Domestic Code 28-306

Overseas Distributor China International Book Trading Corporation

(P.O. Box 399, Beijing, China)

Overseas Code BM 1759

Published on May 20, 2012

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China Pharmaceutical University

Chinese Pharmaceutical Association

Indexed in:

Science Citation Index Expanded (SCIE)

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BIOSIS Toxicology

CAB Abstracts

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Ind Appl Microbiol Abstracts, Environ Sci Poll Manag,

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CHINESE JOURNAL OF NATURAL MEDICINES

(Zhongguo Tianran Yaowu)

CONTENTS

Review

- Podophyllotoxin, a medicinal agent of plant origin: past, present and future
.....GUERRAM Mounia, JIANG Zhen-Zhou, ZHANG Lu-Yong (161)
- Advances on structure-activity relationship of NQO1-target antitumor quinones
.....LIAO Ke, NIU Fang, HAO Hai-Ping, WANG Guang-Ji (170)

Original Papers

Biological and Pharmaceutical Activity

- Protective effect of *Ornithogalum saundersiae* Ait (Liliaceae), against acetaminophen-induced acute liver injury via CYP2E1 and HIF-1 α
.....WAN Ying, WU Yan-Ling, LIAN Li-Hua, NAN Ji-Xing (177)
- β -Cell protective efficacy, hypoglycemic and hypolipidemic effects of extracts of *Achillea millefolium* in diabetic rats
.....Khalid G. Mustafa, Bashir A. Ganai, Seema Akbar, Mohamad Y. Dar, Akbar Masood (185)
- In vitro* antioxidant activity of *Parnassia wightiana* W. extracts
.....WEN Xin-Bao, MIAO Fang, ZHOU Le, ZHANG Miao, HE Quan-Lei (190)

Analysis Studies

- UPLC-MS based metabolic profiling of the phenotypes of *Acanthopanax senticosus* reveals the changes in active metabolites distinguishing the diversities of the plant grown in northeast area of China
.....SUN Hui, HAN Ying, ZHANG Ai-Hua, MENG Xiang-Cai, WANG Zhen-Yue, SUN Wen-Jun, SUN Hai-Feng, WANG Xi-Jun (196)
- Immobilization mediated enhancement of phyllanthin and hypophyllanthin from *Phyllanthus amarus*
.....J. S. Thakur, R. K. Agarwal, M. D. Kharya (207)
- Simultaneous quantitation of eight active components in crude and processed *Radix Scrophulariae* extracts by high performance liquid chromatography with diode array detector
.....CAO Gang, CONG Xiao-Dong, CAI Hao, LI Xiao-Meng, JI Jin-Yu, ZHANG Yun, CAI Bao-Chang (213)

Natural Product Chemistry

- Two new cassane diterpenoids from the seeds of *Caesalpinia sappan* Linn.
.....ZHANG Jing-Yu, WU Fei-Hua, QU Wei, LIANG Jing-Yu (218)
- A new prenylated chalcone from the seeds of *Millettia pachycarpa*
.....SU Xue-Hui, LI Cong-Ying, ZHONG Yu-Jiao, YUAN Zhi-Peng, LI Yan-Fang, LIANG Bing (222)
- A new monoterpenoid indole alkaloid from *Ervatamia chinensis*
.....GUO Ling-Li, ZHANG Yu, HE Hong-Ping, LI Yan, YU Jian-Ping, HAO Xiao-Jiang (226)
- Cytotoxic sesquiterpene lactones from *Vernonia bockiana*
.....LIAO Shang-Gao, WANG Zhen, LI Jing, LIU Ying, LI Yue-Ting, ZHANG Li-Juan, LONG Qing-De, WANG Yong-Lin (230)
- Cucurbitacins from fruits of *Aquilaria sinensis*
.....MEI Wen-Li, LIN Feng, ZUO Wen-Jian, WANG Hui, DAI Hao-Fu (234)
- Cytotoxic activity of five limonoids from *Melieae* cortex and their structure-activity relationship
.....DENG Zhi-Peng, LIU Shao-Chao, CUI Shu-Xiang, ZHOU Ling, YAO Qing-Qiang (238)

