

A union formed by chemi-



BELGIUM

SVENSA

TE NEW TOE!

Austria



Chem PubSoc

Europe

SPAIN













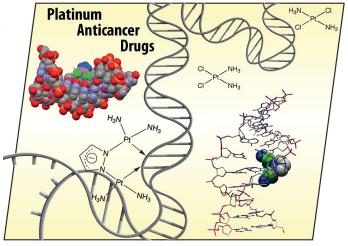








The cover picture shows an impression of the binding of dinuclear platinum compounds to double-stranded DNA. Details are presented in the Microreview by J. Reedijk on p. 1303ff.



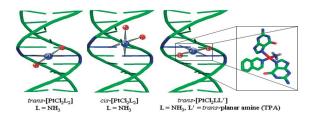
MICROREVIEWS

trans-Platinum Antitumor Agents

S. M. Aris, N. P. Farrell*...... 1293-1302

Towards Antitumor Active *trans*-Platinum Compounds

Keywords: Platinum / Structure—activity relationships / DNA damage / Antitumor agents / Drug design



This microreview summarizes the chemistry and biology of *trans*-platinum compounds containing principally planar amines and succinctly reviews the current status of anticancer relevance of the *trans*-

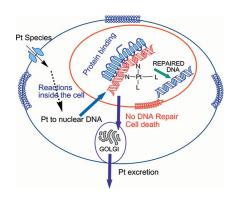
platinum geometry. The potential for novel bifunctional DNA-DNA and DNA-protein adduct formation and their relevance to antitumor activity is succinctly discussed.

Platinum Anticancer Chemistry

J. Reedijk* 1303-1312

Platinum Anticancer Coordination Compounds: Study of DNA Binding Inspires New Drug Design

Keywords: Antitumor agents / Cisplatin / Medicinal chemistry / Platinum / Bioinorganic chemistry / Nucleic acids



The chemistry of platinum anticancer drugs and their binding with nucleic acids is discussed in detail. From mechanistic knowledge at the molecular level, like DNA binding, new drug candidates are discussed. Design and synthesis of new compounds is focused on their bifunctionality. Several biochemical and biophysical studies are reviewed as well.

SHORT COMMUNICATIONS

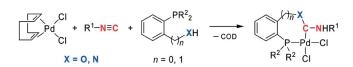
(Phosphanyl-carbene) Pd Complexes

M. R. Eberhard, B. van Vliet,

L. Durán Páchon, G. Rothenberg,

G. Eastham, H. Kooijman, A. L. Spek,

C. J. Elsevier* 1313–1316





A Simple Building-Block Route to (Phosphanyl-carbene)palladium Complexes via Intermolecular Addition of Functionalised Phosphanes to Isocyanides

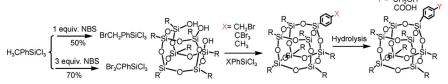
Keywords: Carbenes / Carbene complexes / Palladium / Intermolecular addition

A simple modular synthesis of (phosphanyl-carbene)Pd complexes is presented. This building-block approach to a range of Pd pre-catalysts containing a 'sticky' carbene moiety and a hemi-labile phosphane ligand avoids the synthesis of imidazolium

salts or free carbenes. The design involves a new intermolecular addition of the polar head group of a readily available phosphane-alcohol or -amine to a Pd-coordinated isocyanide.



Octasilsesquioxanes



An effective and milder route to organofunctional trichlorosilanes is reported: mono- and tribrominated trichloro(*p*-tolyl)silane can be selectively prepared by controlling the ratio of *N*-bromosuccinimide (NBS) to trichloro(p-tolyl)silane. From these compounds, several monofunctionalized octasilsesquioxanes (T_8) are obtained, which are very useful precursors for the preparation of new materials.

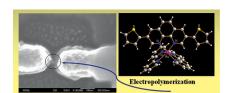
H. Liu,* S. Kondo, N. Takeda, M. Unno* 1317–1319

An Efficient Approach to Monophenyl-Functionalized Octasilsesquioxanes

Keywords: Functionalization / Bromination / Cage silsesquioxanes / Synthesis design

FULL PAPERS

The formation of an oligothiophene polymer bridging gold electrodes with a gap of around 25 nm by in situ electropolymerization, and its removal, is studied by SEM and temperature-dependent electrochemical techniques (I-V curves).



