

paramagnetic to 4.2°K. The evidence suggests that PrAl₂ and CaAl₂ are electronically similar with regard to band structure and both differ in band structure from that in GdAl₂.

Crystal Structure and Magnetic Properties of Ba₂Ni₃F₁₀. M. LEBLANC, G. FERREY, AND R. DE PAPE, Laboratoire des Fluorures et Oxyfluorures Ioniques, Route de Laval, 72017 Le Mans Cedex, France. Ba₂Ni₃F₁₀ is monoclinic (space group *c2/m*) $a = 18.542(7) \text{ \AA}$, $b = 5.958(2) \text{ \AA}$, $c = 7.821(3) \text{ \AA}$, $\beta = 111^\circ 92(10)$. Ba₂Co₃F₁₀ and Ba₂Zn₃F₁₀ are isostructural. The structure has been refined from 995 reflections by full-matrix least-squares refinement to a weighted *R* value of 0.048 (unweighted *R*: 0.047). The three-dimensional network can be described either by complex chains connected to each other by octahedra sharing corners or with a 18L dense packing sequence. The basic unit (Ni₃F₁₀)⁴⁻ is discussed and compared to a different one existing in Cs₄Mg₃F₁₀. Antiferromagnetic properties of Ba₂Ni₃F₁₀ ($T_N = 50 \text{ K}$) are described.

Infrared Spectrum of Several Double Oxides with Corundum Structure Resulting from the Transformation of Lacunar γ Phases with a Spinel Structure. B. GILLOT, F. BOUTON, F. CHASSAGNEUX, AND A. ROUSSET, Laboratoire de Recherches sur la Réactivité des Solides, BP 138, 21004 Dijon, Cedex, France. When aluminum or chromium is substituted by Fe³⁺ ions in α -Fe₂O₃, all the ir bands gradually shift toward high frequencies. Alternatively, for the α phases of type (Fe₂Cr_{4-y}Al_y)O₉ the transition occurs sharply for a composition *y* close to 2. For α phases substituted by (Fe_{6-y}Cr_y)O₉-type chromium a linear variation of frequency with chromium content is observed. From ir data it has been shown that, under given temperature and time conditions, an α phase less rich in chromium than the initial product could be obtained by oxidizing iron chromite. The ir spectrum of the oxidation of pure magnetites whose size is between 1400 and 15 000 Å evolves versus the latter to yield either the γ -Fe₂O₃ or the α -Fe₂O₃ phase which can be formed from γ -Fe₂O₃ or by direct oxidation of Fe₃O₄.

Preparation and Mössbauer Effect of Cr_{1-x}Fe_xOOH (0 ≤ x ≤ 1.0) with the InOOH-Type Structure. K. YABUTA, N. KINOMURA, M. SHIMADA, F. KANAMARU, AND M. KOIZUMI, The Institute of Scientific & Industrial Research, Osaka University, Osaka 565, Japan. The complete solid solutions of Cr_{1-x}Fe_xOOH with the InOOH-type structure were synthesized under high pressures. They were antiferromagnetic with Néel temperatures 570 K for FeOOH, 454 K for Cr_{0.2}Fe_{0.8}OOH, 332 K for Cr_{0.4}Fe_{0.6}OOH, and 160 K for Cr_{0.6}Fe_{0.4}OOH determined by the Mössbauer effect measurements.

Synthese et Caracterisation Cristallographique et Physique d'une Serie de Composes ACu₃Ru₄O₁₂ de Type Perovskite. M. LABEAU, B. BOCHU, J. C. JOUBERT, AND J. CHENAVAS, Laboratoire de Génie Physique, ENSIEG, BP 46, 38402, Saint-Martin-D'Herès, France. A series of ruthenates with the perovskite-like arrangement has been synthesized. The general formula is (ACu₃)(Ru₄)O₁₂ where (ACu₃) and (Ru₄) occupy the *A* and *B* sites of the ideal ABO₃ perovskite structure, respectively; $A = \text{Na}^+, \text{Ca}^{2+}, \text{Sr}^{2+}, \text{Cd}^{2+}, \text{La}^{3+}, \text{Pr}^{3+}, \text{and Nd}^{3+}$. The change of the *A* site charge is balanced by a change of the ruthenium valence (+3, +4, or +5). The stability of the structure is strongly related to the relative size of the cations on the dodecahedral and octahedral sites. The magnetic susceptibility of the compounds with $A = \text{Na}, \text{Ca}, \text{and La}$ is temperature independent above 70°K (for Ca), 120°K (for La) or 220°K (for Na) (Pauli paramagnetism). Resistivity measurements made on sintered samples between 77 and 300°K confirm the metallic nature of these compounds.

Crystal Structure of Na₃PCr₃O₁₃ · 3H₂O, A New Type of Chromophosphoric Anion. M. T. AVERBUCH-POUCHOT, A. DURIF, AND J. C. GUITEL, Laboratoire de Cristallographie, CNRS 166X, 38042 Grenoble Cedex, France. We describe chemical preparation and crystal structure of a new sodium phosphochromate: Na₃PCr₃O₁₃ · 3H₂O. This salt is orthorhombic with $a = 11.72(3)$, $b = 14.89(3)$, $c = 16.59(3) \text{ \AA}$, and $Z = 8$, Space group: *Pbc2₁*. The final *R* value is 0.063 for 2133 independent reflexions. The main feature of this atomic arrangement is the existence of a new type of chromophosphoric anion: PCr₃O₁₃. Infinite chains of NaO₆ run along the *a* direction. A survey of some phosphochromates previously described by the authors is given.

Semiconductivity, Optical Properties, and EPR Spectra of Phenothiazine Derivatives in the Solid State. A. ORTIZ, J. I. FERNÁNDEZ-ALONSO, A. PARDO, AND J. LLABRES, Departamento de Química-Física y Química Cuántica, Universidad Autónoma de Madrid, Cantoblanco, Madrid, 34, Spain. The optical, EPR, and electrical properties of phenothiazine derivatives have been investigated as a function of R₂ substituents. Diffuse reflectance spectra show a charge transfer complex transition between 600 and