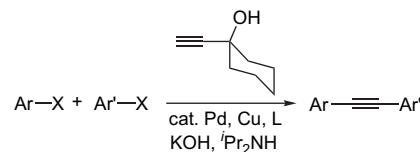


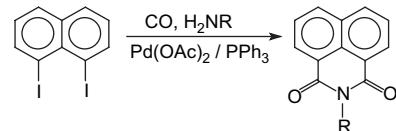
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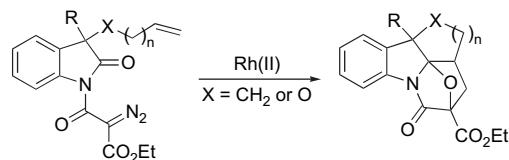
Ethynyl-cyclohexanol: an efficient acetylene surrogate in Sonogashira coupling pp 975–982
 Márton Csékei, Zoltán Novák and András Kotschy*



Facile synthesis of 1,8-naphthalimides in palladium-catalysed aminocarbonylation of 1,8-diido-naphthalene pp 983–987
 Attila Takács, Péter Ács and László Kollár*



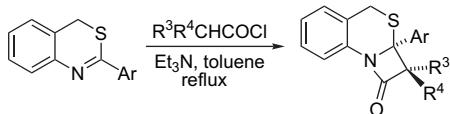
The rhodium(II) carbenoid cyclization–cycloaddition cascade of α -diazo dihydroindolinones for the synthesis of novel azapolycyclic ring systems pp 988–1001
 Dylan B. England, James M. Eagan, Gokce Merey, Olcay Anac and Albert Padwa*



Novel β -lactam condensed 3-thiaquinolines: an efficient synthesis and structural characterization

pp 1002–1011

Péter Csomós, Lajos Fodor,* Gábor Bernáth, Jari Sinkkonen, Jari Salminen, Kirsti Wiinamäki and Kalevi Pihlaja

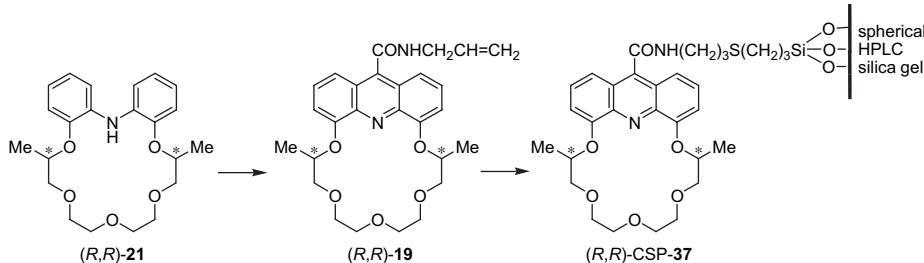


A series of novel monochloro-, dichloro-, and phenyl-substituted (R^3 or/and R^4) β -lactams condensed with 3,1-benzothiazines were obtained by Staudinger reaction and studied by NMR spectroscopy and mass spectrometry.

**Preparation of a new chiral acridino-18-crown-6 ether-based stationary phase for enantioseparation of racemic protonated primary aralkyl amines**

pp 1012–1022

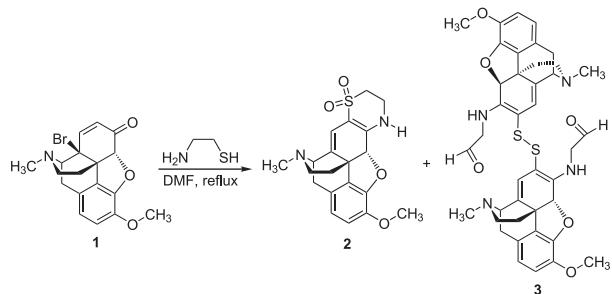
Szilvia Lakatos, József Fetter, Ferenc Bertha, Péter Huszthy,* Tünde Tóth, Viktor Farkas, György Orosz and Miklós Hollói

**Synthesis of 1,4-thiazino- and benzo-1,4-thiazinomorphinans: their acid-catalyzed rearrangement and study of the formation of unexpected oxidation products**

pp 1023–1028

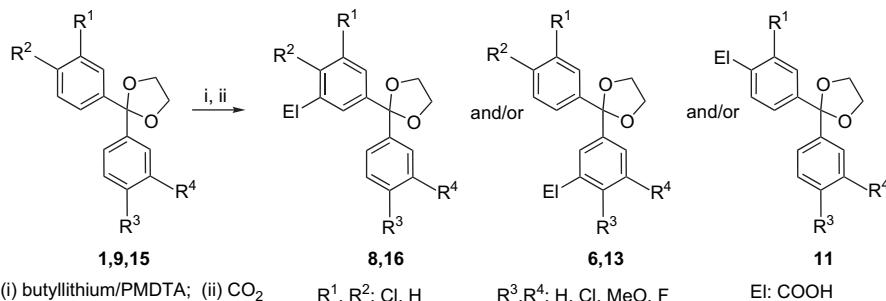
Attila Sipos,* Levente Girán, Harald Mittendorfer, Helmut Schmidhammer and Sándor Berényi

The formation of 1,4-thiazine and benzo-1,4-thiazine rings was performed at the 6,7-positions of the morphinan backbone in order to synthesize systems annulated with a new six-membered ring providing potential pharmacological activity and the opportunity of easy functionalization. An unexpected oxidation of cyclic sulfur was observed in both cases affording either sulfones or open-ringed bis-morphinan-type by-product.

