ABSTRACTS OF FORTHCOMING ARTICLES

novel phase, "X." (Topotactic rearrangement of P to the known acicular polymorph of its benzoxiodole isomer, 3-oxo-3H, 2, 1-benzoxiodol-1-yl *m*-chlorobenzoate, occurs only at elevated temperatures, \sim 55°C). In a much slower, consecutive transformation, during further continuous exposure to X-rays, "X" is photochemically reduced to *o*-iodobenzoic acid (C), which also is formed as a single crystal phase. The resulting single crystal domains of D and E in decomposed pseudomorphs of P are mutually oriented in a manner which reflects their strikingly similar crystal structures. The structure of D has been determined by single crystal X-ray analysis in order to compare the aligned molecular packing arrangements of D and C, and P and D. The mutual alignment of P and E is also presented in this report. All of these topotactic products are aligned so as to conserve the point group symmetry of the parent crystal, P (conservative twinning).

Characterization of CoO-ZnO Solid Solutions. F. PEPE, M. SCHIAVELLO, AND G. FERRARIS. Istituto di Chimica Generale ed Inorganica, Universita di Roma, Rome, Italy. Lattice parameters, magnetic susceptibilities (over the temperature range 98-298°K) and reflectance spectra (uv and visible) of solid solutions of CoO-ZnO up to 10% molar CoO have been studied. It is shown that true solid solution is achieved and that Co²⁺ ions substitute for the Zn²⁺ ions in tetrahedral positions. The effect of the addition of Co²⁺ is to increase the *a* lattice parameter and to decrease the *c/a* ratio. The magnetic moment of Co²⁺ has been found to be $4.54 \times 0.05 \mu_{\rm B}$ by extrapolation. A discussion of the structural and magnetic data in terms of the presence of a C_{3v} effective symmetry is given.

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