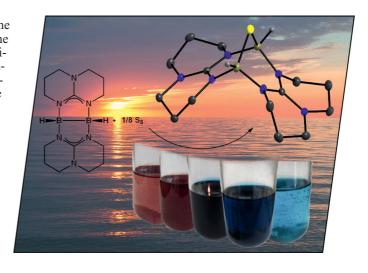


EurJIC is journal ChemPubSoc Europe, a union European of 16 chemical societies formed for the purpose of publishing high- quality science. All owners merged their national journals to form two leading chemistry journals, the European Journal of Inorganic Chemistry and the European Journal of Organic Chemistry.

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

COVER PICTURE

The cover picture shows the crystal structure of the product of oxidative insertion of sulfur into the B-B bond of the doubly base-stabilized diborane(4) $[HB(\mu-hpp)]_2$ (hpp = 1,3,4,6,7,8-hexahydro-2H-pyrimido[1,2-a]pyrimidinate). The colours observed in the course of this reaction are similar to those of the sunset at Flügge beach on the island Fehmarn shown in the background. While the sunset colours are caused by refraction of sunlight, oligosulfide intermediates such as S_8^{2-} and S_3^{-} are responsible for the colours visible during the reaction. The photographs of the flask show the reaction mixture in toluene solutions after 3, 12, 68 and 200 min and after 18 h. Details are discussed in the article by H.-J. Himmel et al. on p. 5201ff. The authors thank Andreas Schuster for the design of the cover picture.

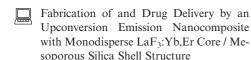


SHORT COMMUNICATION

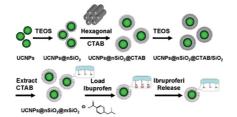
Nanocomposites for Drug Delivery

Y. Yang, Y. Qu, J. Zhao, Q. Zeng, Y. Ran, Q. Zhang, X. Kong,*

H. Zhang* 5195-5199



Keywords: Upconversion / Mesostructure / Drug delivery / Luminescence / Nanoparticles



Nanocomposites with a LaF₃:Yb,Er core and a mesoporous silica shell structure have been developed. These multifunctional photoluminescent composite materials were tested as drug carriers to investigate their drug storage/release properties. The green upconversion photoluminescence of these LaF₃:Yb³⁺, Er³⁺@nSiO₂@mSiO₂ nanocomposites is useful for tracking and monitoring the drug release.

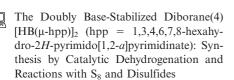
FULL PAPERS

B-B Bond Reactivity

N. Schulenberg, O. Ciobanu, E. Kaifer,

H. Wadepohl.

H.-J. Himmel* 5201-5210



Keywords: Boron / Hydrides / Sulfur / Guanidines



Catalytic dehydrogenation of $[H_2B(\mu-hpp)]_2$ leads to the doubly base-stabilized diborane(4) $[HB(\mu-hpp)]_2$. Oxidative sulfuration of the B-B bond in $[HB(\mu-hpp)]_2$ gives $[HB(\mu-hpp)]_2(\mu-S)$. A mixture of sulfuration and substitution products is formed for reactions with disulfides.

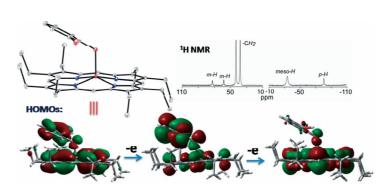
Catecholate Binding

DFT Investigation

A. Chaudhary, R. Patra, S. P. Rath* 5211-5221

Binding of Catechols to Iron(III)-Octaethylporphyrin: An Experimental and

Keywords: Catecholate binding / Hydrogen bonds / Spectroelectrochemistry / Density functional calculations / Structure elucidation



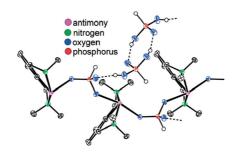
The synthesis, X-ray structures, and properties of new heme analogues Fe^{III} -(OEP)(L) (L: Hcat, 4-NO₂-Hcat, 4-tBu-Hcat, and sal) are reported, in which catechol binds in an η^1 -fashion as an axial

ligand. Spectroelectrochemical studies of one- and two-electron oxidations are also reported. Experimental findings are supported by DFT calculations.