中国天然药物

Chinese Journal of Natural Medicines



双月刊, 2003 年创刊

2012 年 5 月 第 10 卷 第 3 期

主管单位 中华人民共和国教育部
主办单位 中国药科大学
中国药学会
主编 吴晓明 孙汉薰
出版 科学出版社
编辑 《中国天然药物》编辑部

地址 南京童家巷 24 号
邮政编码 210009
电话 86-25-83271565, 83271568
传真 86-25-83271229, 83271568
电子信箱 cjnm@cpu.edu.cn
网址 <http://mc03.manuscriptcentral.com/cjnm>
<http://www.cpcujnm.com/zgtryw>

印刷 南京工大印务有限公司
国内发行 南京市邮政局
国内邮发代号 28-306
国外发行 中国国际图书贸易总公司
(北京 399 信箱, 100044)
国外邮发代号 BM 1759
广告经营许可证 3200004050738
广告部电话 86-25-83271565, 83271568

国际标准刊号 ISSN 1672-3651
国内统一刊号 CN 32-1708/R
第 10 卷 第 3 期 Vol. 10 No. 3
2012 年 5 月 20 日出版
国内定价每期 30.00 元 全年 180.00 元
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■ 责任编辑 丁佐奇 郑晓南
■ 英文审校 史志祥
■ 绘图 巢静妍

综述

- 天然来源的鬼臼毒素研究进展 (英文)
..... GUERRAM Mounia, 江振洲, 张陆勇 (161)
- 靶向 NQO1 抗肿瘤配体关系研究进展 (英文)
..... 廖珂, 牛芳, 郝海平, 王广基 (170)

论文

生物学与药理活性

- 虎眼万年青通过抑制 CYP2E1 和 HIF-1 α 保护对乙酰氨基酚诱导的急性肝损伤 (英文)
..... 宛莹, 吴艳玲, 廉丽花, 南极星 (177)
- 西洋蓍草提取物对糖尿病大鼠的 β 细胞保护活性及降糖、降脂作用 (英文)
..... Khalid G. Mustafa, Bashir A. Ganai, Seema Akbar, Mohamad Y. Dar, Akbar Masood (185)
- 苍耳七提取物的体外抗氧化活性研究 (英文)
..... 温新宝, 苗芳, 周乐, 张森, 何全磊 (190)

药物分析

- 基于 UPLC-MS 的代谢谱区分生长在中国东北地区植物刺五加多样性的研究 (英文)
..... 孙晖, 韩莹, 张爱华, 孟祥才, 王振月, 孙文军, 孙海峰, 王喜军 (196)
- 固定化导提提高苦味叶下珠中的海冰叶下珠脂素和叶下珠次素的含量 (英文)
..... J. S. Thakur, R. K. Agarwal, M. D. Kharya (207)
- HPLC-DAD 法同时测定玄参炮制前后 8 个有效成分的含量 (英文)
..... 曹岗, 丛晓东, 蔡皓, 李晓萌, 李金玉, 张云, 蔡宝昌 (213)

天然产物化学

- 苏木种子中两个新的卡山烷型二萜 (英文)
..... 张敬宇, 吴斐华, 曲玮, 梁敬钰 (218)
- 厚果鸡血藤种子中的一个新异戊烯基萜耳酮 (英文)
..... 苏雷会, 李聪颖, 钟玉蛟, 袁志鹏, 李廷芳, 梁冰 (222)
- 中国狗牙花中一个新单萜吡啶生物碱 (英文)
..... 郭伶俐, 张于, 何红平, 李艳, 郝建平, 郝小江 (226)
- 南川斑鸠菊中的抗肿瘤倍半萜内酯 (英文)
..... 廖尚高, 王珍, 李靖, 刘影, 李月婷, 张丽娟, 龙庆德, 王永林 (230)
- 白木香果实中的葫芦素 (英文)
..... 梅文莉, 林峰, 左文健, 王辉, 戴好富 (234)
- 苦楝皮中 5 个柠檬苦素化合物的细胞毒性以及构效关系 (英文)
..... 邓志鹏, 刘少超, 崔淑香, 周玲, 姚庆强 (238)

