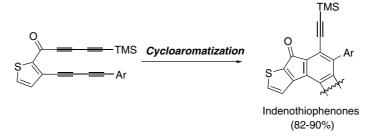


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R
OH
OH
OR
$$R^1CO_2H$$
 μ
 μ
 R^2
 R^2
 R^2
 R^3
 R^2
 R^2
 R^3
 R^4
 R^2
 R^2
 R^3
 R^4
 R^4



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Masashi Hamaguchi,* Naoki Tomida, Eiko Mochizuki and Takumi Oshima

$$\begin{bmatrix} Ar \\ C=O \\ R \end{bmatrix} \xrightarrow{Ar \\ N=N} \begin{bmatrix} CO_2R' \\ N=N \end{bmatrix} \xrightarrow{R'OOC} Ar \\ N=N \\ N-CO_2R' \\ R \end{bmatrix}$$

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Bor-Cherng Hong,* Zhong-Yi Chen, Arumugam Nagarajan, Kottani Rudresha, Vishal Chavan, Wei-Hung Chen, Yea-Fen Jiang, Shuo-Cang Zhang, Gene-Hsiang Lee and Sepehr Sarshar

$$O \cap R$$
, $O \cap R$, O

Straightforward synthesis of enantiopure 2-aminomethyl and 2-hydroxymethyl pyrrolidines with complete stereocontrol

pp 1287-1290

Marco Marradi, Stefano Cicchi, J. Ignacio Delso, Luca Rosi, Tomas Tejero, Pedro Merino* and Andrea Goti*

A practical synthesis of 2-aminomethyl- and 2-hydroxymethyl-3,4-dihydroxypyrrolidines via stereocontrolled addition of TMSCN and LiCH₂OMOM to chiral 3,4 dihydro-2*H*-pyrroline *N*-oxides is reported.

Podospermic acid, 1,3,5-tri-*O*-(7,8-dihydrocaffeoyl)quinic acid from *Podospermum laciniatum* (Asteraceae)

pp 1291-1294

Christian Zidorn,* Bent O. Petersen, Vedrana Udovičić, Thomas O. Larsen, Jens Ø. Duus, Judith M. Rollinger, Karl-Hans Ongania, Ernst P. Ellmerer and Hermann Stuppner

A phytochemical investigation of *Podospermum laciniatum* (L.) DC. (Asteraceae) yielded the new quinic acid derivative podospermic acid (1,3,5-tridihydrocaffeoylquinic acid), which was named after the genus it was isolated from. The structure was established by HR mass spectrometry and extensive 1D and 2D NMR spectroscopy. Podospermic acid is the first naturally occurring dihydrocaffeoylquinic acid derivative. The chemosystematic impact and the radical scavenging activity of the new compound are discussed briefly.

Amino acid binding in water by a new guanidiniocarbonyl pyrrole dication: the effect of the experimental conditions on complex stability and stoichiometry

pp 1295-1298

Carsten Schmuck* and Svea Graupner

The synthesis and binding properties of a new guanidiniocarbonyl pyrrole dication 2 are reported, which efficiently binds alanine carboxylate 7 in water with $\log K_{\rm ass} = 3.9$. However, the experimental conditions for determining the binding constant significantly influence both complex stability and stoichiometry.



A mild and efficient synthesis of α -tosylamino ketones from aryl aziridines in the presence of β -cyclodextrin and NBS in water

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M. Somi Reddy, M. Narender and K. Rama Rao*

Highly selective mono-substitution in Pd-catalyzed cross-coupling reactions of 3,6-dichloropyridazine with organozinc compounds

pp 1303-1305

Dmitriy S. Chekmarev, Alexander E. Stepanov and Alexander N. Kasatkin*

$$\begin{array}{c|c}
CI & R^1ZnX & R^2ZnX \\
Pd(PPh_3)_4 & CI & R^2ZnX
\end{array}$$

$$\begin{array}{c|c}
R^1 & R^2ZnX \\
Pd(PPh_3)_4 & R^2ZnX
\end{array}$$

$$\begin{array}{c|c}
R^1 & R^2 = \text{aryl, benzyl, alkyl}
\end{array}$$

A radical mediated first total synthesis from 'diacetone glucose' and determination of the absolute stereochemistry of xylobovide

pp 1307-1309

G. V. M. Sharma* and T. Gopinath

The first total synthesis by an intramolecular radical cyclisation protocol on a carbohydrate derived 5-hexynyl system, and determination of the absolute stereochemistry of xylobovide are reported.

Amberlyst-15® as a novel and recyclable solid acid for the coupling of aromatic aldehydes with homopropargyl alcohol

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Directed H-bonding inhibition in molecular recognition: an NMR case study of the H-bonding of a dicarboxylic acid with a new mixed diamide receptor having one adjacent pyridine-N-oxide Shyamaprosad Goswami,* Swapan Dey, Annada C. Maity and Subrata Jana

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of receptor 1 with adipic acid

The application of vinamidinium salts to the synthesis of 2,4-disubstituted thiophenes Ryan T. Clemens and Stanton Q. Smith*

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Total synthesis of pawhuskin C: a directed *ortho* **metalation approach** Jeffrey D. Neighbors, Maya S. Salnikova and David F. Wiemer*

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A convenient synthesis of 2-nitroindoles

Sujata Roy and Gordon W. Gribble*

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Acetate aldol reactions of chiral oxocarbenium ions

Sandeep Kanwar and Sanjay Trehan*

pp 1329-1332

Acetate aldol reactions of chiral oxocarbenium ions have been investigated and give the corresponding products with high diastereoselectivity in a predictable manner.

