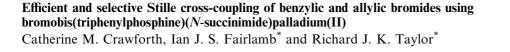
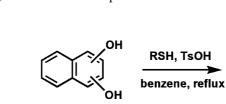
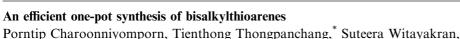


Ian R. Butler,^{*} Alfonso G. Callabero, Glenn A. Kelly, Jennifer R. Amey, Tobias Kraemer, Dafydd A. Thomas, Mark E. Light, Thomas Gelbrich and Simon J. Coles^{*}









Tetrahedron Letters Vol. 45, No. 3, 2004

Contents

SR

SR

Porntip Charoonniyomporn, Tienthong Thongpanchang,^{*} Suteera Witayakran, Yodhathai Thebtaranonth, Karen E. S. Phillips and Thomas J. Katz

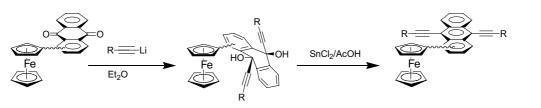


COMMUNICATIONS



pp 457-459

pp 461-465



 $Pd(NCOC_2H_4CO)(PPh_3)_2Br$ (5 mol%) Pr $R^{"SnBu_3}$ Toluene,
60 °C

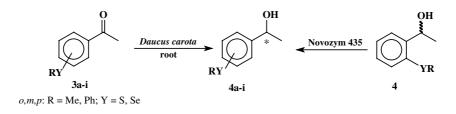
as above

pp 467–472

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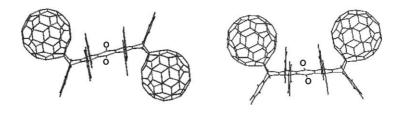
Preparation of chiral organochalcogeno- α -methylbenzyl alcohols via biocatalysis. The role of *Daucus carota* root

João V. Comasseto, Álvaro T. Omori, André L. M. Porto and Leandro H. Andrade*

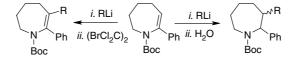


Probing the spatial requirements for [60]fullerene–[60]fullerene π -stacking and the *syn* addition pp 477–481 of [60]fullerenes across acenes

Glen P. Miller^{*} and Jonathan Briggs

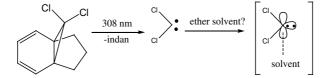


Carbolithiation of ene-carbamates. Application to the synthesis of 2,3-disubstituted ene-carbamates pp 483–484 Franck Lepifre, Bertrand Cottineau, Deborah Mousset, Pascal Bouyssou and Gerard Coudert^{*}



A search for dichlorocarbene ether solvent interactions

Stanislav I. Presolski, Adelajda Zorba, Dasan M. Thamattoor,^{*} Eric M. Tippmann and Matthew S. Platz^{*}



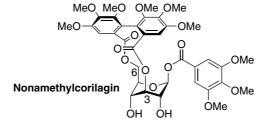
Interactions between dichlorocarbene and ether solvents were probed using laser flash photolysis techniques.

pp 473-476



The first construction of a 3,6-bridged ellagitannin skeleton with ${}^{1}C_{4}/B$ glucose core; synthesis of nonamethylcorilagin

Yasunori Ikeda, Kohei Nagao, Koki Tanigakiuchi, Go Tokumaru, Hitoshi Tsuchiya and Hidetoshi Yamada*



Synthesis of L-α-amino-ω-bromoalkanoic acid for side chain modification Louis A. Watanabe, Binoy Jose, Tamaki Kato, Norikazu Nishino* and Minoru Yoshida

Base-labile tert-butoxycarbonyl (Boc) group on phenols

Kozo Nakamura,* Takero Nakajima, Hiroshi Kayahara, Eisaku Nomura and Hisaji Taniguchi

BocOPh

can avoid side reactions during the deprotection with acids. We note the lability of the Boc to bases and are able to utilize it as a new cleavage condition for synthetic studies.

