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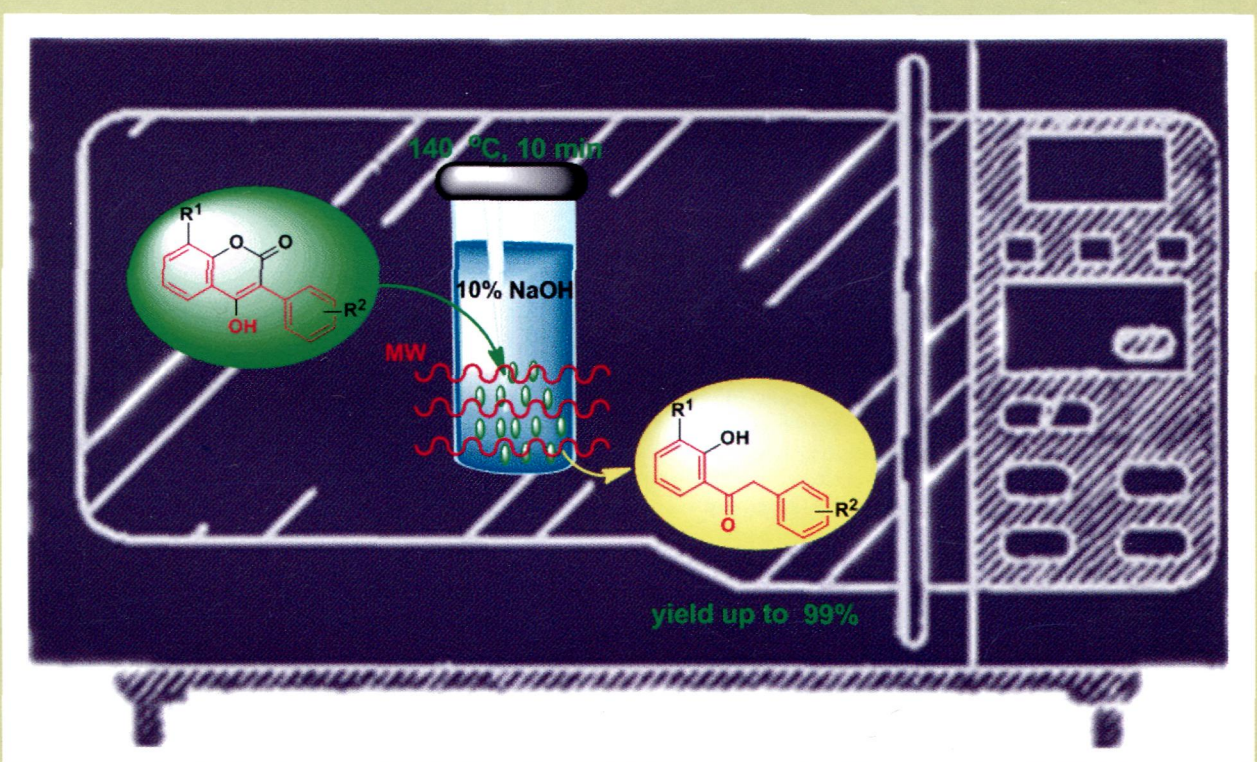

Microwave-Assisted Efficient Synthesis of 2-Hydroxydeoxybenzoins from the Alkali Degradation of Readily Prepared 3-Aryl-4-hydroxycoumarins in Water *pp. 1166–1172*

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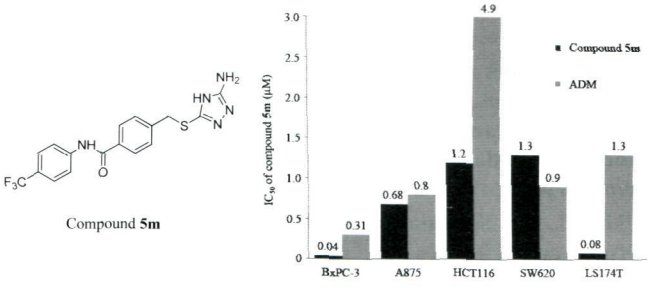
Vol. 61, No. 11 November 2013

Contents

Regular Articles

Synthesis and Biological Evaluation of 1,2,4-Triazole and 1,3,4-Thiadiazole Derivatives as Potential Cytotoxic Agents

