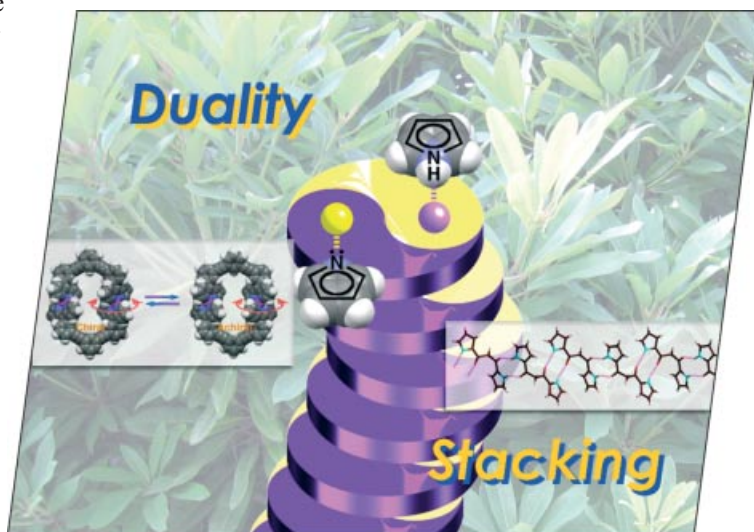




The EUChemSoc Societies have taken the significant step into the future by merging their traditional journals, to form two leading chemistry journals, the *European Journal of Inorganic Chemistry* and the *European Journal of Organic Chemistry*. Three further EUChemSoc Societies (Austria, Czech Republic and Sweden) are Associates of the two journals.

COVER PICTURE

The cover picture shows the two representative characteristics of pyrrole, “duality” as a hydrogen-bonding acceptor (or a metal-coordination ligand) N and a hydrogen-bonding donor (or an anion-binding ligand) NH and “planarity” to form stacking assemblies, depicted by the disks of Yin and Yang. Acyclic π -conjugated oligopyrrole derivatives, though less extensively studied so far, often potentially have even more advantages than cyclic ones. For example, the linear oligopyrroles prepared by the author form infinite and discrete coordination oligomers to give emissive spherical colloids and nanorings, hydrogen-bonding supramolecular assemblies to afford unique crystalline morphologies, and stacking structures to yield organogels that can be controlled by anions. Details are presented in the Microreview by H. Maeda on p. 5313ff. This picture has been provided by Mr. Takashi Hashimoto, one of the members in the author’s group.



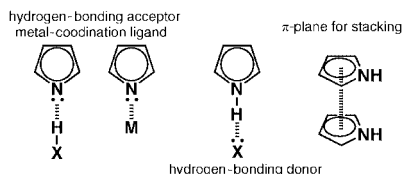
MICROREVIEW

Supramolecular Chemistry

H. Maeda* 5313–5325

Supramolecular Chemistry of Acyclic Oligopyrroles

Keywords: Hydrogen bonding / Metal coordination / π -Conjugation / Pyrroles / Supramolecular chemistry



Acyclic π -conjugated oligopyrrole derivatives, though not extensively studied, often have more advantages as anion receptors and metal-coordination ligands than their cyclic counterparts. They can be used to form versatile complexes, supramolecular assemblies, and nanoscale materials.

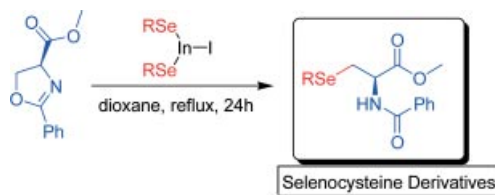
SHORT COMMUNICATIONS

Selenocysteine Derivatives

A. L. Braga,* F. Vargas, F. Z. Galetto,
M. W. Paixão, R. S. Schwab,
P. S. Taube 5327–5331

One-Pot Indium Iodide Mediated Synthesis of Chiral β -Seleno Amides and Selenocysteine Derivatives by Ring-Opening Reaction of 2-Oxazolines

Keywords: Selenium / Nitrogen heterocycles / Oxygen heterocycles / Selenocysteine / Indium



β -Seleno amides and selenocysteine derivatives were synthesized by a straightforward method involving a ring-opening reaction

of 2-oxazolines by indium selenolates that was generated in situ.

Iron–Palladium Association

V. Terrasson, J. Michaux, A. Gaucher,
J. Wehbe, S. Marque, D. Prim,*
J.-M. Campagne* 5332–5335



Iron–Palladium Association in the Preparation of Indoles and One-Pot Synthesis of Bis(indolyl)methanes

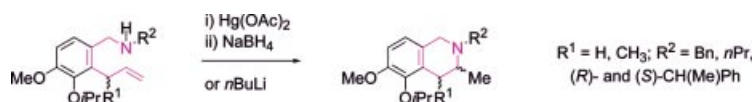
Keywords: Iron / Palladium / Indole / Bis(indolyl)methane



Indoles were prepared by annulation of alkynylanilines with the use of a new FeCl₃–PdCl₂ catalytic combination. One pot access to bis(indolyl)methanes and

trisubstituted indoles through annulation/Friedel–Crafts alkylation and annulation/1,4-Michael addition are described.