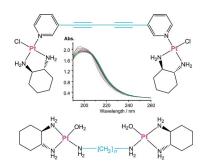
Electropolymerization

W. Huang,* L. Wang, H. Tanaka, T. Ogawa* 1321–1330

Spectral, Structural, and Computational Studies of a New Family of Ruthenium(II) Complexes Containing Substituted 1,10-Phenanthroline Ligands and in situ Electropolymerization of a Phenanthrolineruthenium(II) Complex Bridging Nanogap Gold Electrodes

Keywords: Ruthenium / N ligands / Electrochemistry / Electropolymerization / Thin films / Density functional calculations

The influence of different types of bridging ligands on the thermodynamic and kinetic properties of dinuclear Pt^{II} complexes was investigated for this study. The distance between the Pt^{II} centres and the electron acceptor/donator ability of the bridging ligand have proven to be the crucial factors.



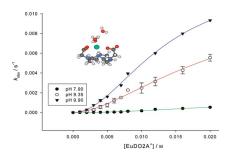
Dinuclear Platinum Complexes

H. Ertürk, R. Puchta, R. van Eldik* 1331–1338

Synthesis, Characterization, Thermodynamic and Kinetic Properties of a New Series of Dinuclear Pt^{II} Complexes

Keywords: Platinum / Antitumor agents / Kinetics

A detailed analysis of the BNPP phosphodiester bond hydrolysis rate constants data vs. lanthanide complex concentration at pH 7.90, 9.35, and 9.90 indicates that dimeric species are more reactive than their monomeric analogues. The overall formation constants for the (Ln-dimer)-BNPP species are about $10^4 \, \mathrm{m}^{-1}$ and the hydrolysis reactivities are in the order Eu > Yb > Er.



Kinetics of Macrocyclic Ln Complexes

C. A. Chang,* B. H. Wu, C.-H. Hsiao 1339–1346

Effects of Concentration of Some Lanthanide(III) Complexes of 1,7-Bis(carboxymethyl)-1,4,7,10-tetraazacyclododecane on Bis(*p*-nitrophenyl)phosphate Hydrolysis

Keywords: Lanthanides / Macrocyclic ligands / Phosphodiesters / Hydrolysis / Kinetics / Reaction mechanisms

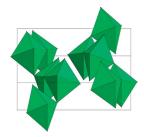
CONTENTS

Cu/ZnO Catalyst Precursors

M. Behrens,* F. Girgsdies, A. Trunschke, R. Schlögl 1347–1357

Minerals as Model Compounds for Cu/ZnO Catalyst Precursors: Structural and Thermal Properties and IR Spectra of Mineral and Synthetic (Zincian) Malachite, Rosasite and Aurichalcite and a Catalyst Precursor Mixture

Keywords: Copper / Zinc / Heterogeneous catalysis / X-ray diffraction / Thermochemistry



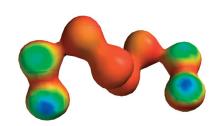
Minerals, single-phase synthetic samples and a typical phase mixture employed as a precursor for Cu/ZnO catalysts are compared and their structural, thermal and IR spectroscopic properties studied. A comprehensive characterisation of this phase mixture is attempted, and the implications of the precursor chemistry (phase composition, Cu/Zn ratio) on the final catalyst are discussed

Sulfur-Sulfur Bond Energies

M. K. Denk* 1358-1368

The Variable Strength of the Sulfur—Sulfur Bond: 78 to 41 kcal — G3, CBS-Q, and DFT Bond Energies of Sulfur (S₈) and Diκlfanes XSSX (X = H, F, Cl, CH₃, CN, NH₂, OH, SH)

Keywords: Sulfur / Radicals / Density functional calculations / Thermochemistry / Bond order / Spin density

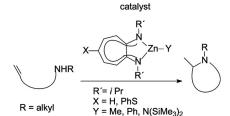


High-precision thermochemical calculations (G3, CBS-Q) and DFT methods were employed to establish the strength of the sulfur–sulfur bond. The delocalization of the spin density in the ${}^{\circ}S-S_6-S^{\circ}$ diradical at the B3LYP/6-311G(2d,p) level is shown in the graphic.

