Spectroscopic and Magnetic Properties of Compounds
Containing the Hydrazinium Cation as a Ligand

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The compounds of formula $M(N_2H_5)_2(SO_4)_2$, with $M = Mn, Fe, Co, Ni, Cu, Zn$ and Cd^{II} , have been investigated by means of infrared spectroscopy, ligand-field spectroscopy, magnetic susceptibility measurements, electron paramagnetic resonance spectra, and Mössbauer measurements.

The spectroscopic investigations indicate a distorted octahedral coordination for the metal ions. The results are in agreement with a previous crystal structure determination of $Zn(N_2H_5)_2(SO_4)_2$. Most of the spectra can be interpreted simply on the basis of tetragonal compressed octahedral geometry. The ligand $N_2H_5^+$ seems to behave as a normal nitrogen-donor ligand, as reflected by its spectroscopic parameters.

Evidence for rhombic distortions came from EPR spectra of the copper and manganese compounds.

Introduction

Coordination compounds containing positively charged ligands occur rather seldom and are subject of growing interest.¹⁻⁵ Examples of such positively charged ligands are $(CH_3)_3N^+-NH_2^4$, $(CH_3)_3N^+-(CH_2)_2-$

Some of these potential ligands are known to occur also as uncoordinated cations in coordination compounds^{4,6} of the type $L^+(MX_n^-)$ such as in $L_2^+(NiCl_4)^{2-}$

An older group of compounds is the series of metal hydrazinium sulfates of formula $M(N_2H_5)_2(SO_4)_2$, that is known, from structural studies, to contain metal ions coordinated distorted octahedrally by two $N_2H_5^+$ ions (*trans*) and two bidentate bridging sulfate anions.^{1,7}

Spectroscopic and magnetic studies on these com-

pounds have not been mentioned in the literature. The purpose of the present study is to fill up this omission and to investigate possible relationships between spectroscopic parameters and structural information for these unusually distorted octahedral compounds. In addition we were interested in the ligand-field spectral parameters of $N_2H_5^+$ compared with N_2H_4 .

Therefore ligand-field spectra, infrared spectra, EPR spectra Mössbauer spectra and magnetic susceptibility measurements were performed on a number of suitable compounds $M(N_2H_5)_2(SO_4)_2$.

Experimental Section

Starting materials. Bishydrazinium sulfate (Aldrich) was used without purification. Metal(II) sulfates were commercially available as the hydrates.

Preparation of the compounds. A solution of the metal sulfates (0.01 moles) in 25 ml of hot water was added to a hot solution of $N_2H_5SO_4$ (0.02 moles) in 15 ml of water. In most cases the compounds $M(SO_4)_2(N_2H_5)_2$ immediately separated as powders, except for $M = Mn^{II}$ where cooling to about 10° C was necessary to obtain the solid compound. The crystals were collected on a glass filter, washed subsequently with water, ethanol and diethylether, and finally dried *in vacuo* (0.1 mm/Hg at about 40°-50° C). The manganese and copper-doped Zn- and Cd compounds were prepared similarly, but with the addition of small amounts (about 1%) of $MnSO_4$ and $CuSO_4$ respectively.

Analyses. The compounds were checked for purity by metal(II) analyses and for a few cases by nitrogen and hydrogen analyses (carried out at organisch chemisch instituut, TNO, P.O. Box 5009, Utrecht).

Measurements. Infrared spectra were recorded on a Unicam SP-1200 spectrophotometer (4000-400 cm^{-1}) and on Hitachi EPI-L (700-200 cm^{-1}), as Nujol mulls between sodium chloride or polythene plates, and also as pressed KBr or CsI discs.

Ligand-field spectra of the solid compounds were obtained by the diffuse reflectance method on a Beckman DK-2A instrument as described elsewhere.⁸

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Table I. Coordination compounds containing the ligand $N_2H_5^+$.

Compound	Colour	X-ray type	metal analyses	
			calculated	found
$Mn(N_2H_5)_2(SO_4)_2$	white	A	17.5	18.3 ^a
$Fe(N_2H_5)_2(SO_4)_2$	pale green	A	18.0	18.2 ^b
$Co(N_2H_5)_2(SO_4)_2$	rose	A	18.6	18.0
$Ni(N_2H_5)_2(SO_4)_2$	green	A	18.5	17.4 ^c
$Cu(N_2H_5)_2(SO_4)_2$	blue	A'	19.7	19.6 ^d
$Zn(N_2H_5)_2(SO_4)_2$	white	A	20.2	20.1
$Cd(N_2H_5)_2(SO_4)_2$	white	A''	30.3	30.3

^a %N 17.28 (calc. 17.90); %H 3.27 (calc. 3.19). ^b %N 17.75 (calc. 17.85); %H 3.34 (calc. 3.19). ^c %N 17.55 (calc. 17.72); %H 3.37 (calc. 3.16). ^d %N (16.96 (calc. 17.42); %H 3.27 (calc. 3.11).

