## 07-Crystallography of Organometallic and Coordination 232 Compounds

This kind of coordination has been observed for coordination compounds of EDTA with different transition metals obtained in presence of water (Solans, X., Font-Altaba, M. & Garcia-Oricain, J. (1984). Afinidad 41, 572-578).

PS-07.04.35 INVESTIGATION OF CRYSTAL STRUCTURE REGULARITIES OF M4[Ru(NO2)6].nH2O ( M=Li,Na,K;

n=12,2,0). By S.A. Gromilov<sup>\*</sup>, V.A. Emelyanov, V.I. Alekseev, I.A. Baidina, A.V. Belyaev. Institute of Inorganic Chemistry, Russian Academy of Sciences, Siberian Branch, Novosibirsk, Russia.

Data on crystal structures of nitrocompounds of Ru(II) - [Ru(NH<sub>3</sub>)<sub>5</sub>NO<sub>2</sub>]·Cl·H<sub>2</sub>O ( Bottomley F., J.Chem.Soc., Dalton, 1972, 19, 2148-2152) and  $M_2[RuNO(NO_2)_4OH]$ , where M is Na (Simonsen S.H. et al., J. Inorg. Chem., 1965, 27, 309-320), or S.H. et al., J. Inorg. Chem., 1965, 27, 309-320), or K (Bokij G.B. et al., J.Struct Chem., 1962, 3, 2, 163-172) are known elsewhere. A procedure of synthesis of hexanitroruthenates(II) of Li (A), Na (B) and K (C) has been developed by us.  $A - \text{Li}_4[\text{Ru}(\text{NO}_2)_6] \cdot 12\text{H}_2^{\text{O}}$   $B - \text{Na}_4[\text{Ru}(\text{NO}_2)_6] \cdot 2\text{H}_2^{\text{O}}$ 

$$C = K_4[Ru(NO_2)_6]$$

X-ray investigation of single and polycrystals has been carried out.

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	a,Å α,deg	β	γ	v,Å <sup>3</sup> z	Simm. Sp.gr.	R,%
A	11,749 90	11,749 90	16,807 90	2320.0	Tetr. I4	3,8
В	8,105 90	8,723 91,87	8,931 90	631,1 2	Monocl. P2 <sub>1</sub> /n	2,9
С	8,595 52,23	8,595 52,23	8,595 52,23	367,0 1	Rmbhdr. R3	3,2
Cryatal		structures of		f the	compounds	

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structures Crystal investigated are based on the complex anions Ru(NO2)4- forming a distorted rhombohedral sub-cell, alkaline metal cations and molecules of crystallized water. The Ru atom has an octahedral environment formed by the N atoms of the nitrite groups. The mean Ru-N distance is 2.065Å. In the crystal structures all 0 atoms of the nitrite groups and water molecules form polyhedra around the cations to link them with each In Athese are other and with the complex anions. octahedra and tetrahedra with mean Li-O distances of 2.02 and 1.95Å, respectively. In B and C the Na and K atoms are coordinated to light O atoms with mean bond distances: Na-0=2.31-2.71, K-0=2.85-3.24Å.

A comparative analysis of the orientations of complex anions and nitrite groups about the probable axe of the 3-D order has been performed.

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PS-07.04.36 MIXED-LIGAND DIETHYLDITHIOCARBAMATES OF Zn(II) AND Cd(II): PREPARATION AND CRYSTAL STRUCTURES. By L.A. Glinskaya\*, R.F. Klevtsova, S.M. Zemskova, S.V. Larionov. Institute of Inorganic Chemistry, Russian Academy of Sciences, Siberian Branch, Russia.

One of the most interesting classes of volati-le coordination compounds of practical use ars diethyldithiocarbamates of metals (Larionov S.V. Zh.Neorg.Chim., 1979,24,6,1446). The mixed-li-gand complexes of Zn(II) and Cd(II) were obta-ined, namely four compounds of CdL<sub>2</sub> and three compounds of ZnL<sub>2</sub>(L=(C<sub>2</sub>H<sub>5</sub>)<sub>2</sub>NCS<sub>2</sub><sup>-</sup>) with nitro-gen heterocycles (Phen, Bipy, Im) and En: I-CdL<sub>2</sub>Phen; II-CdL<sub>2</sub>(2,2'-Bipy); III-CdL<sub>2</sub>Im; IV-[ZnEn3][CdL<sub>3</sub>]<sub>2</sub>; V=ZnL<sub>2</sub>(2,2'-Bipy); VI-[ZnL<sub>2</sub>]<sub>2</sub>(4,4'-Bipy). The crystal data are given in the Table.

	Sp.gr.	a, d	b,ß	c,8	Z	CN
I		11.044 100.38	15.172 99.79	16.967 101.40	4	6(4S+2N)
II	C2/c	18.103	8.381 105.97	16,965	4	6(4S+2N)
III IV V VI	I4.cd Pnaa	20.820 20.348 17.246 17.411	8.576	11.520 28.744 6.792 22.161	4	5(4S+1N) GN, 6S 6(4S+2N) 5(4S+1N)

The crystal structures I-V are built of disc-rete mononuclear and VI - binuclear molecules rete mononuclear and VI - binuclear molecules linked by Van-der-Waals contacts. The central atom environments in the complexe IV form dif-ferent coordination polyhedra: an octahedron and a trigonal prism. The crystal chemical characteristics of complexes studied have been treated following determinations of new mixedligand complexes and literature data.

PS-07.04.37 'CLOSEST PACKING' IN INORGANIC COOR-DINATION CRYSTALS. By N.V. Podberezskaya\*, T.S. Yudanova, S.A. Magarill, E.N. Ipatova, G.V. Romanenko, N.V. Pervukhina, and S.V. Borisov, Institute of Inorganic Chemistry, Siberian Branch of russian Academy of Sciences, Russia.

Regular interest in coordination compounds is accounted for by their wide usage in various chemical processes and technical fields. Crystal structure peculiarities have been considered for sufficiently large classes of inorganic compounds, containing such complex ions as  $(MA_{6\cdot n}X_n)$  or MX<sub>6-n</sub>A<sub>n</sub>), where n=O-6; M-transition metal, A-neutral (H<sub>2</sub>O or NH<sub>3</sub>), and X-acid (F<sup>\*</sup>, Cl<sup>\*</sup>, Br<sup>\*</sup>, I<sup>\*</sup> et al.) ligands (N.V. Podberezskaya, T.S. Yudanova, S.A. Magarill, et al., Probl. Kristallokhim. M.:Nauka, 1990, 82-98; Zh. Strukt. Khimii., 1991, 32, No. 6, 137-150). Complex cations or anions are considered to pack as quasi spheres of radii  $R_{ef}=d(M-X,A)+r_{vdv}(X,A)$ , where d(M-X,A) is the distance between the central atom of a complex ion and X or A atoms, and  $r_{vdv}$  is the van-der-Waal's radius of these atoms. Structures of inorganic coordination compounds may be divided into structure types as fluorite CaF<sub>2</sub>, anti-