Hydroamination

Aminotroponiminate Zinc Complexes with Different Leaving Groups as Catalysts for the Intramolecular Hydroamination of Alkenes

Keywords: Homogeneous catalysis / Heterocycles / Hydroamination / N ligands / Zinc / Organozinc compounds

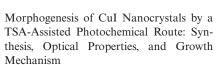


Six different (aminotroponiminato)phenylzinc and bis(trimethylsilyl)amido complexes were prepared and fully characterized. The new complexes were compared with the corresponding zinc methyl complexes to study the influence of the leaving group of the zinc atom in the catalytic hydroamination / cyclization reaction.

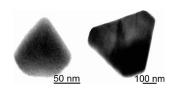
Crystal Engineering

B. C. Zhang, A. J. Xie, Y. H. Shen,* L. B. Yang, Y. P. Huang,

J. J. Lu 1376-1384



Keywords: Copper / CuI / Nanocrystals / Crystal growth / Crystal engineering / Photochemistry / Optical properties

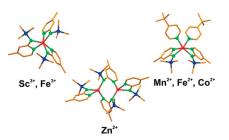


Optical materials consisting of γ -CuI nanocrystals with diamond and platelet morphologies were synthesised by utilising UV-irradiated 12-tungstosilicate as a reducing agent for the reduction of Cu^{2+} and I_2 under ambient conditions. The synthetic processes were carefully followed by UV/Vis absorption and photoluminescence (PL) spectroscopy. A possible mechanism for the formation of diamond and platelet morphologies has been proposed.



Aminopyridinate Complexes

Aminopyridinate complexes of Sc³⁺, Mn²⁺, Fe³⁺, Fe²⁺, Co²⁺ and Zn²⁺ are synthesised and characterised and their magnetic behaviour investigated. One specific aminopyridinato ligand is able to stabilise all first row transition metal complexes, which indicates the high versatility of this ligand class.



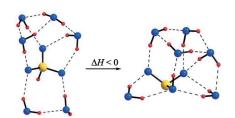
G. Glatz, S. Demeshko, G. Motz, R. Kempe* 1385–1392

First Row Transition Metal Aminopyridinates – the Missing Complexes

Keywords: N ligands / Cobalt / Iron / Manganese / Scandium / Zinc

Microsolvation

A large number of hydrates of oxo S^{IV} species have been studied theoretically. The conversion of SO₂·6H₂O into its isomer H₂SO₃·5H₂O is predicted to be exothermic. The relative stability of sulfonate and hydrogensulfite anions [SHO₃]⁻ is reversed by solvation with more than four water molecules. In a polar continuum less water molecules have the same effect.



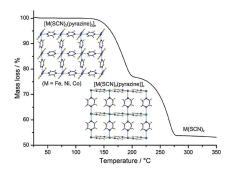
R. Steudel,* Y. Steudel 1393-1405

Sulfur Dioxide and Water: Structures and Energies of the Hydrated Species SO_2 · nH_2O , $[HSO_3]$ -· nH_2O , $[SO_3H]$ -· nH_2O , and H_2SO_3 · nH_2O (n = 0 - 8)

Keywords: Sulfur / Hydrates / Ab initio calculations / Hydrogen bonds / Thermodynamics

Coordination Polymers

Four new (thiocyanato)metal coordination compounds with pyrazine as N-donor ligand are prepared and their structures and thermal and magnetic properties determined. The ligand-rich 1:2 compounds transform quantitatively into new ligand-deficient 1:1 compounds, some of which show a different magnetic behavior, depending on the connection mode of the thiocyanate anions, on heating.



M. Wriedt, I. Jeß, C. Näther* 1406–1413

Synthesis, Crystal Structure, and Thermal and Magnetic Properties of New Transition Metal—Pyrazine Coordination Polymers

Keywords: N ligands / Coordination polymers / Crystal structures / Thermal properties / Magnetic properties

Supporting information on the WWW (see article for access details).

If not otherwise indicated in the article, papers in issue 9 were published online on March 9, 2009

^{*} Author to whom correspondence should be addressed.