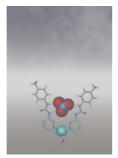
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Cover (far left)

Background: Hemlock (Conium maculatum) (picture William & Wilma Follette @ USDA-NRCS PLANTS Database / USDA NRCS. 1992. Western wetland flora: Field office guide to plant species. West Region, Sacramento, California, USA). Foreground: the structures of conline, one of the hemlock alkaloids, and other compounds that can be obtained from chiral oxime ethers based on (R)/(S)-O-(1-phenylbutyl)hydroxylamine (ROPHy/SOPHy) (pp. 1341–1351).

Inside cover (left)

A silver(I) complex of a simple pyridyl ligand containing a urea derivative templated by nitrate, showing the slight tilt of the nitrate anion (pp. 1352–1353).



Chemical biology articles published in this journal also appear in the Chemical Biology Virtual Journal:

contents

FEATURE ARTICLE

(1341

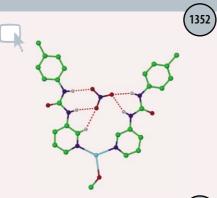
Addition reactions of ROPHy/SOPHy oxime ethers: asymmetric synthesis of nitrogen containing compounds



Christopher J. Moody

Oxime ethers prepared from (R)- or (S)-O-(1-phenylbutyl)-hydroxylamine are versatile intermediates for the asymmetric synthesis of nitrogen containing compounds including amines, amino acids, building blocks of natural products, alkaloids, nitrogen heterocycles, and ferrocene based receptors.

COMMUNICATIONS

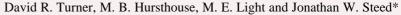


A modular, self-assembled, separated ion pair binding system

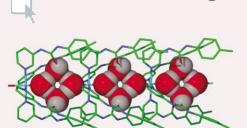
David R. Turner, Elinor C. Spencer, Judith A. K. Howard, Derek A. Tocher and Jonathan W. Steed*

Self-assembly of a discrete ion pair complex in solution and in the solid state with pyridyl–urea ligands requires the presence of both Ag^+ and $\mathrm{NO_3}^-$. In the presence of other anions, dramatically different, infinite solid state systems are observed.

Linear distortion of octahedral metal centres by multiple hydrogen bonds in modular ML_4 systems



A pyridyl urea ligand "L" enclathrates a strongly hydrogen bonded water tetramer in solid state coordination polymers of type ML_4 (M=Co, Ni, Cu, Zn) which display remarkable distortions of the metal coordination environment due to the strength of the hydrogen bonding network.



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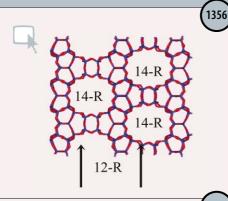
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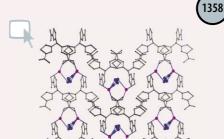
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ITQ-15: The first ultralarge pore zeolite with a bi-directional pore system formed by intersecting 14- and 12-ring channels, and its catalytic implications

Avelino Corma,* Maria Jose Díaz-Cabañas, Fernando Rey, Stavros Nicolopoulus and Khalid Boulahya

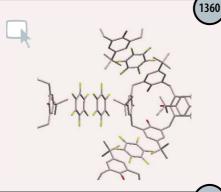
The structure of ITQ-15 is formed by intersecting 14- and 12-ring channels. In its acid form it shows superior activity and selectivity for processing bulky reactants compared with existing 12- or 14-ring zeolites.



Water and tris(5-acetyl-3-thienyl) methane (TATM) assemble into a one-dimensional channel compound

Paul S. Sidhu, Konstantin A. Udachin and John A. Ripmeester

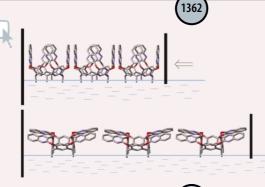
Water induces long-range order into the tris(5-acetyl-3-thienyl)methane host by linking the host acetyl oxygens, giving a one-dimensional channel structure that contains additional disordered water molecules.



Electrostatic and short-range interactions compete in directing the structure of *p-tert*-butylcalix[4]arene inclusion compounds of fluorinated benzenes

Gary D. Enright, Konstantin A. Udachin and John A. Ripmeester*

Some fluorine-substituted benzenes promote p-tert-butylcalix[4]arene guest-host structures where electrostatic interactions dominate the structural motifs by π -stacking with host phenyl rings and sometimes each other.