Protective effect of *Ornithogalum saundersiae* Ait (Liliaceae) against acetaminophen-induced acute liver injury via CYP2E1 and HIF-1 α

WAN Ying ^Δ, WU Yan-Ling ^Δ, LIAN Li-Hua, NAN Ji-Xing *

Key Laboratory for Natural Resource of Changbai Mountain & Functional Molecules, Ministry of Education, College of Pharmacy, Yanbian University, Yanji 133002, China

The hepatoprotective effects of total saponin from *Ornithogalum saundersiae* Ait (Liliaceae) (OC) against acetaminophen (APAP)-induced acute liver injury *in vivo* and *in vitro* were investigated. The results suggest that the hepatoprotective mechanism of OC might be related to antioxidation via blocking the CYP2E1, and mediating reactive oxygen species scavenging and accumulation of hypoxia-inducible factor (HIF)-1 α .

[Research funding] This project was supported by the National Natural Science Foundation of China (Nos. 81160538 and 30960511).

β -Cell protective efficacy, hypoglycemic and hypolipidemic effects of extracts of *Achillea millifolium* in diabetic rats

Khalid G. Mustafa ^{1,2}, Bashir A. Ganai ^{1*}, Seema Akbar ², Mohamad Y. Dar ², Akbar Masood ¹

¹Department of Biochemistry, Faculty of Biological Sciences, University of Kashmir, Srinagar 190006 Kashmir, India;

²Drug Standardisation Research Unit, Regional Research Institute of Unani Medicine (CCRUM), Naseem Bagh, University of Kashmir Campus, Srinagar 190 006 Kashmir, India

The therapeutic uses of *Achillea millifolium* in diabetic rats were evaluated. The results indicate that the extracts showed significant decrease in blood glucose level, TGL, VLDL, cholesterol, SGOT, SGPT, and ALP in diabetic rats, and prevented the β -cells of pancreas from the cytotoxic effects of Alloxan monohydrate.

***In vitro* antioxidant activity of *Parnassia wightiana* W. extracts**

WEN Xin-Bao ¹, MIAO Fang ^{1*}, ZHOU Le ², ZHANG Miao ¹, HE Quan-Lei ¹

¹College of Life Sciences, Northwest A & F University, Yangling 712100, China;

²College of Science, Northwest A & F University, Yangling 712100, China

The total flavonoid contents and *in vitro* antioxidant activity of different extracts from *Parnassia wightiana* and their correlation were studied. The total flavonoid content in ethyl acetate extract (EAE) was the highest, followed *n*-butanol extract (BE), water extract (WE) and chloroform extract (CE). WE and BE showed better antioxidant activity than CE and EAE. Each extract showed concentration dependent antioxidant activity and a high correlation between content of flavonoids and antioxidant activity.

[Research funding] This project was supported by the National Natural Science Foundation of China (Nos. 31000865, 30771454, 31172365, 31170366).

UPLC-MS based metabolic profiling of the phenotypes of *Acanthopanax senticosus* reveals the changes in active metabolites distinguishing the diversities of the plant grown in northeast area of China

SUN Hui ¹, HAN Ying ¹, ZHANG Ai-Hua ¹, MENG Xiang-Cai ^{1*}, WANG Zhen-Yue ², SUN Wen-Jun ¹, SUN Hai-Feng ², WANG Xi-Jun ^{1*}

¹National TCM Key Laboratory of Serum Pharmacology-Chemistry, Heilongjiang University of Chinese Medicine, and Key Pharmacometabolomics Platform of Chinese Medicines, Harbin 150040, China;

²Department of Chinese Herbal Medicinal Resource, Heilongjiang University of Chinese Medicine, Harbin 150040, China