Table II. Infrared spectra of $M(N_2H_5)_2(SO_4)_2$, $(N_2H_5)_2SO_4$ and N_2H_4 .

Assignments	$Mn(N_2H_5)_2(SO_4)_2$ solid state	$(N_2H_5)_2SO_4$ solid	$N_2H_4^{10}$ liquid
weakly H-bridged N-H stretchings	3290 3230 3160 3070 3000	3240 3135 3030 2975	3332 3189
Strongly H-bridged N-H stretchings	2940 2760 2700 2660	 2690 2580	
N-H bendings	1630 1605 1575 1500	1625 1600 1575 1515sh 1495 1410	1608
N-H bending	1310		1324
N-H wagging	1175sh	1240 1185	1283
S-O asymm. stretch	1125 1075	1110 1085	1098
N-H rocking		970	1042
N-N stretching	1005	965	871
S-O symm. stretch	990	988	
Other bands	^a		

^a Bands below 700 cm^{-1} are listed in Table III.

Table III. Far-infrared spectra (700-200 cm^{-1}) of $M(N_2H_5)_2(SO_4)_2$.

Compound	$\nu_1(SO_4^{2-}) + N-H$ def.				N-N bending + $\nu_2(SO_4^{2-})$		Other bands M-N and M-O		
	ν_1	ν_2	ν_3	ν_4	ν_1	ν_2	ν_1	ν_2	ν_3
$Cd(N_2H_5)_2(SO_4)_2$	638	616	607	572	478	456m	360	280	220
$Mn(N_2H_5)_2(SO_4)_2$	638	621	604	558	479	460w	360	285	225
$Fe(N_2H_5)_2(SO_4)_2$	639	621	599	560	482	462vw	378	300	
$Zn(N_2H_5)_2(SO_4)_2$	647	622	606	589	488	464vw	395	300	235
$Ni(N_2H_5)_2(SO_4)_2$	646	620sh	609	593	485		404	305	235
$Cu(N_2H_5)_2(SO_4)_2$	655		614	600	492		420	315	240
$Co(N_2H_5)_2(SO_4)_2$	679	640	610		488		446	335	280sh 235

X-ray powder diagrams were performed on a Guinier camera, using $Cu-K\alpha$ radiation. Mössbauer spectra were obtained at temperatures between 80 and 300° K on commercial ELRON instruments, furnished with a ^{57}Co source diffused into 10 μm Pd foil. Sodium nitroprusside was used as a standard.

Magnetic susceptibilities were performed on a Faraday balance in the 80-300 K region, using $CoHg(SCN)_4$ as a calibrating standard tables. Diamagnetic corrections were taken from standard. EPR spectra of the powdered compounds were obtained at X- and

Q-band frequencies on Varian instruments as described elsewhere.⁹

Results

The compounds under investigation are listed in Table I, with analytical results, colours and X-ray

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Table IV. Ligand-field spectra and calculated parameters (kK) of compounds $M(N_2H_5)_2(SO_4)_2$.

Compounds	T(°K)	Bandmaxima					Dq	Parameters ^a		B
								—Ds	—Dt	
$Fe(N_2H_5)_2(SO_4)_2$	300	11.2	9.1				0.91	0.4 ^a	0.1 ^a	—
	100	11.8	9.3				0.93	0.5	0.1	—
$Co(N_2H_5)_2(SO_4)_2$	300	21.0sh	19.9	18.0sh	9.9	8.5	0.92	0.4 ^b	0.12 ^b	0.80
	100	21.2sh	20.1	18.0sh	10.1	8.7	0.94	0.5	0.11	
$Ni(N_2H_5)_2(SO_4)_2$	300	26.6	15.6	13.5sh	9.4		0.93	0.4 ^c	0.1 ^c	0.87
	100	26.9	15.9	13.6	9.7		0.95	0.5	0.1	0.87
$Cu(N_2H_5)_2(SO_4)_2$	300	14.0	10.0sh							
Cu^{2+} in $Zn(N_2H_5)_2(SO_4)_2$	300	15.3								

^a Ds and Dt obtained in combination with Mössbauer and Magnetic data; the errors are 10% for Ds and 20% for Dt; ^b Obtained from visible spectra only; the errors are 10% for Ds and 10% for Dt. ^c Obtained from the visible spectra; the values for Ds and Dt are upper limits obtained from band widths and a calculation with spin-orbit interaction (assuming $\lambda = -300$ cm^{-1}).