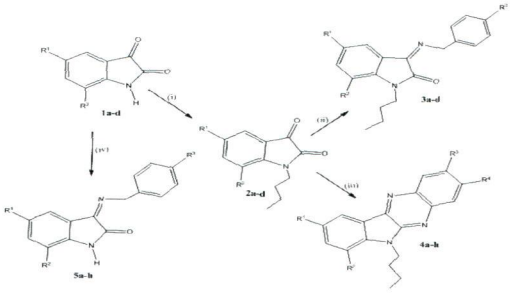
Y.-J. Shi, X.-J. Song, X. Li, T.-H. Ye, Y. Xiong, and L.-T. Yu



pp. 1099–1104

Synthesis and Evaluation of the Cytotoxic Activities of Some Isatin Derivatives

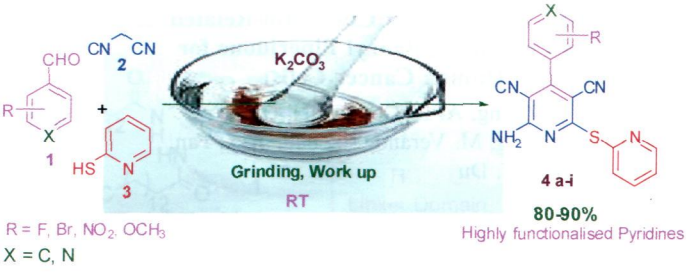
S. S. Reddy, R. Pallela, D.-M. Kim, M.-S. Won, and Y.-B. Shim



pp. 1105–1113

An Efficient Green Multi-Component Reaction Strategy for the Synthesis of Highly Functionalised Pyridines and Evaluation of Their Antibacterial Activities

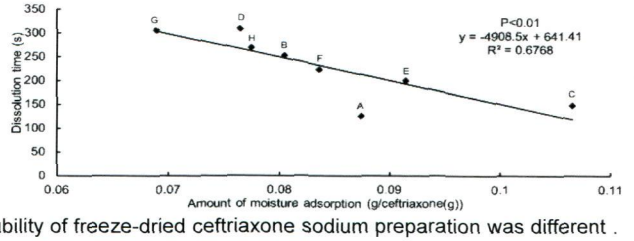
L. Srinivasula Reddy, T. Ram Reddy, R. B. Mohan, A. Mahesh, Y. Lingappa, and N. C. Gangi Reddy



pp. 1114–1120

Comparison of the Dissolution Rate of Ceftriaxone Sodium Preparations for Injection

M. Tange, Y. Hattori, M. Otsuka, M. Yoshida, J. Haginaka, and T. Uchida

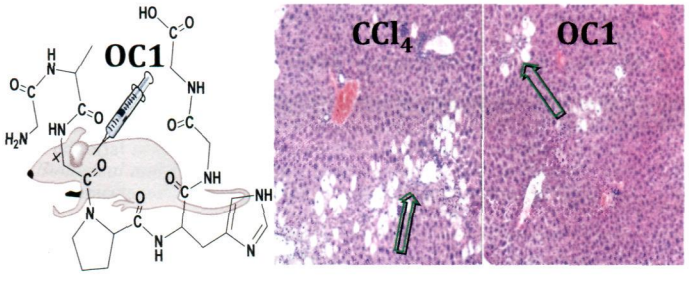


Solubility of freeze-dried ceftriaxone sodium preparation was different . A significant correlation was found between the amount of water adsorbed and the dissolution time.

pp. 1121–1129

Protection of a New Heptapeptide from *Carapax trionycis* against Carbon Tetrachloride-Induced Acute Liver Injury in Mice

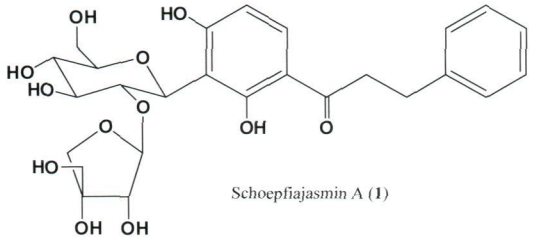
P. Wang, Y. Zhang, Y. An, K. Xu, X. Xu, C. Fu, J. Lin, S. Xu, Q. Li, and H. Lei



pp. 1130–1135

Schoepfiajasmins A–H: *C*-Glycosyl Dihydrochalcones, Dihydrochalcone Glycoside, *C*-Glucosyl Flavanones, Flavanone Glycoside and Flavone Glycoside from the Branches of *Schoepfia jasminodora*

