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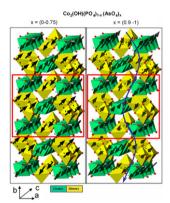
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Regular Articles

Heat capacity and neutron diffraction studies on the frustrated magnetic $Co_2(OH)(PO_4)_{1-x}(AsO_4)_x \ [0 \le x \le 1]$ solid solution

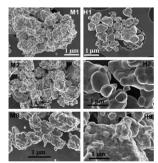
I. de Pedro, J.M. Rojo, J. Rodríguez Fernández, J. Sanchez Marcos, M.T. Fernández-Díaz and T. Rojo *page 1*



Magnetic structures of $\text{Co}_2(\text{OH})(\text{PO}_4)_{1-x}(\text{AsO}_4)_x \ [0 \le x \le 1]$. The ordering of the magnetic moments of Co^{2+} is in *c* direction for the two crystallographic positions (dimers and chains) in all compounds. The unit cell is surrounded by a red line.

Room temperature mechanosynthesis of the $La_{1-x}Sr_xMnO_{3\pm\delta}$ ($0 \le x \le 1$) system and microstructural study

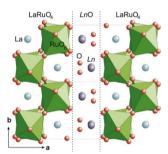
M.J. Sayagués, J.M. Córdoba and F.J. Gotor page 11



SEM micrographs corresponding to milled (M) and Heated (H) samples: LaMnO_{3±δ} (M1 and H1) La_{0.75}Sr_{0.25}MnO_{3±δ} (M2 and H2,) and SrMnO_{3±δ} (M8 and H8).

Regular Articles—Continued

Single crystalline and rare earth substituted La₂RuO₅ investigated by x-ray diffraction and EXAFS spectroscopy S. Riegg, A. Reller and S.G. Ebbinghaus page 17

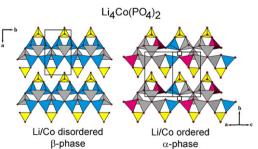


The crystal structure of $La_{2-x}Ln_xRuO_5$ (Ln = Pr, Nd, Sm, Gd, Dy) is shown viewed along the **c**-axis. The alternating stacking of LaRuO₄ and LnO layers leads to the formation of zig-zag layers of corner sharing RuO₆ octahedra. The La sites in the LaRuO₄ layers are represented by light blue spheres, while the La/Ln sites in the LnO layers are colored dark blue. EXAFS investigations reveal a cationic ordering with roughly 65% of the substituting Ln ions occupying the LnO layers.

Synthesis, structures and properties of the new lithium cobalt(II) phosphate $Li_4Co(PO_4)_2$

R. Glaum, K. Gerber, M. Schulz-Dobrick, M. Herklotz, F. Scheiba and H. Ehrenberg

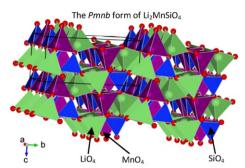
page 26



The complex formation and decomposition behavior of Li₄Co (PO₄)₂ with temperature has been elucidated. The crystal structure of its α -phase was determined from single crystal data, HT-XRPD allowed derivation of a structure model for the β -phase. Both modifications belong to the Li₃PO₄ structure family.

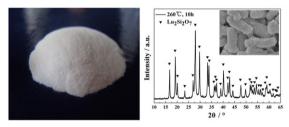
Continued

Crystal chemistry of the *Pmnb* polymorph of Li₂MnSiO₄ R.J. Gummow, N. Sharma, V.K. Peterson and Y. He *page 32*



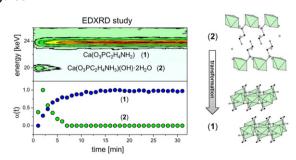
Polyhedral representation of the crystal structure of Li_2MnSiO_4 in the *Pmnb* space group. LiO₄, MnO₄ and SiO₄ tetrahedra are shown in green, purple and blue, respectively.

Hydrothermal synthesis of lutetium disilicate nanoparticles Xiaoping Tang, Yanfeng Gao, Hongfei Chen and Hongjie Luo page 38



An image for the as-prepared $Lu_2Si_2O_7$ powders (left) and XRD pattern (right) (inset shows the SEM graph of powders).

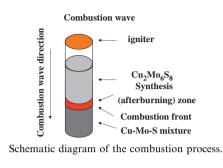
High-throughput and in situ EDXRD investigation on the formation of two new metal aminoethylphosphonates – $Ca(O_3PC_2H_4NH_2)$ and $Ca(OH)(O_3PC_2H_4NH_3) \cdot 2H_2O$ Corinna Schmidt, Mark Feyand, André Rothkirch and Norbert Stock *page 44*



The detailed in situ energy dispersive X-ray diffraction (EDXRD) investigation on the formation of the new inorganic–organic hybrid compound $Ca(O_3PC_2H_4NH_2)$ leads to the discovery of a new crystalline intermediate phase. Both crystal structures were elucidated using X-ray powder diffraction data.