Phenols are deprotected with weak bases from their tert-butoxycarbonyl (Boc) derivatives. Boc deprotection with bases

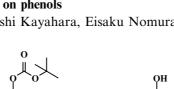
Asymmetric enolate alkylation via templation with chiral synthetic receptors Brenda J. Postnikova and Eric V. Anslyn*

[enolate]

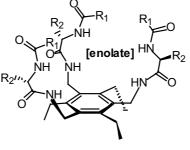
pp 495-499

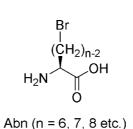
pp 501-504

447



PhOH





pp 491-494

Solution structure of bis(acetoxy)iodoarenes as observed by ¹⁷O NMR spectroscopy Giovanni Cerioni^{*} and Gianluca Uccheddu

The 17 O NMR spectra of the title compounds show equivalence for all the oxygens in CDCl₃ solution. The possibility of an ion pair structure is discussed.

N(iPr)2

Synthesis of a formamidine-protected 5'-amino-2',5'-dideoxyguanosine phosphoramidite and preparation of 5'-acylamidooligonucleotides

Jan A. Rojas Stütz and Clemens Richert*

from 2-*N*-(dibutylformamidino)-2'-deoxyguanosine via a three-pot route.

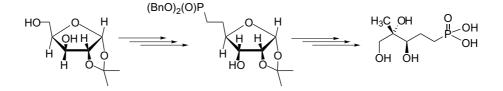
A phosphoramidite of 5'-amino-2',5'-dideoxyguanosine suitable for automated DNA synthesis was prepared

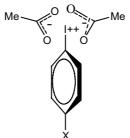
Hoda El Bitar, Van Hung Nguyen, Anthony Gramain, Thierry Sévenet and Bernard Bodo*

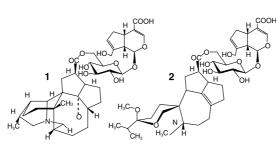
Daphcalycinosidines A and B, new iridoid-alkaloids from Daphniphyllum calycinum

The structures of daphcalycinosidines A (1) and B (2) were determined by spectral methods and chemical modifications.

(3*R*,4*S*)-3,4,5-Trihydroxy-4-methylpentylphosphonic acid, an isosteric phosphonate analogue pp 519–521 of 2-*C*-methyl-D-erythritol 4-phosphate, a key intermediate in the new pathway for isoprenoid biosynthesis Guillaume Hirsch, Catherine Grosdemange-Billiard, Denis Tritsch and Michel Rohmer^{*}







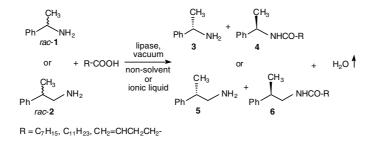
pp 505-507

pp 509-513



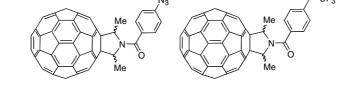
Lipase-catalyzed enantioselective reaction of amines with carboxylic acids under reduced pressure in non-solvent system and in ionic liquids

Roxana Irimescu and Katsuya Kato^{*}



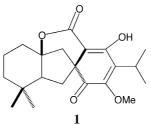
Synthesis of C₆₀ derivatives for photoaffinity labeling

Eiji Okada, Yuka Komazawa, Masaaki Kurihara, Hideshi Inoue, Naoki Miyata, Haruhiro Okuda, Toshie Tsuchiya and Yoko Yamakoshi^{*}



A trypanocidal diterpene with novel skeleton from Dracocephalum komarovi

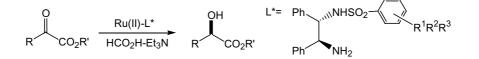
Nahoko Uchiyama, Michiho Ito, Fumiyuki Kiuchi, Gisho Honda,* Yoshio Takeda, Olimjon K. Khodzhimatov and Ozodbek A. Ashurmetov



A new diterpene, komarovispirone (1) with a spiro-octahydroindene skeleton, was isolated from Dracocephalum komarovi, which showed trypanocidal activity against epimastigote of Trypanosoma cruzi.