Design and characterisation of novel hexadentate 3-hydroxypyridin-4-one ligands

pp 1333-1336

Sirivipa Piyamongkol, Tao Zhou, Zu D. Liu, Hicham H. Khodr and Robert C. Hider*

One-pot, four-component reaction of isocyanides, dimethyl acetylenedicarboxylate, and cyclobutene-1,2-diones: a synthesis of novel spiroheterocycles

pp 1337-1339

Vijay Nair,* Rajeev S. Menon, Ani Deepthi, B. Rema Devi and A. T. Biju

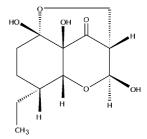
Isocyanides, dimethyl acetylenedicarboxylate, and cyclobutene-1,2-diones react in one-pot to afford novel spirocyclic compounds with double insertion of the isocyanide.

An anti-herpes simplex virus-type 1 agent from Xylaria mellisii (BCC 1005)

pp 1341-1344

Pattama Pittayakhajonwut,* Rapheephat Suvannakad, Surang Thienhirun, Samran Prabpai, Palangpon Kongsaeree and Morakot Tanticharoen

A structurally unique polyketide, mellisol and 1,8-dihydroxynaphthol 1-O- α -glucopyranoside, have been isolated from the fungus *Xylaria mellisii*. The relative stereostructure of mellisol was determined on the basis of X-ray crystallographic data.





Solid supported synthesis of structurally diverse dihydropyrido[2,3-d]pyrimidines using microwave irradiation

pp 1345-1348

Anu Agarwal and Prem M. S. Chauhan*

A library of dihydropyrido[2,3-d]pyrimidines has been synthesized on solid support using microwave irradiation.

One-step synthesis of 2-arylbenzothiazole ('BTA') and -benzoxazole precursors for in vivo imaging of $\beta\text{-amyloid}$ plaques

pp 1349-1351

David Alagille, Ronald M. Baldwin and Gilles D. Tamagnan*

Selective alkylation and Suzuki coupling as an efficient strategy for introducing functional anchors to the ethylene-bis(indenyl) ligand

pp 1353-1356

Anthony P. Panarello, Oleksiy Vassylyev and Johannes G. Khinast*



A convenient method for the preparation of primary amines using tritylamine

pp 1357-1360

Vassiliki Theodorou,* Valentine Ragoussis, Alexandros Strongilos, Evangelos Zelepos, Argyro Eleftheriou and Maria Dimitriou

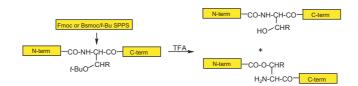
$$TrNH_2 + RX \longrightarrow TrNHR \xrightarrow{1. TFA} RNH_2 \cdot HC1 \qquad \text{where X: Cl, Br, I, OTs}$$

A convenient method for the preparation of primary amines from alkyl halides or alkyl tosylates by the use of tritylamine has been established. Primary amines were obtained in good yields by treating the *N*-tritylamines with trifluoroacetic acid.

Dramatically enhanced $N \rightarrow O$ acyl migration during the trifluoroacetic acid-based deprotection step in solid phase peptide synthesis

pp 1361-1364

Louis A. Carpino,* Eberhard Krause, Calin Dan Sferdean, Michael Bienert and Michael Beyermann*





Synthetic applications of the amine-base treatment in the ozonolysis of substituted-allyl silyl ethers or -allyl esters via a novel ene-diol type rearrangement

pp 1365-1368

Yung-Son Hon* and Ying-Chieh Wong

$$\begin{array}{c} O\Sigma \\ R \end{array} \begin{array}{c} 1. \ O_3 \ , \ CH_2Cl_2, \ -78 \ ^{o}C \\ 2. \ 1.2 \ equiv. \ Base \\ \end{array} \begin{array}{c} O\Sigma \\ R \end{array} \begin{array}{c} R = Aryl \quad Base = Et_3N \\ R = Alkyl \quad Base = DBU \\ \Sigma = R_3Si, \ Ac, \ Piv, \ Bz \end{array}$$

The ozonolysis of substituted-allyl silyl ethers or -allyl esters followed by treatment with bases gave the corresponding α -silyloxy ketones or α -acyloxy ketones in good yields. The reaction was proposed to proceed via a novel ene-diol rearrangement of the corresponding α -silyloxy aldehydes or α -acyloxy aldehydes intermediates.

Synthesis of flavanones using nanocrystalline MgO

pp 1369-1371

B. M. Choudary,* K. V. S. Ranganath, Jagajit Yadav and M. Lakshmi Kantam

Photoinduced Bergman cycloaromatization of imidazole-fused enediynes

pp 1373-1375

Zhengrong Zhao, Justin G. Peacock, Daniel A. Gubler and Matt A. Peterson*

A family of simple amide-derived air-stable P,O-ligands for Suzuki cross-coupling of unactivated aryl chlorides

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Wei-Min Dai* and Ye Zhang

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*Corresponding author

(1) Supplementary data available via ScienceDirect

COVER

The first asymmetric total synthesis of a marine natural product, methyl sarcoate, has been achieved featuring the asymmetric Michael addition, the dithiane coupling, the Kosugi-Migita-Stille coupling, and the ring-closing metathesis. *Tetrahedron Letters* **2005**, *46*, 1263–1267.

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