**Lithiation of 2-(chloraryl)-2-aryl-1,3-dioxolanes with butyllithium activated by N,N,N',N'',N''' -pentamethyldiethylenetriamine**

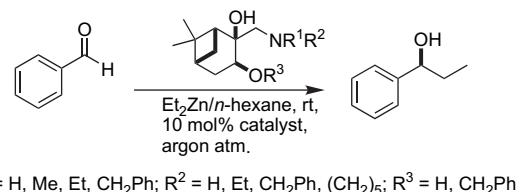
pp 1029–1033

Márta Porcs-Makkay,* Anna Komáromi, Gyula Lukács and Gyula Simig



Synthesis and application of monoterpene-based chiral aminodiols
Zsolt Szakonyi, Anasztázia Hetényi and Ferenc Fülöp*

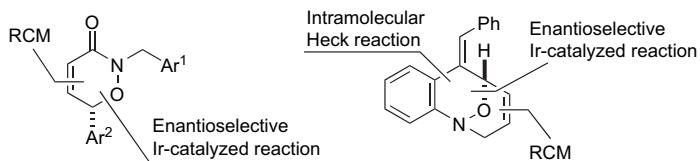
pp 1034–1039



Primary, secondary and tertiary aminodiols derived from (–)- α -pinene were prepared and applied as chiral catalysts in the enantioselective addition of diethylzinc to aromatic aldehydes, resulting in (1S)-1-phenyl-1-propanol with up to ee 84%.

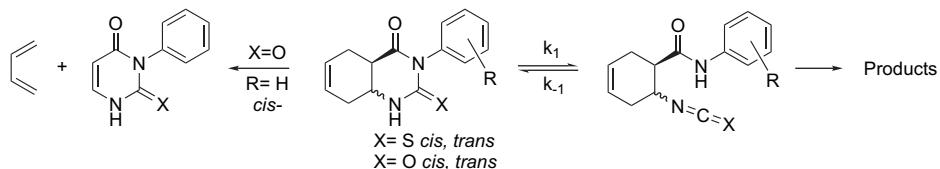
Enantioselective synthesis of [1,2]-oxazinone scaffolds and [1,2]-oxazine core structures of FR900482
Valluru Krishna Reddy, Hideto Miyabe, Masashige Yamauchi and Yoshiji Takemoto*

pp 1040–1048



Flash vacuum pyrolysis (fvp) of some hexahydroquinazolin-4(1*H*)-ones
Walter J. Peláez, Zsolt Szakonyi, Ferenc Fülöp and Gloria I. Yranzo*

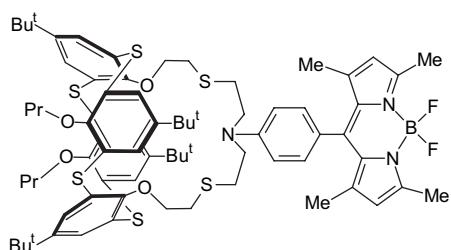
pp 1049–1057



Synthesis, optical and electroanalytical characterizations of a thiocalix[4](N-phenylazacrown-5)ether-BODIPY ionophore

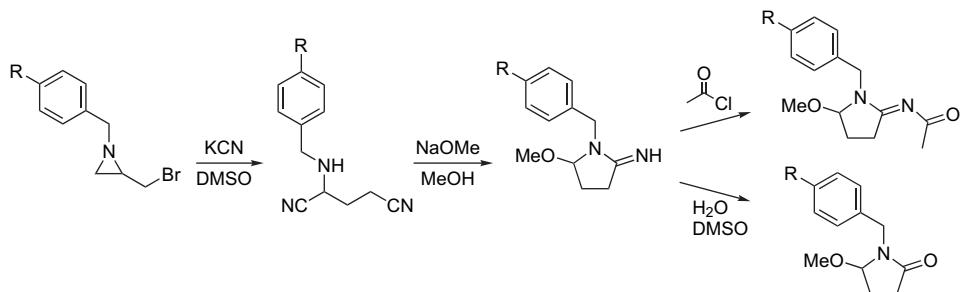
Viktor Csokai, Mihály Kádár, Diem Lan Ha Mai, Olívia Varga, Klára Tóth, Miklós Kubinyi, Alajos Grün and István Bitter*

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Novel synthesis of 2-aminopentanedinitriles from 2-(bromomethyl)aziridines and their transformation into 2-imino-5-methoxypyrrolidines and 5-methoxypyrrrolidin-2-ones