Transfer hydrogenation of activated ketones using novel chiral Ru(II)-N-arenesulfonyl-1,2-diphenylethylenediamine complexes

Damjan Sterk, Massoud S. Stephan and Barbara Mohar^{*}



pp 531-533

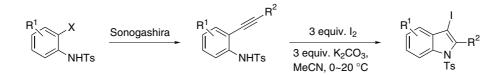
pp 535-537

pp 527-529

449

A simple, two-step synthesis of 3-iodoindoles Muhammad Amjad and David W. Knight^{*}

pp 539-541



Efficient Sonagashira couplings provide rapid access to *N*-tosylsulfonamides, which undergo smooth 5-*endo*-dig iodocyclisations to give good to excellent yields of potentially useful 3-iodoindoles.

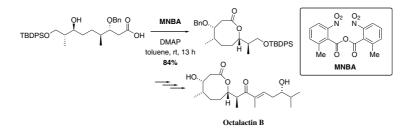
Asymmetric total synthesis of octalactin B using a new and rapid lactonization

Isamu Shiina,* Hiromi Oshiumi, Minako Hashizume, Yu-suke Yamai and Ryoutarou Ibuka

pp 543-547

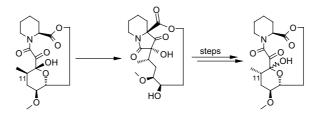
pp 549-551

pp 553-556



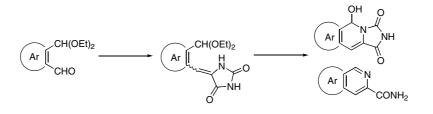
Selective transformation of ascomycin into 11-epi-ascomycin

Karl Baumann,^{*} Markus Bacher, Annelaure Damont and Andrea Steck

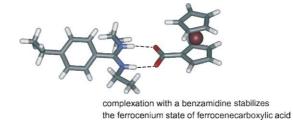


Synthesis of carbamoylpyridine and imidazo[1,5-a]pyridin-1,3-diones via ortho-acetalhydantoin intermediates

Jean M. Chezal, Emmanuel Moreau, Nicolas Desbois, Yves Blache, Olivier Chavignon and Jean C. Teulade*



Graeme Cooke,^{*} Florence M. A. Duclairoir, Arno Kraft, Georgina Rosair and Vincent M. Rotello



Synthesis of calix[4]arene library substituted with peptides at the upper rim

pp 561-564

Hideaki Hioki,^{*} Yumiko Ohnishi, Miwa Kubo, Emi Nashimoto, Yukinori Kinoshita, Miho Samejima and Mitsuaki Kodama

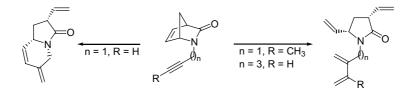
A fluorescence-labeled calix[4]arene library substituted with peptides at the upper rim was synthesized. Screening of the library for binding a dye-labeled oligopeptide indicated that some peptidocalix[4]arenes selectively bind the oligopeptide. The chemosensitivity of the library members for a target peptide was also investigated.

T : Tripeptide F : Fluorophore

3375 members

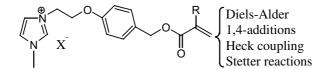
Control of product distribution in the domino metathesis reactions of *N*-alkynyl 2-azabicyclo[2.2.1]hept-5-en-3-ones. A convenient synthesis of functionalized γ-lactams and indolizidinones

Odón Arjona,* Aurelio G. Csákÿ, Vanessa León, Rocío Medel and Joaquín Plumet*



Synthesis and preliminary use of novel acrylic ester-derived task-specific ionic liquids Siddam Anjaiah, Srivari Chandrasekhar and René Grée^{*}

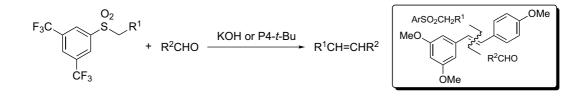
pp 569-571



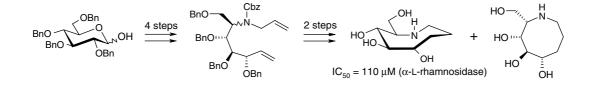
The synthesis and reactions of new task-specific ionic liquids (TSILs) 1 and 2 are described.