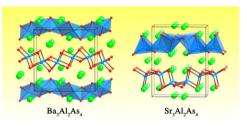
Ultra fast elemental synthesis of high yield copper Chevrel phase with high electrochemical performance

Gregory Gershinsky, Ortal Haik, Gregory Salitra, Judith Grinblat, Elena Levi, Gilbert Daniel Nessim, Ella Zinigrad and Doron Aurbach page 50



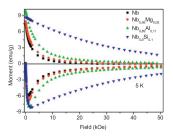
Synthesis and structural characterization of the ternary Zintl phases $AE_3Al_2Pn_4$ and $AE_3Ga_2Pn_4$ (AE = Ca, Sr, Ba, Eu; Pn = P, As)

Hua He, Chauntae Tyson, Maia Saito and Svilen Bobev page 59



 $AE_3Al_2Pn_4$ and $AE_3Ga_2Pn_4$ (AE=Ca, Sr, Ba, Eu; Pn=P, As) crystallize in two different structures—Ca₃Al₂P₄, Sr₃Al₂As₄, Eu₃Al₂P₄, Eu₃Al₂As₄, Ca₃Ga₂P₄, Sr₃Ga₂P₄, Sr₃Ga₂As₄, and Eu₃Ga₂As₄, are isotypic with the previously reported Ca₃Al₂As₄ (space group C2/c (No. 15)), while Ba₃Al₂P₄ and Ba₃Al₂As₄ adopt a different structure known for Na₃Fe₂S₄ (space group *Pnna* (No. 62)). The polyanions in both structures are made up of $TrPn_4$ tetrahedra, which by sharing common corners and edges, form $\frac{2}{\infty}[TrPn_2]^{3-}$ layers in the former and $\frac{1}{\infty}[TrPn_2]^{3-}$ chains in Ba₃Al₂P₄ and Ba₃Al₂As₄.

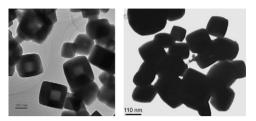
Superconductivity in quaternary niobium oxynitrides containing main group elements (M = Mg, Al, Si) Y. Ohashi, S. Kikkawa, I. Felner, M.I. Tsindlekht, D. Venkateshwarlu, V. Ganesan and J.V. Yakhmi page 66



The doped Si oxide accompanied with some amount of cation vacancy in cubic NbN lattice induces relatively large magnetic hysteresis on isothermal hysteresis loops at 5 K of the dc magnetization up to 5 T among the four niobium oxynitrides containing main group elements, $Nb_{1.00}(N_{0.98}O_{0.02})$; $(Nb_{0.95}Mg_{0.05})(N_{0.92}O_{0.08})$; $(Nb_{0.89}Al_{0.11})(N_{0.84}O_{0.16})$, and $(Nb_{0.87}Si_{0.09}\Box_{0.04})(N_{0.87}O_{0.13})$.

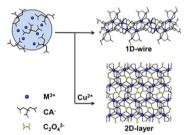
Effect of solvents on morphologies of PbTe nanostructures: Controllable synthesis of hollow and solid PbTe nanocubes by a solvothermal method

Wenzhong Wang, Lijuan Wang and Qing Zhou page 72



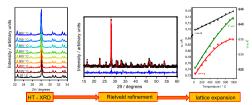
A facile solution-phase route has been developed to synthesize hollow and solid PbTe nanocubes. The possible growth mechanism of hollow and solid PbTe nanocubes was discussed in detail.

New metal-organic frameworks of $[M(C_6H_5O_7)(C_6H_6O_7)(C_6H_7O_7)(H_2O)]$. H₂O (M = La, Ce) and $[Ce_2(C_2O_4)(C_6H_6O_7)_2]$. 4H₂O Sheng-Feng Weng, Yun-Hsin Wang and Chi-Shen Lee *page* 77



 $[M(C_6H_5O_7)(C_6H_6O_7)(C_6H_7O_7)(H_2O)]$. H₂O (M = La(1a), Ce(1b)) and $[Ce_2(C_2O_4)(C_6H_6O_7)_2]$. 4H₂O (2)—with 1D and 2D structures were synthesized and characterized.

Thermal expansion of Ba₂ZnSi₂O₇, BaZnSiO₄ and the solid solution series BaZn_{2-x}Mg_xSi₂O₇ ($0 \le x \le 2$) studied by high-temperature X-ray diffraction and dilatometry Marita Kerstan, Matthias Müller and Christian Rüssel *page 84*



XRD-patterns of $Ba_2ZnSi_2O_7$ were recorded at different temperatures (left). For each XRD-pattern a Rietveld-refinement was performed, the image in the middle shows the XRD-pattern measured at room temperature (circles), the Rietveld calculation (red line) and the difference between them (blue line). The lattice parameters derived hereof were plottet against the temperature and fitted to a polynomial (right picture). From those polynomials the lattice expansion was calculated.

Structural variation and optical properties of ZnO-LiGaO₂ pseudo-binary system

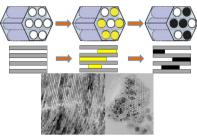
Takahisa Omata, Masao Kita, Kosuke Tachibana and Shinya Otsuka-Yao-Matsuo *page 92*



The structure of ZnO varied upon alloying with LiGaO₂.

Highly ordered magnetic mesoporous silicas for effective elimination of carbon monoxide Jiho Lee and Jeong Ho Chang

page 100



Strategy for the preparation of highly abundant Fe nanoparticle embedded MS catalyst by hydrogen reduction process and HR-TEM images of cross-sectional and top view.

Luminescence of the elpasolite series $M_2^I M^{II} M Cl_6$ ($M^I = Cs$, Rb; $M^{II} = Li$, Na; M = Lu, Y, Sc, In) doped with europium using synchrotron radiation excitation

Peter A. Tanner, Chang-Kui Duan, Guohua Jia and Bing-Ming Cheng *page 105*

24000 23500 22500 22500 22500 22500 0.8 0.9 1.0 1.1 Latice parameter (nm)

Luminescence of Eu^{3+} and Eu^{2+} in elpasolite hosts under synchrotron radiation is observed and assigned.

Corrigendum

Corrigendum to "Oxygen excess in the "114" cobaltite hexagonal structure: The ferrimagnet CaBaCo₄O_{7.50}" [J. Solid State Chem. 184 (2011) 2588–2594] V. Pralong, V. Caignaert, T. Sarkar, O.I. Lebedev, V. Duffor and B. Raveau *page 109*

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