Antimony and Bismuth Phosphites

The syntheses of molecular organoantimony and organobismuth phosphinate, phosphite and mixed phosphite-phosphonate complexes, where the central antimony or bismuth ion is stabilized by the NCN pincer-type ligand [2,6-(Me₂NCH₂)₂-C₆H₃] $^-$, are described. The decomposition pathway for bismuth phosphinate complexes is also reported.



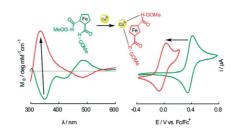
T. Svoboda, R. Jambor, A. Růžička, Z. Padělková, M. Erben, L. Dostál* 5222-5230

NCN Chelated Organoantimony(III) and Organobismuth(III) Phosphinates and

Phosphites: Synthesis, Structure and Reactivity

Keywords: Antimony / Bismuth / Phosphorus / Chelates / X-ray diffraction

Metal coordination enables control over ferrocene helicity. Synthesis of a 1,n'-ferrocenoyl—histidine conjugate is reported. Metal coordination to the His imidazoles drastically alters the helicity of the ferrocene group from a *P*-helical to an *M*-helical conformation.



Bioorganometallic Metal Chelators

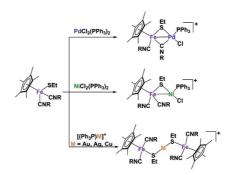
L.-Y. Cheng, Y.-T. Long,* H. Tian, H.-B. Kraatz* 5231–5238

Spectroscopic and Electrochemical Investigations into the Interactions of Metal Ions with a Ferrocenoyl-Histidine Peptide Conjugate

Keywords: Cyclic voltammetry / NMR spectroscopy / N ligands / Metal recognition / Chiral induction

Heteronuclear Complexes

A new series of heteronuclear thiolate iron complexes including heterobinuclear Fe-SEt-M (M = Pd or Ni) units and heterotrinuclear Fe-SEt-M-SEt-Fe (M = Au, Ag, or Cu) species with isocyanide ligands have been prepared, and their characterizations are also investigated.

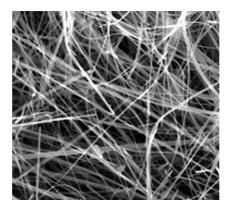


Synthesis of a New Family of Heteronuclear Thiolate Iron Complexes that Contain Isocyanide Ligands

Keywords: Transition metals / Heteronuclear complexes / Isocyanide ligands / Thiolate ligands

Vanadium Oxide Nanorods

We describe methods of hydrothermal synthesis of vanadium oxide nanorods produced from polycrystalline V_2O_5 . Analysis of the sensor properties of the nanorods revealed a significant response to triethylamine with a short response time, about 32 s, with a sensitivity of 30%.



Synthesis, Structure, and Sensor Properties of Vanadium Pentoxide Nanorods

Keywords: Vanadium oxides / Nanostructures / Hydrothermal synthesis / Resistivetype gas sensor

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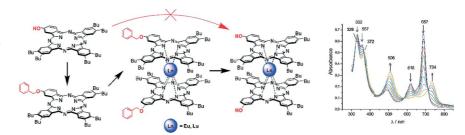
Sandwich Complexes

V. E. Pushkarev,* A. Yu. Tolbin,

N. E. Borisova, S. A. Trashin, L. G. Tomilova* 5254-5262

A₃B-Type Phthalocyanine-Based Homoleptic Lanthanide(III) Double-Decker π-Radical Complexes Bearing Functional Hydroxy Groups: Synthetic Approach, Spectral Properties and Electrochemical Study

Keywords: Phthalocyanines / Lanthanides / Sandwich complexes / Double-decker complexes / Pi radicals



Stable, phenolic hydroxy group containing π -radicals: Directly inaccessible hydroxy-substituted lanthanide(III)—bis(phthalocy-anine) sandwich complexes were obtained from the corresponding benzyloxy-pro-

tected precursors. A combination of spectroscopic and electrochemical methods unambiguously showed that these promising structural building blocks are stable π -radical species.