FULL PAPERS



The synthesis of 3-methyl- and 3,4-dimethyltetrahydroisoquinolines from aromatic aminoalkenes, using intramolecular hydroamination by reductive aminomercuration or sub-stoichiometric amounts of *n*-butyllithium, is reported.

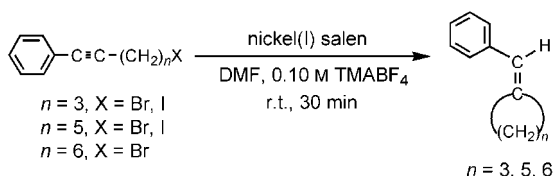
droamination by reductive aminomercuration or sub-stoichiometric amounts of *n*-butyllithium, is reported.

Intramolecular Hydroamination

R. Pathak, P. Naicker, W. A. Thompson, M. A. Fernandes, C. B. de Koning, W. A. L. van Otterlo* 5337–5345

Intramolecular Hydroaminations Mediated by Reductive Mercuration or *n*-Butyllithium To Afford 3-Methyl- and 3,4-Dimethyl-1,2,3,4-tetrahydroisoquinolines

Keywords: Hydroamination / Tetrahydroisoquinolines / Reductive mercuration / *n*-Butyllithium



Phenyl-conjugated acetylenic halides (e.g. 7-bromo- or 7-iodo-1-phenyl-1-heptyne) are catalytically reduced by nickel(I) salen electrogenerated at a glassy carbon cathode

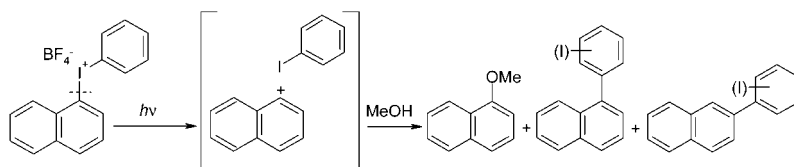
to afford benzylidenecycloalkanes, along with diphenylalkadiynes, alkynes, and alkenynes.

Synthesis of Benzylidenecycloalkanes

M. S. Mubarak,* T. B. Jennermann, M. A. Ischay, D. G. Peters* ... 5346–5352

Catalytic Reduction of Phenyl-Conjugated Acetylenic Halides by Nickel(I) Salen: Cyclization versus Coupling

Keywords: Catalysis / Nickel / Acetylenic halides / Intramolecular cyclization / Radical coupling



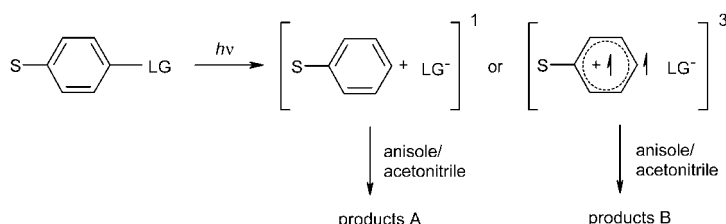
The reactivity of photogenerated naphthyl cations has been investigated with the aid of product studies and MO calculations.

Spin-Isomeric Naphthyl Cations

M. Slegt, F. Minne, H. Zuilhof, H. S. Overkleeft, G. Lodder* 5353–5363

Photochemical Generation and Reactivity of Naphthyl Cations: *cine* Substitution

Keywords: Photochemistry / Naphthyl cations / Spin multiplicity / *cine* substitution/ Ab initio calculations



The reactivities of singlet and triplet phenyl cations were studied by use of their chemo-

selectivities and the regioselectivities of their attacks on methoxybenzene.

Spin-Selective Phenyl Cation Trapping

M. Slegt, H. S. Overkleeft, G. Lodder* 5364–5375

Fingerprints of Singlet and Triplet Phenyl Cations

Keywords: Photochemistry / Phenyl cations / Spin multiplicity / Regioselectivity / Chemoselectivity

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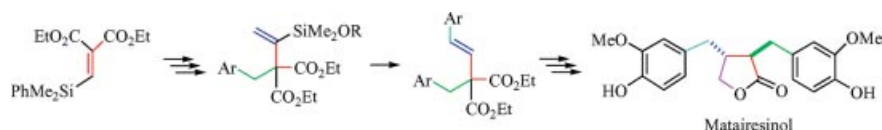
Dibenzylbutyrolactone Lignans