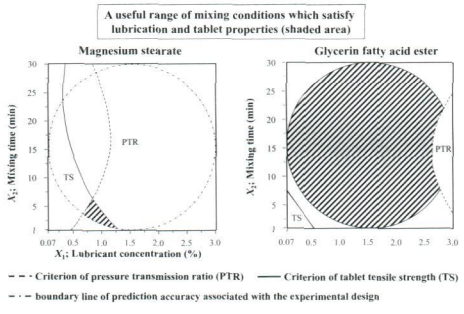
K. Ukida, T. Doi, S. Sugimoto, K. Matsunami, H. Otsuka, and Y. Takeda



pp. 1136–1142

Determination of Useful Ranges of Mixing Conditions for Glycerin Fatty Acid Ester by Multiple Regression Analysis

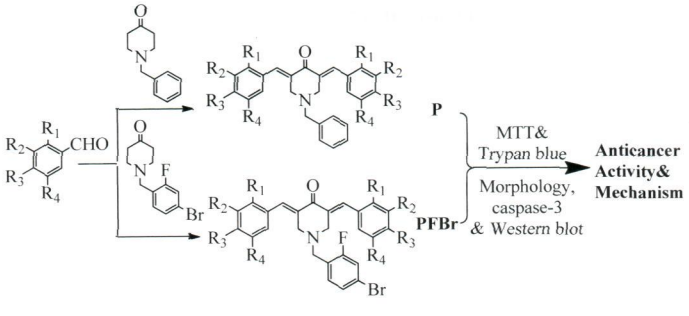
T. Uchimoto, Y. Iwao, H. Hattori, S. Noguchi, and S. Itai



pp. 1143–1148

Synthesis and Evaluation of Curcumin-Related Compounds Containing Benzyl Piperidone for Their Effects on Human Cancer Cells

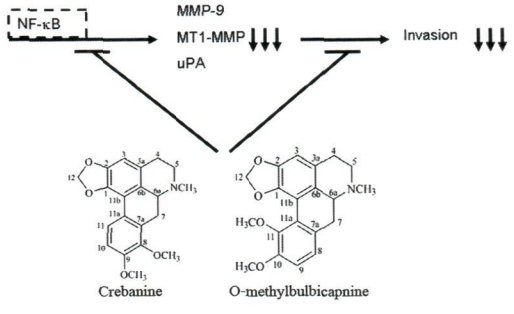
D.-Y. Zhou, K. Zhang, A. H. Conney, N. Ding, X.-X. Cui, H. Wang, M. Verano, S. Zhao, Y.-X. Fan, X. Zheng, and Z.-Y. Du



pp. 1149–1155

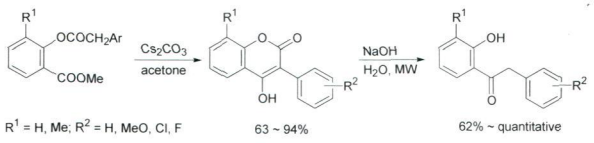
Anti-invasion Effect of Crebanine and *O*-Methylbulbocapnine from *Stephania venosa via* Down-Regulated Matrix Metalloproteinases and Urokinase Plasminogen Activator

S. Yodkeeree, P. Wongsirisin, W. Pompimon, and P. Limtrakul



pp. 1156–1165

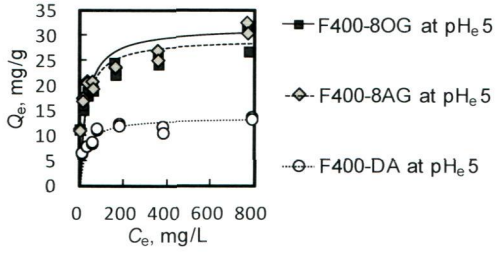
Microwave-Assisted Efficient Synthesis of 2-Hydroxydeoxybenzoins from the Alkali Degradation of Readily Prepared 3-Aryl-4-hydroxycoumarins in Water
Z.-Z. Zhou, G.-H. Yan, W.-H. Chen, and X.-M. Yang



pp. 1166–1172

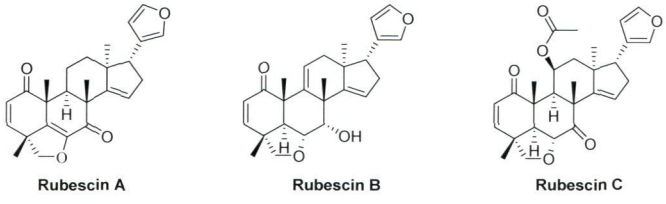
Notes

Effect of Surface Property of Activated Carbon on Adsorption of Nitrate Ion
T. Iida, Y. Amano, M. Machida, and F. Imazeki



