Applied Organometallic Chemistry

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Section: Bioorganometallic Chemistry

The anion-binding properties of ferrocene-based receptors bearing phenol group are evaluated for F⁻, Cl⁻, Br⁻, I⁻, AcO⁻ and $H_2PO_4^-$ by UV-vis, ¹H NMR titration and cyclic voltammetry experiments. Results indicate that the anion binding abilities can be effectively tuned by introducing nitro group in the ortho position of phenyl ring of the receptors, and the most obvious effect is for $H_2PO_4^-$.

X. F. Shang, H. Lin, X. F. Xu, P. Jiang and H. K. Lin* 821–825

Ferrocene-based derivative bearing phenol group recognitive sites: efficient $H_2PO_4^-$ receptor

Retracted.

M. M. Naseer* and Z. H. Chohan 826-835

Synthesis, characterization and reactivity towards first-row d-transition metals and biological significance of new pyridinyl derived N-substituted sulfonamides

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Identification statement

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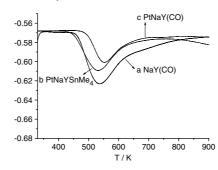
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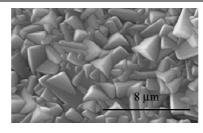
Section: Materials, Nanoscience and Catalysis

The grafted Me₃Sn-complex alters the electronic properties of Pt atoms, and thus results in decreased CO desorption temperature on the grafted sample.



Preparation and catalytic properties of a bimetallic Sn-Pt complex in the supercages of NaYzeolite by use of surface organometallic chemistry

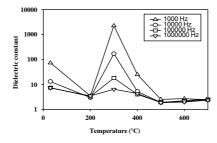
Synthesized continuous NaA membranes were prepared using a 0.5 μ m NaA crystal seed via vacuum seeding. The optimum conditions were 363 K synthesis temperature for 15–20 min via microwave heating. The flux and separation factor obtained were 1.56 kg/m² h and 1760.52, respectively, for the substrate without an intermediate layer.



Effects of synthesis parameters on zeolite membrane formation and performance by microwave technique

Interestingly, the substrate with an intermediate layer showed better flux and separation factor at 1.69 kg/m² h and 6532.72, respectively.

A perovskite lead zirconate was synthesized, using lead glycolate and sodium tris (glycozirconate) as the starting precursors, by the sol-gel process. The structure obtained was the orthorhombic form when calcined at low temperature at 300 °C for 1 h and transformed to the monoclinic and cubic forms of the perovskite phase at higher temperatures above the Curie



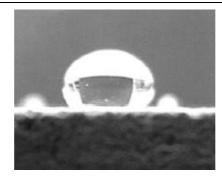
N. Tangboriboon, A. Jamieson, A. Sirivat and S. Wongkasemjit* 849-857

A novel route to perovskite lead zirconate from lead glycolate and sodium tris(glycozirconate) via the sol-gel process

temperature as verified by X-ray data. The lead zirconate synthesized and calcined at 300 °C for 1 h has the highest dielectric constant, the highest electrical conductivity, and the dielectric loss tangent of 2267, 3.058 $\times~10^{-4}~(\Omega.m)^{-1}$, and 2.484 at 1000 Hz, respectively.

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The surface of quarry stone was modified by continuous plasma polymerization of hexamethyldisiloxane. The hydrophilic surface of the quarry stone was made hydrophobic and impermeable to water. Three different reaction times were analyzed. Contact angle and FT-IR analyses show that the hydrophobic character of the surface is due to methyl groups on the surface. The change in the contact angle with temperature and the wetting temperature ($T_{\rm w}$) were measured.



Surface modification of quarry stone by hexamethyldisiloxane plasma treatment

Three aromatic diamines, 2,6-di-(4-aminophenyloxy) toluene (A), 2,2'-di-[4-(4-aminophenyloxy) phenyl] propane (B) and 4,4'-di (4-aminophenyloxy) biphenyl (C), were obtained by

Synthesis and physicochemical studies of ferrocene-containing materials

reacting 2,6-dihydroxytoluene, bisphenol-A and 4,4'-dihydroxybiphenyl with *p*-nitrochlorobenzene. The synthesized diamines were then polymerized with 1,1'-ferrocenedicarboxylic acid chloride with low-temperature polycondensation to produce organometallic aromatic polyamides (aramids). The synthesized monomers and polymers were characterized by their solubilities, elemental analysis, FTIR spectroscopy and ¹H-NMR spectroscopy. The inherent viscosities, differential scanning calorimetry and thermogravimetry were also used for polymer characterization.