R. Singh, G. C. Singh,
S. K. Ghosh* 5376–5385



A Novel Approach Towards Dibenzylbutyrolactone Lignans Involving Heck and Radical Reactions: Application to (±)-Matairesinol Synthesis

Keywords: Olefination / Stereoselective synthesis / Radical cyclization / Dibenzylbutyrolactone / Matairesinol



Functionalized 1,4-diaryl butenes, precursors to the dibenzylbutyrolactone lignan skeleton, have been synthesized by a stereoselective Heck reaction of iodoarenes with in situ generated vinylated malonates. The

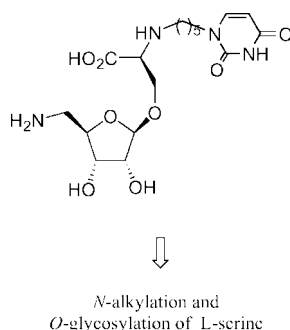
lactone ring has been constructed by a diastereoselective radical cyclization. This strategy has been applied in the synthesis of (±)-matairesinol.

Enzyme Inhibitors

L. Le Corre, C. Gravier-Pelletier,*
Y. Le Merrer* 5386–5394

Towards New *MraY* Inhibitors: A Serine Template for Uracil and 5-Amino-5-deoxy-ribose Scaffolding

Keywords: Translocase *MraY* inhibitors / Antibiotics / Glycosylation / Reductive amination / Amino acids



The bacterial translocase *MraY* is a good target for the development of new antibiotics since it is ubiquitous and essential for bacterial growth. The goal of this work was the synthesis of simplified analogues of natural inhibitors of this enzyme. The structure of the targeted compound retains the uracil and aminoribose parts linked by a serinyl template.

Asymmetric Catalysis

L. Eberhardt, D. Armspach,* D. Matt,*
L. Toupet, B. Oswald 5395–5403

Efficient, Rhodium-Catalyzed Hydrogenation of α -Dehydroamino Acid Esters with Chiral Monodentate Aminophosphanes Bearing Two Binaphthyl Groups

Keywords: Asymmetric catalysis / Hydrogenation / Rhodium / Phosphoramidites / Binaphthyl / Azepines / Binol



A chiral monodentate ligand containing a dinaphtho-fused 1,3,2-dioxaphosphine unit N–P-bonded to a dinaphtho-fused azepine ring system has been synthesised

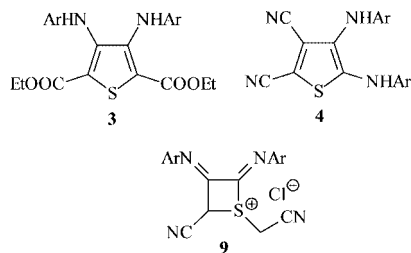
and tested in the hydrogenation of methyl 2-(acetylamino)-3-arylpropenoates; *ee* values of up to 99 % were obtained.

Push-Pull Thiophenes

G. Buehrdel, R. Beckert,* E. Birckner,
U.-W. Grummt, B. Beyer, S. Kluge,
J. Weston, H. Goerls 5404–5409

A Simple One-Pot Synthesis of Solvato-fluorescent Push-Pull Thiophenes

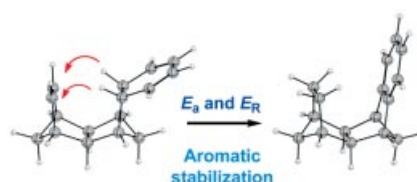
Keywords: Thiophene / Fluorescence / Bis-imidoyl chlorides / Cyclization reaction / Rearrangement




Cyclization of thiodiglycolate with bis-imidoyl chlorides yields the expected symmetric thiophenes of type **3**. Surprisingly, exchanging thiodiglycolate for thiodiacetonitrile yields asymmetric 2,3-diamino-4,5-dicyanothiophenes **4**, which show strongly solvent-dependent fluorescence (solvatofluorescence). The substitution pattern was confirmed by single-crystal X-ray analysis and possibly originates from a 1,2-rearrangement of a thietanium salt intermediate of type **9**.

Aromatic Stabilization Energies

The double group transfer reactions in *syn*-sesquinorbornanes, which involves the concomitant formation of an aromatic ring, are suggested as benchmarks for energetic aromaticity criteria.



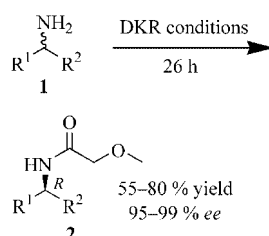
G. Frenking, F. P. Cossío, M. A. Sierra, I. Fernández* 5410–5415

Double Group Transfer Reactions as Indicators of Aromatic Stabilization 


Keywords: Aromaticity / Double group transfer reactions / Aromatic stabilization energy / Density functional calculations

Dynamic Kinetic Resolution

A modified version of the Bäckvall system for the dynamic kinetic resolution (DKR) of amines is reported, allowing complete conversion of various amine substrates with high *ee* within 26 h for most substrates.

