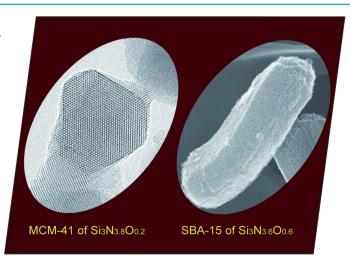


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Other ChemPubSoc Europe journals are Chemistry – A European Journal, ChemBioChem, ChemPhysChem, ChemMedChem, ChemSusChem and ChemCatChem.

COVER PICTURE

The cover picture shows a new family of mesoporous silicon (oxy)nitrides prepared by the nitridation of mesoporous silicas with ammonia. The (oxy)nitrides obtained maintained their respective regular pore structures. The nitrogen contents could be controlled by the amounts of ammonia supplied. The use of a plug-flow reactor instead of a usual boat-type reactor is a key to the successful synthesis of highly nitrided samples. Details are discussed in the article by M. Iwamoto et al. on p. 2235 ff.



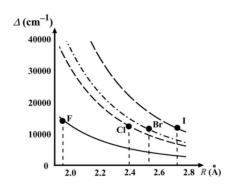
SHORT COMMUNICATIONS

Ligand Field Theory

H. Bolvin*..... 2221-2223

Ab Initio Investigation of Spectroscopic Parameters for ${\rm CrX_6}^{3-}$: The Unexpected Role of Bond Length Variation – Size Really Matters

Keywords: Ab initio calculations / Ligand field theory / Spectrochemical series / Excited states / Transition metals / Halides



Within the CrX_6^{3-} series, it is shown that π effects decrease down the series and that the decrease in the ligand field parameter, Δ , is due to the lengthening of the bond.

Cancer Therapy

K. L. Ciesienski, L. M. Hyman,

D. T. Yang, K. L. Haas, M. G. Dickens,

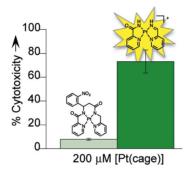
R. J. Holbrook,

K. J. Franz* 2224-2228



A Photo-Caged Platinum(II) Complex That Increases Cytotoxicity upon Light Activation

Keywords: Platinum / Cisplatin / Caged compounds / Phototherapy / Photoactivation / Prodrugs / Cytotoxicity



Photoinitiated drug release is an attractive strategy for increasing drug efficacy only at the irradiated site while minimizing toxicity to surrounding healthy tissue. Here we show that a neutral, nontoxic Pt^{II} complex that is inert to ligand exchange reactions can be converted by UV light to a charged complex with light-dependent cytotoxicity.

Mixed-Valence Cobalt Clusters

L.-L. Zheng, J.-D. Leng, R. Herchel,

Y.-H. Lan, A. K. Powell,

M.-L. Tong* 2229-2234



Anion-Dependent Facile Route to Magnetic Dinuclear and Dodecanuclear Cobalt Clusters

Keywords: Azides / Cobalt / Cluster compounds / Magnetic properties



A dodecanuclear $\text{Co}^{\text{III}}_{3}\text{Co}^{\text{II}}_{9}$ and a dinuclear $\text{Co}^{\text{II}}_{2}$ cluster were synthesized facilely from reactions of different cobalt(II) salts with the ligand (1*H*-benzimidazol-2-yl)-

methanol. The cobalt ions in the unprecedented Co_{12} supercluster are linked into a disclike structure through $\mu_3\text{-}O_L$, $\mu\text{-}O_L$, $\mu_3\text{-}O^2\text{-}$, $\mu_{1.1}\text{-}N_3^-$ and $\mu_{1,1,1}\text{-}N_3^-$ bridges.

FULL PAPERS

Mesoporous Silicon (Oxy)Nitride

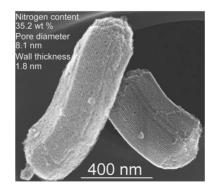
F. Hayashi, K.-i. Ishizu,

M. Iwamoto*...... 2235-2243



Effect of Pore Structure on the Nitridation of Mesoporous Silica with Ammonia

Keywords: Mesoporous materials / Nitrides / Nanostructures / Heterogeneous catalysis / Ceramics

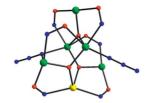


Mesoporous silicon (oxy)nitrides with regular pore structures were prepared by nitridation of mesoporous silica MCM-41, SBA-15, and MCM-48 with ammonia. The nitrogen contents were 35–39 wt.-%. The reaction rates were dependent on the surface areas. Characterization revealed no collapse of the regular pore structure through the nitridation.



Polynuclear Manganese Complexes

Two new hexanuclear, mixed-valence, Mn^{II/III} cage-like clusters have been obtained from the amalgamation of azide and methyl 2-pyridyl ketone oxime ligands in Mn non-carboxylate chemistry. The complexes both possess small (S = 5/2) spin ground states that suggest the presence of spin frustration effects.



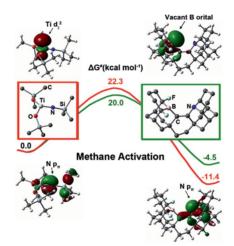
C. Lampropoulos, T. C. Stamatatos, M. J. Manos, A. J. Tasiopoulos, K. A. Abboud, G. Christou* ... 2244-2253

New Mixed-Valence Mn^{II/III}₆ Complexes Bearing Oximato and Azido Ligands: Synthesis, and Structural and Magnetic Characterization

Keywords: Mixed-valent compounds / Polynuclear complexes / Manganese / Azides / Magnetic properties

Methane Activation

Whereas the transition-metal (TM) complex maintains an active site through symmetry-unmatched orbitals, the designed molecule uses linkages to prohibit electron donation from the N to the B center, thus preserving an effective active site for Lewis acid/base effects. Some of these molecules have kinetics and thermodynamics in activating methane comparable with or better than those of the TM complex.



