

3: isosteviol

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1: serofendic acid A

2: serofendic acid B



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The reaction of the carbenoid generated from 3-diazo-4-hydroxycoumarin with alcohols was run with the title catalyst, obtaining moderate to excellent yields of the corresponding adducts.

Observations on the regioselectivity of some Baeyer–Villiger Tetrahedron Letters 43 (2002) 3641 reactions Michael Harmata* and Paitoon Rashatasakhon Department of Chemistry, University of Missouri-Columbia, Columbia, MO 65211, USA



McMurry intermolecular cross-coupling between an ester and a ketone: scope and limitations

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Scope and limitations of an unusual McMurry cross-coupling reaction between a benzilic ester and a hindered ketone is described. \sim



Enantioselective reduction of aryl ketones using $LiBH_4$ and TarB-X: a chiral Lewis acid

Tetrahedron Letters 43 (2002) 3649

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A high yielding preparation of α -trimethylsilyloxyphosphonates by silylation of α -hydroxyphosphonates with HMDS catalyzed by iodine

Tetrahedron Letters 43 (2002) 3653

n= 1 to 28

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A general, versatile, high yielding and convenient procedure for the immediate conversion of various α -hydroxyphosphonates to α -trimethylsilyloxyphosphonates under neutral conditions using HMDS in the presence of a catalytic amount of iodine is described.





Diazo coupling: an alternative method for the upper rim amination of thiacalix[4]arenes

Tetrahedron Letters 43 (2002) 3665

Pavel Lhoták,* Jiri Morávek and Ivan Stibor

Department of Organic Chemistry, Institute of Chemical Technology, Technická 5, 166 28 Prague 6, Czech Republic The direct nitration of thiacalix[4]arene is not possible due to concomitant oxidation of the sulphur bridges. Alternatively, thiacalixarene reacts smoothly with diazonium salts to form tetrasubstituted azo derivatives that can be reduced to give upper rim amino-substituted thiacalixarene derivatives.







2-(3-Aminopropyl)-4-pentenoic acid as a bio-compatible/cleavable linker for solid-phase organic synthesis

Tetrahedron Letters 43 (2002) 3677

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2-(3-Aminopropyl)-4-pentenoic acid lithium salt (1) was prepared and used as a bio-compatible, cleavable linker in solid-phase organic synthesis. The products were released from solid-phase through cycloelimination.



Short solution phase preparation of fused azepine-tetrazoles via a UDC (Ugi/de-Boc/cyclize) strategy

Tetrahedron Letters 43 (2002) 3681

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This letter reveals a novel application of the TMSN3 modified Ugi reaction for the solution phase synthesis of fused azepine-tetrazole libraries. \mathbf{R} , \mathbf{R} , \mathbf{P} , \mathbf{O} , \mathbf{H} , \mathbf{R} .





Lanthanum metal-assisted deoxygenative coupling of alcohols

Tetrahedron Letters 43 (2002) 3689

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$$\begin{array}{ccc} \text{R-OH} & + & \text{La} & \xrightarrow{\text{Me}_3 \text{SiCl}/^{\text{cat.}} \text{I}_2/^{\text{cat.}} \text{Cul}} & \text{R-R} \\ \textbf{1} & & \text{CH}_3 \text{CN} & \textbf{2} \end{array}$$



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Tetrahedron Letters 43 (2002) 3715

The Cp₂TiMe₂-catalyzed intramolecular hydroamination/cyclization of aminoalkynes

Igor Bytschkov and Sven Doye*

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Unusual oxidative rearrangement of 1,5-diazadecalin

Tetrahedron Letters 43 (2002) 3747

Xiaolin Li, Zhenrong Xu, Erin F. DiMauro and Marisa C. Kozlowski*

Department of Chemistry, Roy and Diana Vagelos Laboratories, University of Pennsylvania, Philadelphia, PA 19104, USA

Upon treatment with $(PhIO)_n$ or $PhI(OAc)_2$, 1,5-diaza-*cis*-decalin undergoes oxidation at the more hindered position followed by Grob type fragmentation to yield the ring-expanded bislactam.





Preparation and conversion of chiral *O*-isopropylidene-protected 4-aminocyclohexenol to various key intermediates toward narcissus alkaloids Tetrahedron Letters 43 (2002) 3757

Shanmugham Elango, Ying-Chuan Wang, Chien-Liang Cheng and Tu-Hsin Yan*

Department of Chemistry, National Chung-Hsing University, Taichung 400, Taiwan, Republic of China











Allylic strain effects on the stereochemistry of the alkylation reaction of mycophenolic acid chiral enolates

Tetrahedron Letters 43 (2002) 3777

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An attempted enantioselective synthesis of all the four individual diastereomers of α , β -dimethylated mycophenolic acid using Evans chiral auxiliaries is reported. The 1,3-allylic strain effect on the enolates used favor the formation of the *syn* acids.



Aryl epoxide-halohydrin transformations: stereochemistry of reactions of aryl epoxides with lithium halide-acetic acid reagent

Tetrahedron Letters 43 (2002) 3781

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