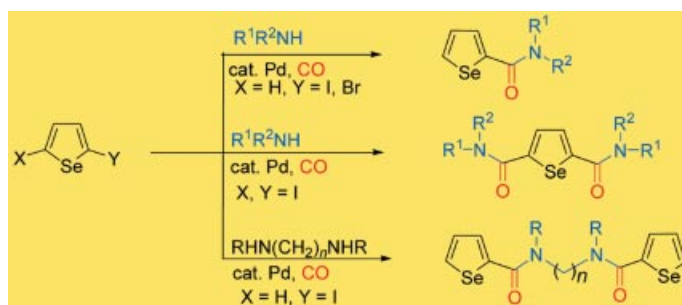


M. A. J. Veld, K. Hult, A. R. A. Palmans,* E. W. Meijer* 5416–5421


Fast DKR of Amines Using Isopropyl 2-Methoxyacetate as Acyl Donor 

Keywords: Dynamic kinetic resolution (DKR) / Amines / Enzyme catalysis / Asymmetric synthesis / Ruthenium

Pd-Catalyzed Carbonylation



P. Prediger, R. Brandão, C. W. Nogueira, G. Zeni* 5422–5428

Palladium-Catalyzed Carbonylation of 2-Haloselenophenes: Synthesis of Selenophene-2-carboxamides, Selenophene-2,5-dicarboxamides and *N,N'*-Bridged Selenophene-2-carboxamides 

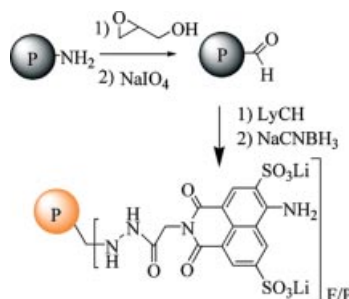
Keywords: Selenophene / Carbonylation / Palladium / Amides / Carbon monoxide

The palladium-catalyzed carbonylation reaction of haloselenophenes and amines under carbon monoxide atmosphere was performed. The description of a new route

to prepare selenophene-2-carboxamides, selenophene-2,5-dicarboxamides and *N,N'*-bridged selenophene-2-carboxamides is reported.

Bioconjugation Using Glycidol

The reactions between glycidol and either the nonglycosylated protein BSA or the glycoprotein avidin (AV), followed by mild oxidation of the intermediate glycol moieties, afforded reactive aldehyde functionalities. As an application, the functionalized proteins were derivatized with the fluorescent dye Lucifer Yellow CH, affording highly fluorescent bioprobes.



J.-M. Heldt, N. Fischer-Durand, M. Salmain, A. Vessières,* G. Jaouen 5429–5433

The Use of Glycidol to Introduce Aldehyde Functions Into Proteins – Application to the Fluorescent Labelling of Bovine Serum Albumin and Avidin

Keywords: Amination / Protein modifications / Lucifer Yellow CH / Fluorescent probes / Nucleophilic substitution

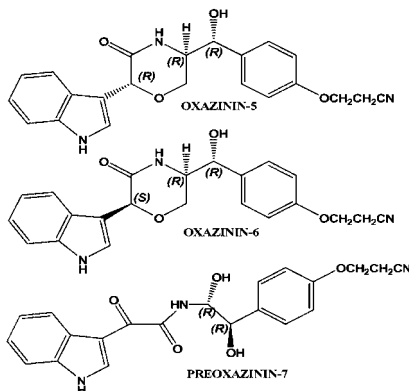
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Marine Biotoxins

P. Ciminiello,* C. Dell'Aversano,
E. Fattorusso,* M. Forino, L. Grauso,
F. U. Santelia, L. Tartaglione,
V. I. Moutsos, E. N. Pitsinos,
E. A. Couladouros 5434–5439

Stereostructural Determination by a Synthetic and NMR-Based Approach of Three Oxazinins Isolated from Adriatic Mussels

Keywords: Oxazinin / Marine biotoxins / Adriatic mussels / NMR spectroscopy / Synthetic methods



Two oxazinins, namely oxazinin-5 and -6, along with a related linear precursor (pre-oxazinin-7) were isolated from toxic mussels collected along the Northern Adriatic coasts. The planar structure of these novel compounds was achieved through extensive NMR spectroscopic analysis, whereas a synthetic approach was crucial for their absolute stereochemical elucidation.

CORRECTION

C. B. Singh, V. Kavala, A. K. Samal,
B. K. Patel* 5441

Aqueous-Mediated *N*-Alkylation of Amines

Keywords: Amines / Alkylation / Tertiary amines / Aqueous media / Surfactants

If not otherwise indicated in the article, papers in issue 31 were published online on October 16, 2007