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Coordination Compounds of Phosphoryl Containing Pendant Macrocyclic Ligands with Alkali and Transition Metals

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COORDINATION COMPOUNDS OF PHOSPHORYL CONTAINING PENDANT MACROCYCLIC LIGANDS WITH ALKALI AND TRANSITION METALS

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Electroneutral phosphoryl-containing pendant polyazamacrocycles L^I-L^{IV}, proved to be efficient complexing agents for alkali and transition metal ions

The ligands involve different coordination sites: the "hard" base, oxygen atoms, and less "hard" nitrogen atoms in aminogroups. The stability constants (K) of [ML] + complexes (M is "hard" acid Li, Na, K, Cs ions) were detected in the THF. The discrimination coefficients, which characterize the selectivity of ligands, were calculated. The value of K was found to depend on the macrocycle cavity size and the pendant chain length.

Mono- and polynuclear Co(II), Ni(II) and Cu(II) complexes with LI-LIV ligands were synthesized. In the binuclear complexes the ligands act as bridges between two cations and all P=O and amino groups from coordinate bonds, whereas in the case of MLX, complexes only some of the donor groups are involved in coordination. The electronic structure and coordination polyhedrons geometry of metal ions is determined by the manner of ligand coordination.