UPLC-MS technology-based metabolic profiles was used to discriminate phenotypes of *Acanthopanax senticosus*. By processing the metabolomic data with principal component analysis (PCA), partial least squares-discriminant analysis (PLS-DA) and orthogonal projection to latent structures (OPLS) analysis, eleven secondary metabolites were found to be responsible for classing the two varieties of *Acanthopanax senticosus*. They might be taken as potential biological markers of sexual type. These marker metabolites include flavone, coumarin, terpenes and organic acids, which are the main active constituents related to the therapeutic effects of *Acanthopanax senticosus*. As a result, by monitoring these highly expressed metabolites and through selective cultivation, an *Acanthopanax senticosus* with higher amounts of medicinally relevant constituents could be created.

Immobilization mediated enhancement of phyllanthin and hypophyllanthin from *Phyllanthus amarus*

J. S. Thakur*, R. K. Agarwal, M. D. Kharya

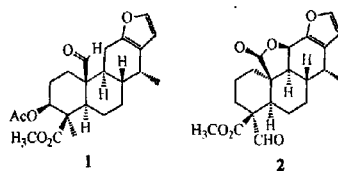
Plant Biotechnology Laboratory, Department of Pharmaceutical Sciences, Dr. Hari Singh Gour Central University, Sagar (M.P.) 470003, India

A cost effective process was developed for enhancing phyllanthin and hypophyllanthin utilizing the immobilization technique. The study revealed that an increase in the content of phyllanthin and hypophyllanthin was elicitor concentration dependent and silver nitrate treatment gave a maximum yield of hepatoprotective bioactives as compared to the other abiotic elicitors used.

Two new cassane diterpenoids from the seeds of *Caesalpinia sappan* Linn.

ZHANG Jing-Yu, WU Fei-Hua, QU Wei, LIANG Jing-Yu*

Department of Natural Medicinal Chemistry, China Pharmaceutical University, Nanjing 210009, China



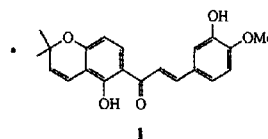
Phanginin L (1) and phanginin M (2) are two new cassane diterpenoids isolated from the seeds of *Caesalpinia sappan*. Compound 1 showed moderate cytotoxicity against HepG2 cell line with IC_{50} of $9.13 \mu\text{g}\cdot\text{mL}^{-1}$.

A new prenylated chalcone from the seeds of *Milletia pachycarpa*

SU Xue-Hui, LI Cong-Ying, ZHONG Yu-Jiao, YUAN Zhi-Peng, LI Yan-Fang*, LIANG Bing

Department of Pharmaceutics & Bioengineering, School of Chemical Engineering, Sichuan University, Chengdu 610065, China

The chemical constituents from the seeds of *Milletia pachycarpa* were investigated. Compound 1, 3-hydroxy-4-methoxylonchocarpin, is a new prenylated chalcone, and compounds 2-4 (4-methoxylonchocarpin, isobavachromene, dorsipoinsettifolin) were isolated from this genus for the first time. Compounds 1-4 selectively showed moderate inhibition against one or more of the tested cell lines.



[Research funding] This project was supported by the Key Program of the National Natural Science Foundation of China (No. 30730111).

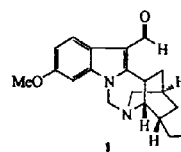
A new monoterpenoid indole alkaloid from *Ervatamia chinensis*

GUO Ling-Li^{1,2}, ZHANG Yu², HE Hong-Ping², LI Yan², YU Jian-Ping^{1*}, HAO Xiao-Jiang^{2*}

¹School of Life Sciences, Guizhou University, Guiyang 550025 China;

²State Key Laboratory of Phytochemistry and Plant Resources in West China, Kunming Institute of Botany, Chinese Academy of Sciences, Kunming 650201, China

The chemical constituents of the whole plants of *Ervatamia chinensis* was studied. Compound 1 (ervachinin E) is a new monoterpenoid indole alkaloid and with moderate anti-tumor activity, while five known compounds are reported from this species for the first time.



[Research funding] This project was supported by the National Natural Science Foundation of China (No. 31100259).