$\mathbf{Z}\mathbf{n}^{\text{II}}\text{-induced conformational control of amphiphilic cavitands in Langmuir monolayers$

Markus Frei, Federica Marotti and François Diederich*

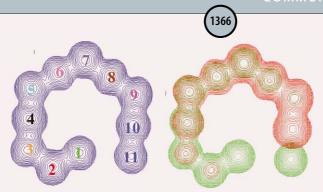
Reversible switching from the $C_{4\nu}$ -symmetric vase to the $C_{2\nu}$ -symmetric kite conformation of resorcin[4]arene cavitands was induced by stoichiometric Zn^{II} ion coordination. Langmuir monolayers were obtained of both conformers with the area per molecule increasing dramatically from 120 Å² for the vase to 270 Å² for the kite form.

hydrogen I $[M] - CI + MeOH, + NH_4PF_6; - NH_4CI$ $[M] = Ru(\eta^5 - C_5H_5)(PPh_3)_2$

A nucleic acid base derivative tethered to a ruthenium carbene complex: hydrogen bonded dimers in both the solid state and solution?

Hayrullo Hamidov, John C. Jeffery and Jason M. Lynam*

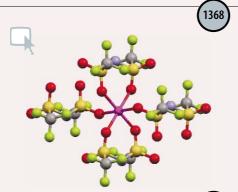
A ruthenium carbene complex containing a tethered uracil group has been prepared and shows a dimeric structure in the solid state: evidence for persistence of this structure in solution has been obtained.



A high-throughput NMR-based ee-assay using chemical shift imaging

Manfred T. Reetz,* Patrick Tielmann, Andreas Eipper, Alfred Ross* and Götz Schlotterbeck

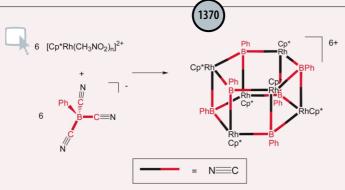
A high-throughput screening system for determining enantiopurity, based on NMR chemical shift imaging, allows up to 5600 ee-determinations per day.



Metal bis{(trifluoromethyl)sulfonyl}amide complexes: highly efficient Friedel-Crafts acylation catalysts

Martyn J. Earle,* Ullastiina Hakala, Barry J. McAuley, Mark Nieuwenhuyzen, Alwar Ramani and Kenneth R. Seddon

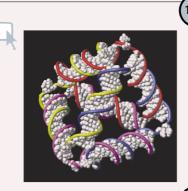
A range of metal bis{(trifluoromethyl)sulfonyl}amide complexes, including many previously unreported ones, have been synthesised, most of which have been found to be excellent Friedel-Crafts acylation catalysts in the absence of solvent; these reactions have also been carried out in ionic liquids, which allow the catalysts to be recycled and reused.



Synthesis and characterization of the hexagonal prismatic cage {THF⊂[PhB(CN)₃]₆[Cp*Rh]₆}⁶⁺

Matthew L. Kuhlman, Haijun Yao and Thomas B. Rauchfuss*

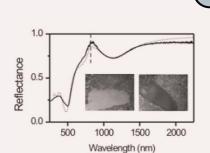
Condensation of $Cp*Rh(CH_3NO_2)_n^{2+}$ and the tricyanoborate $[PhB(CN)_3]^-$ affords the hexagonal bipyramidal cage $\{[PhB(CN)_3]_6[Cp*Rh]_6\}^{6+}$, demonstrating that tetrahedral tricyanide building blocks can lead to cage structures.



The single-step synthesis of a DNA tetrahedron

Russell P. Goodman,* Richard M. Berry and Andrew J. Turberfield

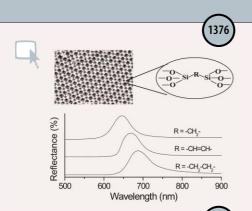
A tetrahedral nanostructure whose edges are DNA double helices self-assembles spontaneously when four oligonucleotides are annealed in solution.



Optical properties of mesoporous II-VI semiconductor compound films

I. S. Nandhakumar,* T. Gabriel, X. Li, G. S. Attard, M. Markham, D. C. Smith and J. J. Baumberg

The optical properties and birefringence of nanostructured mesoporous CdTe films were measured by UV–VIS reflectance spectroscopy and optical microscopy under cross—polarized illumination respectively.



Synthesis, characterization and optical properties of ordered macroporous organosilicas

Zuocheng Zhou, Xiaoying Bao and X. S. Zhao*

Ordered macroporous organosilica materials have been prepared wherein the organic spacer and pore size may be tuned to allow generation of different phases with different refractive indexes.