Table V. Mössbauer spectra and magnetic measurements of $Fe(N_2H_5)_2(SO_4)_2$.

Temperature (°K)	Mössbauer data		Magnetic data		After a $\theta = -8$ correction
	C.S. ^a	Q.S. ^a	χ (10^4 cgs) ^b	μ (B.M.) ^c	
340	1.42	2.99		5.24	
290	1.46	3.16	11690	5.20	5.28
254	1.48	3.29			
242			13910	5.22	5.28
220	1.50	3.41	15270	5.18	5.28
198	1.52	3.49	16850	5.16	5.27 ³
175			18820	5.15	5.27
146	1.55	3.60	22280	5.12	5.25
135			23900	5.10	5.23
117	1.57	3.68	27090	5.06	5.21
105	1.57	3.70	29920	5.02	5.19
78	1.59	3.76	37400	4.86	5.11

^a Mössbauer parameters are in mm/s, relative to sodium nitroprusside; C.S. values are accurate to ± 0.01 ; Q.S. values are accurate to ± 0.02 mm/s; Mössbauer temperatures are accurate to $\pm 3^\circ$. ^b Susceptibilities are corrected for diamagnetism by 300×10^{-6} cgs units. ^c Magnetic moments are accurate to ± 0.05 B.M.; relative errors are ± 0.01 B.M. since the measurements are from the same sample. ^d The value reported in ref. 1.

Table VI. EPR spectra of Cu^{2+} and Mn^{2+} in $M(N_2H_5)_2(SO_4)_2$.

Compound	Frequency	g-values ^a			Other parameters		
		g_3	g_2	g_1	A_3 ^c	A_2	A_1
$Cu(N_2H_5)_2(SO_4)_2$	X	2.330	2.095	2.055	u	u	u
	Q	2.325	2.090	2.050	u	u	u
Cu^{2+} in Zn compound	X	2.335	2.135	2.060	100	u	u
	Q	2.330	2.130	2.050	100	u	u
Cu^{2+} in Cd compound	X	2.345	2.105	2.055	u	u	u
	Q	2.340	2.095	2.045	u	u	u
		g_{iso} ^b			A_{iso} ^b	$D(cm^{-1})$ ^e	λ ^f
$Mn(N_2H_5)_2(SO_4)_2$	X,Q	2.01			95(5)	u	u
Mn^{2+} in Zn compound	X,Q	2.01			95(2)	0.025	0.25
Mn^{2+} in Cd compound	X,Q	2.01			94(3)	0.030	0.25

^a g-values are accurate to ± 0.005 ; ^b the g-value is ± 0.01 ; ^c The value for A_3 is ± 5 Gauss; ^d The uncertainty is indicated between parentheses; ^e The D-values are ± 0.005 cm^{-1} ; ^f λ is defined as E/D and is accurate to ± 0.05 ; ^g u is unresolved.

types. The division into the X-ray types was made according to the line patterns, that are very similar in d -values and relative intensities. Only the Cu^{II} and Cu^{II} compounds show slightly different line patterns. The infrared spectra of all compounds are very similar in the high-frequency region. For one compound the observed infrared bands are listed and compared with N_2H_4 and with $(N_2H_5)_2SO_4$ in Table II. Table III gives the far-IR spectra of the com-

pounds ($700-200$ cm^{-1}) together with some tentative assignments.

In Table IV the ligand-field maxima of the coloured compounds are presented along with some calculated parameters.

Magnetic susceptibility and Mössbauer results of $Fe(N_2H_5)_2(SO_4)_2$ are listed in Table V, whereas the EPR data of the Mn and Cu compounds are given in Table VI.

Discussion

Structure of the compounds. A two-dimensional X-ray analysis by Prout and Powell in 1961 has shown that $\text{Zn}(\text{N}_2\text{H}_5)_2(\text{SO}_4)_2$ contained Zn^{II} ions coordinated by four oxygens of four different SO_4 groups and two monodentate N_2H_5 groups, in an irregular octahedron.

The SO_4 groups are bidentate to two different Zn ions, yielding infinite chains in one direction. The distances around Zn are: Zn-N 2.08 Å, Zn-O 2.10 - 2.38 Å. Hydrogen bonds with N_2H_5 ligands hold the chains together.

The study of Hand and Prout⁷ on similar compounds and the results of the isomorphy study in Table I, indicate that all compounds of the present investigation have the same basic geometry. As will be shown below, the spectroscopic results are in agreement with this picture. According to the manner of description, distortion of the octahedron around the metal ion can be considered with respect to the nature of the ligands or with respect to the bondlengths.