Phosphite-Substituted Chromophores

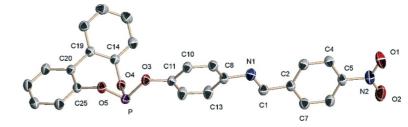
M. B. Murphy-Jolly,* S. B. Owens Jr., J. L. Freeman, G. M. Gray,*

C. M. Lawson,

D. P. Shelton 5263-5271

Syntheses, Crystal Structures and Photophysical Measurements of Phosphite-Substituted Schiff Base and Azobenzene Ligands

Keywords: Nonlinear optics / Schiff bases / Azo compounds / P ligands / Transition metals

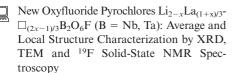


A novel class of phosphite-containing chromophores, O_2N -1- C_6H_4 -4-CH=N-1- C_6H_4 -4- $OP(OC_6H_4)_2$ (2) and O_2N -1- C_6H_4 -4-X=N-1- C_6H_4 -4- $OP(OC_{10}H_6)_2$ [X = CH (3), N (4)], and their transition-metal com-

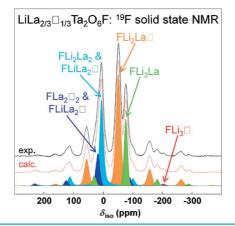
plexes, cis-Mo(CO)₄(2)₂ (5), PdCl₂(2)₂ (6), and cis-Mo(CO)₄(3)₂ (7) were synthesized and fully characterized. Second-order NLO properties of the ligands and complexes were investigated by HRS measurements.

Oxyfluoride Pyrochlores

C. Galven, C. Legein,* M. Body, J.-L. Fourquet, J.-Y. Buzaré, F. Le Berre,* M.-P. Crosnier-Lopez 5272–5283



Keywords: Pyrochlores / Local structure / X-ray diffraction / NMR spectroscopy / Electron microscopy / Structure elucidation



Average and local orders are characterized by XRD, TEM and 19 F solid-state NMR spectroscopy in two new oxyfluoride pyrochlores $\text{Li}_{2-x}\text{La}_{(1+x)/3}\square_{(2x-1)/3}\text{B}_2\text{O}_6\text{F}$ (B = Ta, Nb). The local environments of the F⁻ions, surrounded by Li⁺, La³⁺ ions and vacancies that occupy the same crystallographic site, are identified and quantified.

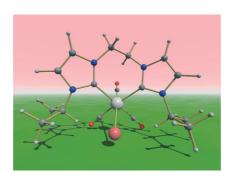
Biscarbene Complexes of Rhenium

O. Hiltner, F. J. Boch, L. Brewitz, P. Härter, M. Drees, E. Herdtweck, W. A. Herrmann,*

F. E. Kühn* 5284-5293

Bridged fac-Tricarbonylrhenium(I)—Biscarbene Complexes: Synthesis, Characterization, and Molecular Dynamics

Keywords: Carbonyl ligands / N ligands / Carbene ligands / Rhenium / Density functional calculations

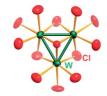


Six new Re complexes with chelating N-heterocyclic carbene (NHC) ligands were synthesized and fully characterized. NMR techniques and DFT calculations were used to explain their dynamical behavior.



Cluster Formation in IL

Ionic liquids (IL) are shown to be suitable reaction media for the convenient room-temperature synthesis of transition metal clusters. The new cluster compound $Sn[SnCl][W_3Cl_{13}]$ comprises a $[W_3Cl_{13}]^{3-}$ cluster with $C_{3\nu}$ symmetry as well as an unusual $[SnCl]^+$ cation. The compound is a paradigm of a material displaying extreme structural polarity.



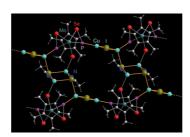
E. Ahmed,	M. Groh,	
M. Ruck*		5294-5297

Room-Temperature Synthesis of the Highly Polar Cluster Compound Sn[SnCl][W₃Cl₁₃]

Keywords: Cluster compounds / Ionic liquids / Tungsten / Merohedral twinning / Bijvoet differences

Coordination Polymers

New coordination polymers $[(Cp^*_2Mo_2P_2-Se_3)(CuX)_3(CH_3CN)]_n$ $(Cp^* = C_5Me_5; X = Cl, Br, I)$ have been prepared. A competitive coordination of Cu towards P and Se is observed that is completed by weak intermolecular Se···Se interactions to give 2D networks. Addition of P_4Se_3 as a competitive reagent gave $[(Cp^*_2Mo_2P_2Se_3)(P_4Se_3)-(CuX)_2]_n$, a new type of organometallic—inorganic hybrid polymer.