3,5-Bis(trifluoromethyl)phenyl sulfones in the modified Julia olefination: application to the synthesis of resveratrol

Diego A. Alonso, Carmen Nájera* and Montserrat Varea

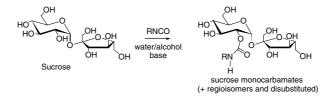


Synthesis and biological evaluation of the first example of an eight-membered iminoalditol pp 579–581 Guillaume Godin, Elodie Garnier, Philippe Compain,^{*} Olivier R. Martin,^{*} Kyoko Ikeda and Naoki Asano



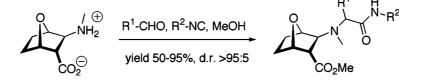
Preparation of amphiphilic sucrose carbamates by reaction with alkyl isocyanates in water-alcohol mixtures

Delphine Christian, Juliette Fitremann, Alain Bouchu and Yves Queneau*



U-4C-3CR versus U-5C-4CR and stereochemical outcomes using suitable bicyclic β -amino acid derivatives as bifunctional components in the Ugi reaction

Andrea Basso,^{*} Luca Banfi, Renata Riva and Giuseppe Guanti^{*}



Suitable bicyclic β -amino acids have been employed as bifunctional components in Ugi condensations. The effects of the *cis/trans* configuration and *N*-alkylation of the amino acid on product distribution and stereochemistry have been investigated, the *N*-alkylated *trans* derivative furnishing excellent control of the diastereoselectivity.

pp 583-586

pp 573–577

pp 587-590

Xyloccensin L, a novel limonoid from Xylocarpus granatum

Jun Wu,* Si Zhang, Qiang Xiao, Qingxin Li, Jianshe Huang, Lijuan Long and Liangmin Huang

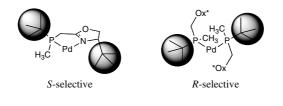
The isolation and structure elucidation of xyloccensin L from the stem bark of Xylocarpus granatum is described. Xyloccensin L is a highly oxidized heptacyclic A, B, D-seco limonoid with an α-8, 30-epoxy ring and a rare oxygen bridge between C1 and C29.

Effect of lithium chloride on allylation of zirconacyclopentadienes Lian Leng, Chanjuan Xi,* Chao Chen and Chunbo Lai

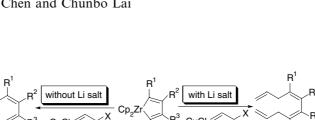
Efficient indole N-detosylation using thioglycolate Charlotte M. Haskins and David W. Knight*

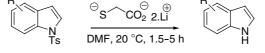
P-stereogenic P/N hybrid ligands: a remarkable switch in enantioselectivity in palladium-catalyzed asymmetric allylation

Hiroshi Danjo, Masato Higuchi, Mitsuhiro Yada and Tsuneo Imamoto*



pp 591-593





pp 595-598

pp 599-601

pp 603-606

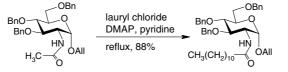
Regioselective synthesis of bis(silyl enol ethers) and bis(conjugated enones) through electron transfer pp 607–609 from Mg metal

Hirofumi Maekawa, Manabu Sakai, Tetsuro Uchida, Yoshio Kita and Ikuzo Nishiguchi*

$$\begin{array}{c} O \\ R^{1} \\ R^{2} \end{array} \xrightarrow{R^{3}} \begin{array}{c} Me_{3}SiCl / Mg / DMF \\ \hline Yield : 48 \sim 72\% \end{array} \xrightarrow{R^{2} R^{1} R^{1}} \\ Me_{3}SiO \\ R^{3} R^{3} OSiMe_{3} \end{array}$$