(1378)

Phase selectively soluble dendrimer-bound osmium complex: a highly effective and easily recyclable catalyst for olefin dihydroxylation

Wei-Jun Tang, Nian-Fa Yang,* Bing Yi, Guo-Jun Deng, Yi-Yong Huang and Qing-Hua Fan^*

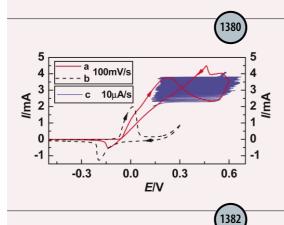
A new switched biphasic catalysis system for highly effective olefin dihydroxylation has been described, in which the dendritic osmium catalyst preferred to dissolve in the non-polar organic layer and could be easily separated from the polar diol products through phase separation induced by addition of water at the end of the reaction.



Transition of oscillatory mechanism for methanol electro-oxidation on nano-structured nickel hydroxide film (NNHF) electrode

Wei Huang, Zelin Li,* Youdi Peng and Zhenjiang Niu

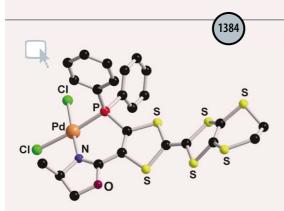
Instead of CO_{ad} formation and removal as on Pt electrodes, coupling of charge transfer with diffusion and convection mass transfer accounts for the oscillation found in the methanol electro-oxidation on the NNHF electrode.



Allylation and highly diastereoselective syn or anti crotylation of N-toluenesulfonylimines using potassium allyl- and crotyltrifluoroborates

Sze-Wan Li and Robert A. Batey*

Air and moisture stable potassium allyl- and crotyltrifluoroborates undergo addition to *N*-sulfonyl and *N*-sulfinyl aldimines in the presence of Lewis acids, to provide the corresponding homoallylic amines in high yields and excellent diastereoselectivity.



Tetrathiafulvalene based phosphino-oxazolines: a new family of redox active chiral ligands

Céline Réthoré, Marc Fourmigué* and Narcis Avarvari*

Electroactive chiral ethylenedithio-tetrathiafulvalene-oxazolines (EDT-TTF-OX) and -phosphinooxazolines (EDT-TTF-PHOX) have been synthesized and characterized, together with the racemic palladium complex (EDT-TTF-PHOX)PdCl₂.

Bis(phenanthroline)-ethylenediamine conjugate displays excimer fluorescence upon binding with DNA

Keigo Hayashi, Hitoshi Akutsu, Hiroaki Ozaki and Hiroaki Sawai*

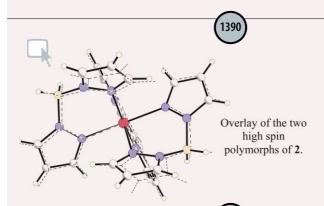
1,2-Bis(1,10-phenanthrolin-2-yl)ethylenediamine **1** displays a unique excimer fluorescence at 528 nm by binding with DNA, while its monomer fluorescence at 400 nm shows a decrease upon binding with DNA. **1** works as a groove-binder to DNA.

X N Pd/dppf Na₂CO₃ R CN

Potassium hexacyanoferrate(II)—a new cyanating agent for the palladium-catalyzed cyanation of aryl halides

Thomas Schareina, Alexander Zapf and Matthias Beller*

The use of potassium hexacyanoferrate(II) as a cyanating agent for the general synthesis of benzonitriles is described. Under optimized reaction conditions good yields of the desired products and unprecedented catalyst productivities have been achieved.



Thermal and light induced polymorphism in $\text{iron}(\Pi)$ spin crossover compounds

Amber L. Thompson, Andrés E. Goeta,* José A. Real,* Ana Galet and M. Carmen Muñoz

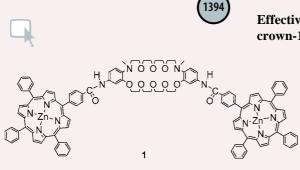
The crystal structures of the thermal and light induced spin states of $\{Fe[H_2B(pz)_2]_2L\}$ $\{[H_2B(pz)_2]^-=$ dihydrobis(pyrazolyl)borate, L=2,2'-bipyridine (1), bipy and 1,10-phenanthroline, phen (2)), demonstrate that 2 exhibits "thermal and light induced polymorphism," while 1 does not.