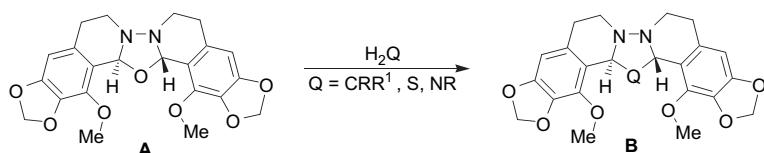
Matthias D'hooghe, Andries Van Nieuwenhove, Willem Van Brabandt, Mario Rottiers and Norbert De Kimpe*



Synthesis of new condensed nitrogen heterocyclic systems

pp 1071–1076

Dezső Korbonits*, Benjamin Podányi, Árpád Illár, Kálmán Simon, Miklós Hanusz, István Hermecz*

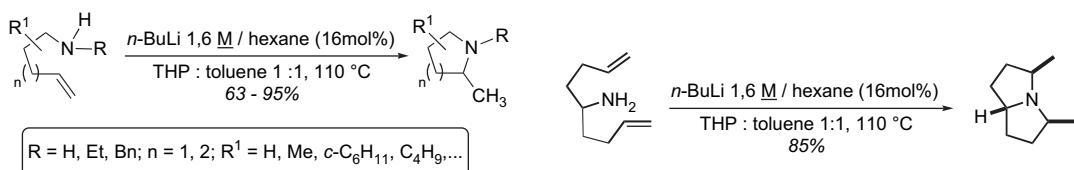


Substitution of the central O atom of the double intramolecular pseudosalt **A** by C, S, or N nucleophiles afforded representatives of three new heterocyclic systems **B**.

Highly efficient, base-catalysed, intramolecular hydroamination of non-activated olefins

pp 1077–1087

Coralie Quinet, Pierre Jourdain, Christophe Hermans, Ali Ates, Isabelle Lucas, István E. Markó*

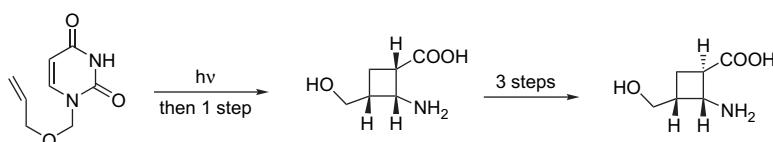


Mono, fused bicyclic and bridged bicyclic amines can be readily prepared by the *n*-BuLi-catalysed intramolecular hydroamination of non-activated olefins.

Efficient synthesis of 3-hydroxymethylated *cis*- and *trans*-cyclobutane β-amino acids using an intramolecular photocycloaddition strategy

pp 1088–1093

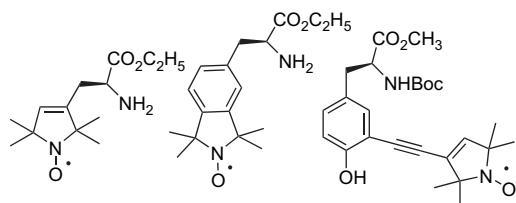
Aurélie Mondière, Runhui Peng, Roland Remuson*, David J. Aitken*



Synthesis and resolution of new paramagnetic α -amino acids

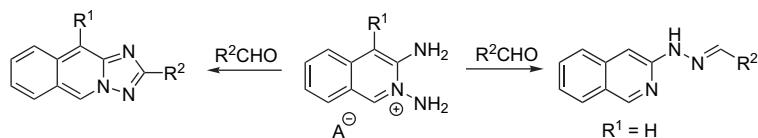
Tamás Kálai, József Schindler, Mária Balog, Elemér Fogassy, Kálmán Hideg*

pp 1094–1100

**A new synthesis of the linearly fused [1,2,4]triazolo[1,5-*b*]isoquinoline ring. Observation of an unexpected Dimroth rearrangement**

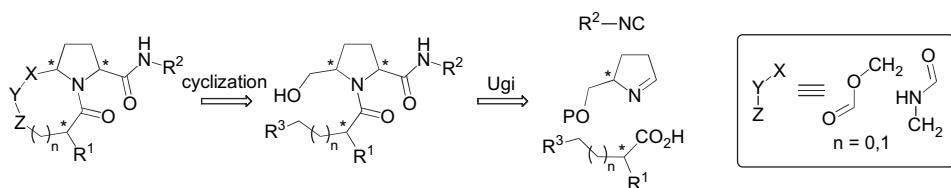
László Filák, Zsuzsanna Riedl, Orsolya Egyed, Mátyás Czugler, Cuong N. Hoang, Joachim G. Schantl, György Hajós*

pp 1101–1113

**A convergent synthesis of enantiopure bicyclic scaffolds through multicomponent Ugi reaction**

Luca Banfi, Andrea Basso, Giuseppe Guanti, Silvia Merlo, Claudio Repetto, Renata Riva*

pp 1114–1134



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