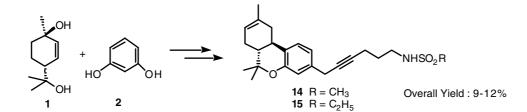
An unprecedented *N*-transacylation reaction on 2-acetamido-2-deoxy- α -D-glucopyranosides Yingxia Li,^{*} Chunxia Li, Peng Wang, Shidong Chu, Huashi Guan and Biao Yu^{*}

pp 611-613

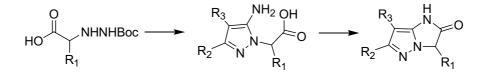


A novel methodology for the synthesis of 1-desoxy- Δ^8 -tetrahydrocannabinol (THC) analogues Hang Sun, Anu Mahadevan and Raj K. Razdan^{*}

pp 615-617



Solution phase synthesis of imidazo[1,2-b]pyrazol-2-one, an interesting 5,5-fused heterocyclic ring system pp 619–621 Benjamin E. Blass,^{*} Anil Srivastava, Keith R. Coburn, Amy L. Faulkner, John J. Janusz, James M. Ridgeway and William L. Seibel

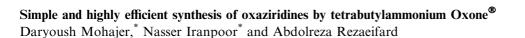


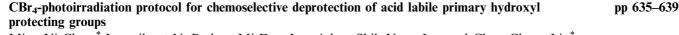
Prolinoamino acids as a tool to stabilize β -turns with the side chain of natural amino acids Jean Quancard, Philippe Karoyan, Olivier Lequin, Emmanuel Wenger, André Aubry, Solange Lavielle and Gérard Chassaing*



On the use of succinic anhydride as acylating agent for practical resolution of aryl-alkyl alcohols pp 627-630 through lipase-catalyzed acylation

Nassima Bouzemi, Hanane Debbeche, Louisa Aribi-Zouioueche* and Jean-Claude Fiaud*





Ming-Yi Chen,* Laxmikant N. Patkar, Mi-Dan Jan, Adam Shih-Yuan Lee and Chun-Cheng Lin*

-CH=N-CHa

100%

conversior

conversion

-CH=N 100%

 $HO \xrightarrow{OH} R \qquad HO \xrightarrow{OO} OR \\ (PO)_2 \qquad OR \\ HO \xrightarrow{O} B \qquad Fmoc. N \xrightarrow{R} OMe$ or TBDMS CBr₄/MeOH hv 20 examples



Et₂O + conv. 50% (±) (-) 92 % ee saponification (+) 92 % ee

-с́н-́м-сн

35%E, 57% Z

100% E

CBr₄-photoirradiation protocol for chemoselective deprotection of acid labile primary hydroxyl

Bu₄NHSO₅

CH₂CN, rt., <50 min

455

pp 631-634

Two novel 9,11-*seco*-11-norabietanes from the roots of *Taiwania cryptomerioides* Chiou-Feng Chyu, Yi-Ming Chiang, Hsiu-Chuan Lin and Yueh-Hsiung Kuo^{*}

Two novel 9,11-seco-11-norabietanes, namely taiwanlactones A (1) and B (2), were isolated from the roots of *Taiwania* cryptomerioides. The absolute configuration of 1 was elucidated by a modified Mosher's method. The biotransformation mechanisms of 1 and 2 were proposed.

OTHER CONTENTS

Contributors to this issue Instructions to contributors

*Corresponding author (*i*)⁺ Supplementary data available via ScienceDirect

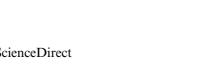
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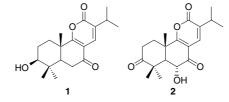
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CONTEN







pp 641-643

p I pp III–V