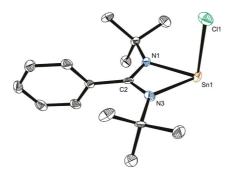
M. Bodensteiner, M. Dušek, M. M. Kubicki, M. Pronold, M. Scheer, J. Wachter,* M. Zabel 5298-5303

A New Building Block for Organometallic—Inorganic Hybrid Polymers: The Mixed Group 15/16 Element Ligand Complex $[Cp*_2Mo_2(\mu,\eta^{2:2}-PSe)_2(\mu-Se)]$ ($Cp*=C_5Me_5$)

Keywords: Phosphorus / Selenium / Copper / Coordination modes / Polymers

Tin Compounds

This article addresses the synthesis and characterization of amidinato-stabilized monomeric chlorostannylene. Furthermore, the chlorostannylene was converted into its amide and triflate derivatives by substitution reaction. To verify the Lewis basicity of chlorostannylene it was treated with Fe₂(CO)₉, whereupon it formed the Lewis acid—base adduct.



S. S. Sen, M. P. Kritzler-Kosch, S. Nagendran, H. W. Roesky,* T. Beck, A. Pal, R. Herbst-Irmer 5304–5311

Synthesis of Monomeric Divalent Tin(II) Compounds with Terminal Chloride, Amide, and Triflate Substituents

Keywords: Tin / N ligands / NMR spectroscopy / X-ray diffraction

Alkene Epoxidation

Experimental and theoretical studies on the epoxidation of a range of alkenes with aqueous $\rm H_2O_2$ and dititanium-containing 19-tungstodiarsenate(III) $\rm [Ti_2(OH)_2As_2-W_{19}O_{67}(H_2O)]^{8-}$ (1) as catalyst revealed a mechanism that involves a reversible interaction between $\rm H_2O_2$ and the Ti–OH group of 1 to produce a titanium hydroperoxo complex followed by electrophilic oxygen atom transfer to the alkene.



Epoxidation of Alkenes with H₂O₂ Catalyzed by Dititanium-Containing 19-Tungstodiarsenate(III): Experimental and Theoretical Studies

gsor-

Keywords: Polyoxometalates / Titanium / Epoxidation / Density functional calculations / Hydrogen peroxide

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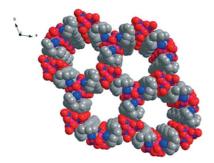
Ln Metal-Organic Frameworks

Q. Wang, K.-Z. Tang, W.-S. Liu, Y. Tang,* M.-Y. Tan 5318-5325



Synthesis, Crystal Structures, and Luminescent Properties of Noninterpenetrating (6,3) Type Network Lanthanide Metal—Organic Frameworks Assembled by a New Semirigid Bridging Ligand

Keywords: Lanthanides / Metal—organic frameworks / Semirigid bridging ligands / Homometallic compounds / Luminescence



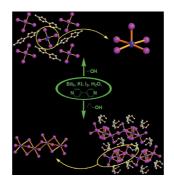
A new semirigid bridging ligand, 2,5-di methyl-1,4-bis[{(2'-benzylaminoformyl)-phenoxy}methyl]benzene (L), was designed to assemble a series of rare noninterpenetrating (6,3) type network metal—organic frameworks with lanthanide cations, namely [{Ln₂(NO₃)₆L₃}·(H₂O)₂·(CHCl₃)]_n. The luminescent properties of the Sm^{III}, Eu^{III}, Tb^{III}, and Dy^{III} MOFs were studied in detail.

Viologen Iodobismuthates



Iodine-Induced Solvothermal Formation of Viologen Iodobismuthates

Keywords: Semiconductors / Solvothermal synthesis / Bismuth / Iodine / Viologens



Four viologen iodobismuthate complexes were generated from the solvothermal reactions of BiI₃, I₂, KI, 4,4'-bipyridine, alcohols and a small amount of water in MeCN. The optical, electrical conductivity and dielectric properties of the products were investigated.

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

If not otherwise indicated in the article, papers in issue 32 were published online on November 2, 2010

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