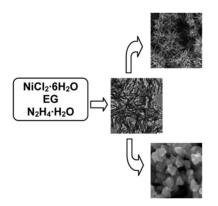
G. Lu, L. Zhao, H. Li, F. Huang, Z.-X. Wang* 2254-2260

Reversible Heterolytic Methane Activation of Metal-Free Closed-Shell Molecules: A Computational Proof-of-Principle Study

Keywords: Ab initio calculations / C-H activation / Metal-free methane activation / Frustrated Lewis pair / Lewis acid/base ef-

Ni Nanoflowers

3D hierarchical flower/star-like Ni nanostructures were formed. Once the reagents are mixed, the rod-like morphology of the intermediate products [Ni(N₂H₄)₂]Cl₂ are first synthesized. Depending on the experimental temperature, nickel nanostructures varying from a flower-like superstructure to nanostars can be achieved by the oriented growth of preferred nickel crystal planes.

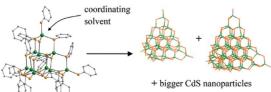


P. Li, N. Wang,* R. Wang* 2261-2265

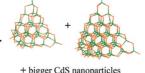
Flower-Like Nickel Nanocrystals: Facile Synthesis, Shape Evolution, and Their Magnetic Properties

Keywords: Nickel / Nanostructures / Solution synthesis / Magnetic properties

Cadmium Sulfide Nanoparticles



The molecular cluster [Cd₁₀S₄(SPh)₁₆]⁴⁻ undergoes rearrangement reactions in coordinating solvents resulting in larger clus-



ters and eventually formation of CdS nanoparticles. The mechanism of this transformation was investigated.

H. Peterlik, U. Schubert* 2266-2275

M. Bendova, M. Puchberger, S. Pabisch,

Studies on the Formation of CdS Nanoparticles from Solutions of (NMe₄)₄- $[Cd_{10}S_4(SPh)_{16}]$

Keywords: Cadmium / Cadmium sulfide / Nanoparticles / Cluster compounds / S ligands / Rearrangement



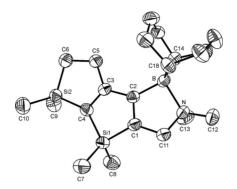
CONTENTS

Fused Silacarbacycles

B. Wrackmeyer,* O. L. Tok, E. V. Klimkina, W. Milius...... 2276-2282

Fused Silacarbacycles Containing a Silole Unit: 1,2-Hydroboration and 1,1-Organoboration of Alkynyl(vinyl)silanes

Keywords: Alkynylsilanes / Hydroboration / Organoboration / Siloles / NMR spectroscopy



Alkynyl(vinyl)silanes containing two alkynyl groups react with 9-borabicyclo[3.3.1]-nonane (9-BBN) by consecutive 1,2-hydroboration of the vinyl group and twofold intramolecular 1,1-organoboration to afford 1,6-disilapentalene derivatives, fused silacarbacycles with a silole unit.

Task-Specific Ionic Liquids

Synthesis and Characterization of Task-Specific Ionic Liquids Based on Peroxydisulfate and Their Application in Oxidation Reactions

Keywords: Ionic liquids / Peroxydisulfate salts / Imidazolium salts / Oxidation / Alcohols

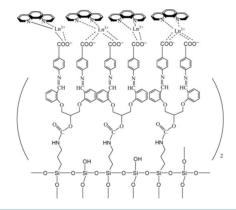
Four task-specific ionic liquids based on peroxydisulfate anions were synthesized and characterized. These ionic liquids have the desired properties for given oxidation reactions of alcohols and thiols, and they show excellent performance as both oxidant and medium in solvent- and catalyst-free reaction systems.

Luminescent Lanthanide Hybrids

J.-L. Liu, B. Yan,* L. Guo 2290-2296

Photoactive Ternary Lanthanide-Centered Hybrids with Schiff-Base Functionalized Polysilsesquioxane Bridges and N-Heterocyclic Ligands

Keywords: Organic—inorganic hybrid composites / Lanthanides / Schiff bases / Luminescence / Polysilsesquioxane bridges



This work focuses on the synthesis of a series of silica-based organic—inorganic hybrid materials, containing different Schiffbase organic compounds, through a covalent self-assembly process. The resulting materials exhibit regular, uniform microstructures and the organic and inorganic compounds are covalently linked through Si–O bonds.

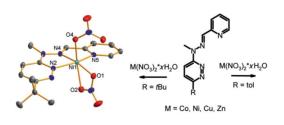
Schiff Base Pyridazine Complexes

K. R. Grünwald, G. Saischek, M. Volpe, F. Belaj, N. C. Mösch-Zanetti* 2297–2305

Pyridazine-Based Ligands and Their Coor-

Pyridazine-Based Ligands and Their Coordinating Ability towards First-Row Transition Metals

Keywords: Nitrogen heterocycles / Schiff bases / Tridentate ligands / Chelates



A new class of pyridazine-based ligands was prepared and their coordination of first-row transition metals was tested. Evidence from X-ray diffraction and electro-

chemical analysis shows the formation of 1:1 and 1:2 complexes depending on the electronic influence exerted by the peripheral substituents on the particular ligand.

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

If not otherwise indicated in the article, papers in issue 14 were published online on April 30, 2010

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