85-95% ee

A novel asymmetric route to succinimides and derived compounds: synthesis of the lignan lactone (+)-hinokinin

D. Jonathan Bennett, Paula L. Pickering and Nigel S. Simpkins*

A novel approach to chiral succinimides and derived compounds has been developed that involves chiral lithium amide desymmetrisation of an *N-ortho-tert*-butylphenyl succinimide to generate a putative atropisomeric intermediate enolate, alkylation of which enables access to the lignan lactone (+)-hinokinin.



electrophile

Effective cation-assisted chirality induction using a dibenzo-diaza-30-crown-10 with bis(zinc(Π) porphyrin) units

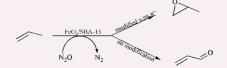
Yuji Kubo,* Yusuke Ishii, Toshiharu Yoshizawa and Sumio Tokita

The first rationalized diaza crown ether-bis(porphyrin) conjugate with a large ring-based conformation flexibility has been synthesized, showing an effective K^+ -assisted chirality induction.

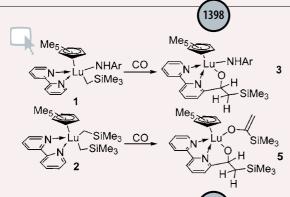


Iron-catalysed propylene epoxidation by nitrous oxide: dramatic shift of allylic oxidation to epoxidation by the modification with alkali metal salts

Xiaoxing Wang, Qinghong Zhang, Qian Guo, Yinchuan Lou, Lüjuan Yang and Ye Wang^*



The modification of FeO $_x$ /SBA-15 with K $^+$ dramatically shifted the allylic oxidation of C $_3$ H $_6$ to epoxidation by N $_2$ O



C-H activation of a 2,2'-bipyridine ligand within (mono)pentamethylcyclopentadienyl lutetium complexes

Thomas M. Cameron,* John C. Gordon, Brian L. Scott and William Tumas

We report the activation of a 2,2'-bipyridine ligand within a class of (mono)cyclopentadienyl lanthanide complexes when reacted with carbon monoxide.

1400

Palladium catalyzed amination of vinyl chlorides: a new entry to imines, enamines and 2-amino-1,3-butadienes

José Barluenga,* M. Alejandro Fernández, Fernando Aznar and Carlos Valdés

$$R^{1} = Ar, -CH = CH - R$$

Vinyl chlorides are used in palladium catalyzed cross-coupling reactions with amines to give rise to enamines, imines and 2-amino-1,3-butadienes with very high yields and chemoselectivity.





Plasmachemical surface functionalised beads: versatile tailored supports for polymer assisted organic synthesis

Jas Pal Badyal, Audrey M. Cameron, Neil R. Cameron, Diane M. Coe, Richard Cox, Benjamin G. Davis,* Leslie J. Oates, Gisle Oye, Christos Spanos and Patrick G. Steel*



Plasmachemical surface modification of porous polystyrene beads with allylamine or diaminopropane provides reactive amine functionality exclusively at accessible surface sites, allowing faster reactions than classically prepared materials.



Fe–Ce–ZSM-5 a new catalyst of outstanding properties in the selective catalytic reduction of NO with NH₃

G. Carja, G. Delahay,* C. Signorile and B. Coq

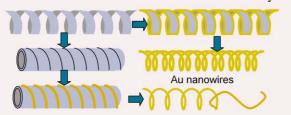
$$2 \text{ NO} + 2 \text{ NH}_3 + 1/2 \text{ O}_2 \xrightarrow{\text{Fe-Ce-ZSM-5}} 2 \text{ N}_2 + 3 \text{ H}_2 \text{O}$$

A Fe–Ce–ZSM-5 catalyst elaborated from a new synthesis route exhibits very high NO conversion (75–100%) in the selective catalytic reduction of NO by NH_3 in a wide temperature window (523–823 K), even in the presence of H_2O and SO_2 .

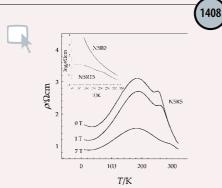


Preparation of helical gold nanowires on surfactant tubules

Ryo Takahashi and Tsutomu Ishiwatari*



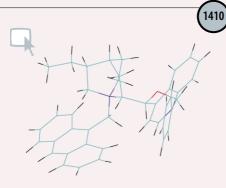
A novel approach has been found for the synthesis of gold nanowires shaped as single, double, and multiple helical structures, using surfactant tubules as template.