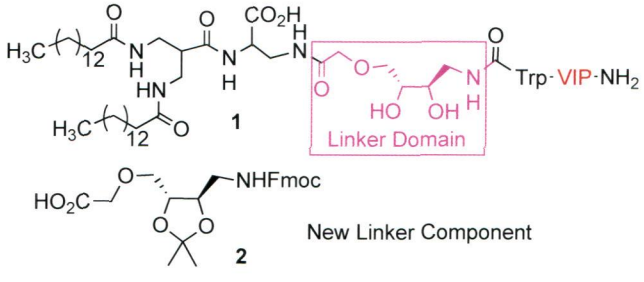
pp. 1173–1177

Rubescins A, B and C: New Havanensin Type Limonoids from Root Bark of *Trichilia rubescens* (Meliaceae)
A. Tsamo Tontsa, P. Mkounga, F. N. Njayou, J. Manautou, M. Kirk, P. G. Hultin, and A. E. Nkengfack



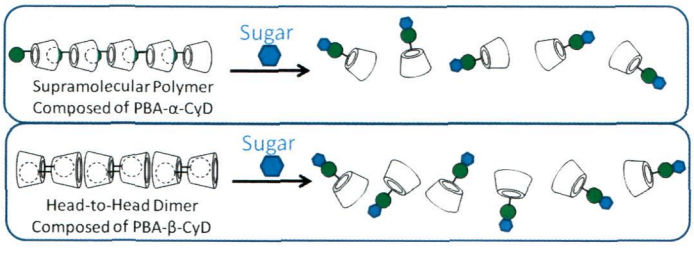
pp. 1178–1183

Synthesis of VIP-Lipopeptide Using a New Linker to Modify Liposomes: Towards the Development of a Drug Delivery System for Active Targeting
T. Masaka, T. Matsuda, Y. Li, Y. Koide, A. Takami, K. Yano, R. Imai, R. Ichihara, N. Yagi, H. Suzuki, H. Hikawa, K. Terada, and Y. Yokoyama



pp. 1184–1187

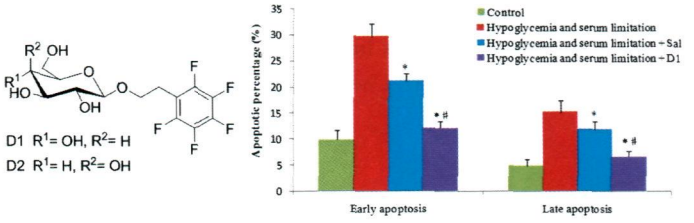
Sugar-Sensitive Supramolecular Structures Based on Phenylboronic Acid-Modified Cyclodextrins
K. Nakamura, T. Seki, Y. Egawa, R. Miki, Y. Oda, T. Yamanoi, and T. Seki



pp. 1188–1191

Synthesis and Neuroprotective Effects of the Fluorine Substituted Salidroside Analogues in the *PC12 Cell Model Exposed to Hypoglycemia and* Serum Limitation

X. Yang, W. Xu, W. Zhao, Y. Zhao, Y. Yang, and Y. Ling



pp. 1192–1196

About the cover: An operationally simple, green and efficient approach for the synthesis of 2-hydroxydeoxybenzoins bearing diverse substituents was developed by the microwave-assisted alkali degradation of 3-aryl-4-hydroxycoumarins in water. The latter compounds were readily prepared from the intramolecular Claisen condensation reaction of methyl 2-(2-arylacetoxy)benzoates in the presence of Cs_2CO_3/acetone, in excellent yields without laborious workup procedures. This method is highly atom-economic and applicable for the large-scale synthesis of 2-hydroxydeoxybenzoins. See the article by Zhou *et al.* on page 1166 of this issue.