The new ansa-complexes were used on ethylene homopolymerization, α -olefin homopolymerization, ethylene/ α -olefin copolymerization. When ^tBu was introduced into the para position on phenyl groups in (Ph₂C(Cp)(Ind) ZrCl₂), the activity and the incor-

3. R= ^tBu, M=Ti 4. R= ^tBu, M=Zr 5. R= ^tBu, M=Hf 6. R= MeO, M=Zr 7. R= MeO, M=Hf

X. Yang, Y. Zhang and J. Huang* 870-879

α-olefin homopolymerization and ethylene/1-hexene copolymerization catalysed by novel ansa-group IV complexes/MAO system

poration of 1-hexene on copolymerization were both increased.

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Section: Main Group Metal Compounds

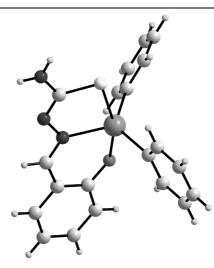
Bis(N,N-dialkyldithiocarbamato)antimony(III) alkylenedithiophosphate of the type $[R_2NCS_2]_2SbS(S)\overline{POGO}$ [where $NR_2 = N(CH_3)_2$, $N(C_2H_5)_2$ and $N(CH_2)_4$; $G = -CH_2 - C(C_2H_5)_2 - CH_2 -$, $-CH_2 - C(CH_3)_2 - CH_2 -$, $-CH(CH_3) - CH(CH_3) - CH(CH_3)$ and $-C(CH_3)_2 - C(CH_3)_2 -$] were synthesized and characterized by physico-chemical, spectral [UV, IR and NMR (1H , ^{13}C and ^{31}P)] and thermal (TG, DTA and DSC) analysis. The TG decomposition step of the complex indicated the formation of Sb_2S_3 as final product. The melting point of these complexes was confirmed by DSC analysis. These complexes were screened for antibacterial and antifungal activity using the disk diffusion method. All the complexes showed good affinity as antibacterial and antifungal agents, which increased with increasing concentration.

$$R_2N$$
 S_3
 S_6
 S_1
 S_6
 S_1
 S_6
 S_1

H. P. S. Chauhan* and U. P. Singh 880–889

Synthetic, spectral, thermal and antimicrobial studies on some bis(N,N'-dialkyl-dithiocarbamato)antimony(III) alkylenedithiophosphates

A series of organotin compounds have been characterised spectroscopically and by X-ray crystallography. The tin atom is penta-coordinated in each case and the majority of structures associate via eight-membered {N-C-N-H}₂ synthons. The efficacy of these compounds against a range of bacteria, fungi, plant pathogens and human cancer cell lines is also described.



M. S. Sarma, S. Mazumder, D. Ghosh, A. Roy*, A. Duthie and E. R. T. Tiekink 890 – 905

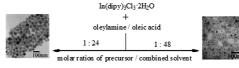
Synthesis, spectroscopic characterization and biocidal activity of some diorganotin(IV) complexes of salicylaldehydethiosemicarbazones and related ligands. Molecular and supramolecular structures of $[R_2Sn(OArCH=N-N=CSNH_2)]$, where R=Me, Ph and $Ar=-C_6H_4$, $-C_6H_3(5-Br)$ and $C_6H_3(5-CI)$, and of $[Me_2Sn\{OC_6H_3(5-Br)CH=N-N=CSNH_2\}OH_2]$

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	Synthesis of inorganic materials
Book Review	N. T. Lucas 907 – 908
	Carbon-rich compounds: from molecules to materials

Section: Materials, Nanoscience and Catalysis

By thermo-decomposition of the $ln(dipy)_3Cl_3\cdot 2H_2O$ (dipy = α , α -dipyridyl) precursor in the presence of stabilizing surfactant, quasi-



P. Zhu, W. Wu, J. Zhou, W. Zhang* 909-912

Preparation of size-controlled In_2O_3 nanoparticles

monodisperse and size-controlled In_2O_3 nanoparticles were synthesized. The PL emission at 378 nm was found at room temperature.

Section: Main Group Metal Compounds

Three diorganotin(IV) complexes of N-(2-hydroxy-4-nitrophenyl)-salicylidene-imine derivatives have been prepared and the new compounds characterized by C, H, N analysis, mass, IR, $^1\text{H-NMR}$ and $^{13}\text{C-NMR}$ spectroscopy. The crystal structure of (C₆H₅)₂Sn(OC₆H₃OC₂H₅CH=NC₆H₃NO₂O) was characterized by single crystal X-ray diffraction analysis. A coordination geometry nearly half-way between trigonal-bipyramidal- and square pyramidal- arrangement (trigonality index: $\tau=0.49$) was inferred. In the solid state, $\pi-\pi$ interactions between the aniline fragments of neighbouring molecules exist with a centroidal distance of 3.734 (5) Å and a slip angle of 19.1°.

Synthesis and characterization of diorganotin(IV) complexes of Schiff bases with ONO-type donors and crystal structure of [N-(2-hydroxy-4-nitrophenyl)-3-ethoxysalicylideneiminato]diphenyltin(IV)