Unexpected effect of Ru-substitution in lightly doped manganites

Lorenzo Malavasi,* M. Cristina Mozzati, Cristina Tealdi, M. Rosa Pascarelli, Carlo B. Azzoni and Giorgio Flor

In this communication we report about an unexpected effect of Ru-doping on La_{1-x}Na_xMnO_{3+\delta} manganites. These results are explained by considering a compensation mechanism based on oxygen content variation.



Computational screening of combinatorial catalyst libraries

James L. Melville, Benjamin I. Andrews, Barry Lygo and Jonathan D. Hirst*

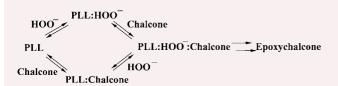
A computational catalyst design methodology is presented, utilizing combinatorial synthesis in parallel with chemometric analysis. The technique has been successfully applied to a library of cinchonidine-based phase transfer catalysts.



Kinetics of chalcone oxidation by peroxide anion catalysed by poly-L-leucine

Giacomo Carrea, Stefano Colonna, Alastair D. Meek, Gianluca Ottolina* and Stanley M. Roberts

The kinetics of the enantioselective chalcone oxidation by H₂O₂, catalysed by poly-L-leucine, follows a steady state random bireactant mechanism with one of the pathways - HOO- binding first - kinetically preferred.



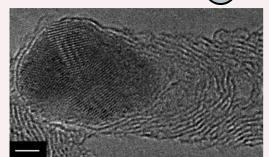
1412

Asymmetric Michael addition reactions of 2-silyloxyfurans catalyzed by binaphthyldiimine-Ni(II) complexes $= Me. R^2 = H$

Hiroyuki Suga,* Takeo Kitamura, Akikazu Kakehi and Toshihide Baba

N,N'-Bis(2-quinolylmethylene)-1,1'-binaphthyl-2,2'-diamine–Ni(II) complex and analogous complexes were found to be efficient chiral Lewis acid catalysts for the asymmetric Michael addition reactions between 2-silyloxyfurans and 3-alkenoyl-2-oxazolidinones.





Growth of aligned carbon nanofibres over large areas using colloidal catalysts at low temperatures

B. Kleinsorge, V. B. Golovko, S. Hofmann, J. Geng, D. Jefferson, J. Robertson and B. F. G. Johnson*

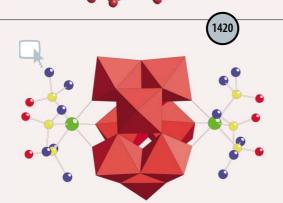
Highly purified cobalt colloids have been employed as a catalyst to grow aligned carbon nanofibres at temperatures as low as 300 °C by dc plasma enhanced chemical vapour deposition over large areas.

(1418

Supertetrahedral decametallic Ni(II) clusters directed by $\mu_6\text{-tris}\text{-alkoxides}$

Rachel Shaw, Ian S. Tidmarsh, Rebecca H. Laye, Barbara Breeze, Madeleine Helliwell, Euan K. Brechin,* Sarah L. Heath, Mark Murrie, Stefan Ochsenbein, Hans-Ulrich Güdel* and Eric J. L. McInnes*

Reactions of triol ligands with $Ni(\Pi)$ diketonates in superheated alcohol or acetonitrile solutions give decametallic $Ni(\Pi)$ clusters with highly regular supertetrahedral cores.



A novel isopolytung state functionalized by ruthenium: $[HW_9O_{33}Ru^{II}_{\ 2}(dmso)_6]^{7-}$

Lihua Bi, Firasat Hussain, Ulrich Kortz,* Masahiro Sadakane and Michael H. Dickman

The ruthenium-supported isopolyanion $[HW_9O_{33}Ru^{II}_2(dmso)_6]^{7-}$ (1) is composed of a nonatungstate wheel stabilized by two Ru(dmso)₃ groups, representing the first structurally characterized Ru-coordinated polyoxotungstate and a novel class of isopolyanions supporting photochromic moieties.

1422

Rearrangement of pyrrolines derived from the Birch reduction of electron-deficient pyrroles: radical ring-expansion to substituted tetrahydropyridines

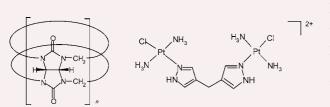
Peter G. Turner, Timothy J. Donohoe* and Rick P. C. Cousins

The Birch reduction of pyrroles can be quenched with dihalomethanes, leading to compounds which are excellent precursors for radical ring expansion reactions.



1424

Multi-nuclear platinum complexes encapsulated in cucurbit[n]uril as an approach to reduce toxicity in cancer treatment



Nial J. Wheate, Anthony I. Day, Rodney J. Blanch, Alan P. Arnold, Carleen Cullinane and J. Grant Collins*

Encapsulation of *trans*-[{PtCl(NH₃)₂}₂ μ -dpzm]²⁺ in cucurbituril slows the rate of reaction of the platinum complex with guanosine without significantly affecting its cytotoxicity.



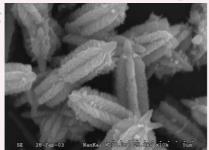
Selective production of hydrogen from partial oxidation of methanol over silver catalysts at low temperatures

Liuye Mo, Xiaoming Zheng and Chuin-Tih Yeh*

$$CH_3OH + 1/2O_2 \xrightarrow{\text{Ag/CeO}_2 - ZnO} CO_2 + 2H_2$$

Hydrogen can be selectively produced from the partial oxidation of methanol over Ag/CeO₂–ZnO catalyst at a low temperatures <200 °C.





Rotor-like ZnO by epitaxial growth under hydrothermal conditions

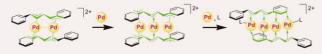
X. P. Gao,* Z. F. Zheng, H. Y. Zhu, G. L. Pan, J. L. Bao, F. Wu and D. Y. Song

Rotor-like ZnO was grown from a mixture of rod-like ZnO and a saturated $\text{Zn}(\text{OH})_4^{2-}$ solution under moderate hydrothermal conditions at 100 °C, in which the rod-like ZnO crystal plane acts as a matrix core, and the branched nanorods showed fast epitaxial growth.



Stepwise growth of polypalladium chains in 1,4-diphenyl-1,3-butadiene sandwich complexes

Yasuki Tatsumi, Tomoki Naga, Hiromitsu Nakashima, Tetsuro Murahashi* and Hideo Kurosawa*



stepwise growing

A series of 1,4-diphenyl-1,3-butadiene multinuclear palladium sandwich complexes are synthesized by stepwise growth of palladium chains within a sandwich framework.



p-Type macroporous silicon having three-dimensional structure

Sung-Ho Lee* and Ryutaro Maeda

zeliti i a do 5500

Novel macroporous silicon with three-dimensional structure was fabricated using an organic-based electrolyte, dimethylformamide (DMF), in the p-type silicon. The obtained three-dimensional macroporous structure grew wholly along the <100> orientation of the p-type silicon wafer.





Cross-metathesis between ethane and toluene catalyzed by [(≡SiO)₂TaH]: the first example of a cross-metathesis reaction between an alkane and an aromatic

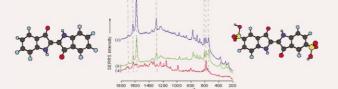
Mostafa Taoufik, Ekkehard Schwab, Michael Schultz, Dominic Vanoppen, Marc Walter, Jean Thivolle-Cazat* and Jean-Marie Basset*

The silica-supported tantalum hydride [(\equiv SiO)₂TaH] **1** catalyzes at moderate temperature (150–250 °C) the cross-metathesis reaction between toluene and ethane, to form mainly ethylbenzene and xylenes.

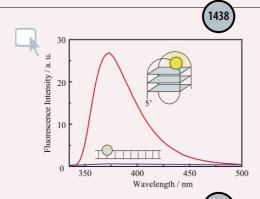
(1436

Analysis of the conversion of indigo into indigo carmine dye using SERRS

Iqbal T. Shadi, Babur Z. Chowdhry,* Martin J. Snowden and Robert Withnall*



In a novel application SERRS has been used, by employing a silver sol, to monitor and analyse the conversion of indigo into the indigo carmine dye.



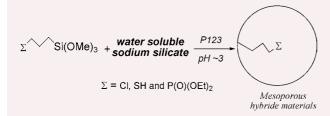
Fluorescence properties of 2-aminopurine in human telomeric DNA

Takumi Kimura, Kiyohiko Kawai, Mamoru Fujitsuka and Tetsuro Majima*

Ap was used to monitor the duplex to quadruplex conformational change in human telomeric DNA based on the fluorescence intensities and lifetimes of Ap in the duplexes and quadruplexes.



Direct syntheses of functionalized mesostructured silica by using an inexpensive silica source



Robert J. P. Corriu,* Ahmad Mehdi, Catherine Reyé and Chloé Thieuleux

A general and economic one step synthesis methodology for the preparation of organically functionalised mesostructured silica was obtained by the co-condensation of water soluble sodium silicate and different function organotrialkoxysilanes.

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