Regarding the spectrochemical nature of the ligands, the octahedron is tetragonal compressed with a rhombic distortion due to the unequal O-M bond lengths. Regarding the bondlengths, on the other hand, the octahedron can be considered as tetragonal elongated, with a rhombic component due to the different spectrochemical positions of oxygen and nitrogen.

As will be shown below, most of the measurements can be understood on the basis of the compressed octahedron, without rhombic distortion. Only the EPR spectra of the Cu compounds can be best understood on the basis of an octahedron elongated along the Cu-O direction.

This slightly deviating behaviour of the Cu compound may be related to the deviating X-ray patterns of the Cu^{II} and Cd^{II} compounds that may be due to the nature of the metal ions (Cu is rather small and Cd is rather large in ionic radius). The results are not in agreement with a structure $\text{M}(\text{N}_2\text{H}_4)_2(\text{HSO}_4)_2$, containing bidentate N_2H_4 , as suggested by Bhargave *et al.*¹¹

Infrared Spectra. The reported absorption bands in Table II and III can all be ascribed to vibrations of either the N_2H_5^+ species, the SO_4^{2-} species, or the MO_4N_2 skelet.

Regarding the vibrations of the N_2H_5 groups it is observed that N-H stretches occur as low as 2660 cm^{-1} . This is indicative for strong hydrogen-bond formation with the SO_4^{2-} anions. The two bands at 3290 and 3230 cm^{-1} on the other hand are due to much weaker hydrogen-bonded N-H bands, presumably from the NH_2 side of the ligand that is further away from the anions.¹ There is some doubt about the position of the N-N stretching; possibly the highest one of the doublet near 1000 cm^{-1} is responsible for this vibration. This is quite high, although hydrazine is also known to have increased N-N stretches upon coordination. In addition the Irving-Williams sequence of metal ions holds.

Although the S-O stretching vibrations principally should yield information about the bonding of the SO_4^{2-} anions, the manner of coordination cannot be derived from the present spectra, because of the broad bands and the overlap with the N-H vibrations. On the other hand, the spectra are not in contradiction with the structure determination of the Zn compound.

N-H deformation vibrations and the N-N bending are observed below 700 cm^{-1} . The former occur just around the split $\nu_4(\text{SO}_4)$ vibrations near 600 cm^{-1} , yielding a rather complex line pattern of at least four lines. The N-N bending is observed near 480 cm^{-1} close to the $\nu_2(\text{SO}_4)$ vibration (forbidden in Td symmetry). It is observed from Table III that the Irving-Williams sequence of metal ions, *i.e.* $\text{Mn} < \text{Fe} < \text{Co} < \text{Ni} < \text{Cu} > \text{Zn} > \text{Cd}$, also holds for this vibration. This is not unexpected, and has been observed before for many other ligand vibrations.¹³

Below 450 cm^{-1} we observe the vibrations due to the MO_4N_2 skelet, *i.e.* the M-O and M-N stretchings. Due to the complexity of the species, it is impossible to speak of separate M-N and M-O frequencies, and in fact the band at highest frequency falls close to the region commonly observed for metal-oxygen vibrations^{12,13} and is also in the region of M-N found for hydrazine complexes.¹⁴ Again the Irving-Williams sequence holds.

Ligand-field Spectra. The observed reflectance spectra of the solid compounds are in agreement with a basically octahedral geometry around the metal ions. However, most bands are very broad and some of them even show splittings. A few splittings become better resolved at low temperatures.

Assuming compressed tetragonal geometry¹ and using the procedures for the evaluation of parameters given before,⁸ we could calculate values for Dq, Ds, Dt and B for the several compounds. As the octahedron is compressed along the tetragonal axis, the values for Ds and Dt are negative.

For the iron compound, only the value for $4\text{Ds} + 5\text{Dt}$ could be determined from the ligand-field spectra (*i.e.* equal to the separation between the two ligand-field bands). Combination with the results obtained from the Mössbauer spectra and the magnetic susceptibility measurements, however, yielded separate values for Dt and Ds, as indicated in Table IV.

The parameters for the cobalt compound could be calculated accurately because of the fact that more (split) bands are observed.

The spectra of the nickel compound showed no splittings at all, even not at liquid nitrogen temperatures. After taking a value of -300 cm^{-1} for the spin-orbit coupling constant, from the observed line width an upper limit for Ds and Dt could be determined with the aid of a full matrix diagonalization procedure.

As is seen from Table IV the parameters for the

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Fe, Co and Ni compounds agree very well with each other. For $\text{Cu}(\text{N}_2\text{H}_5)_2(\text{SO}_4)_2$ and for the copper doped in the zinc compound, no parameters could be calculated due to the lacking of sufficient bands.

As can be seen from Table IV, both Dq and $-Ds$ increase upon cooling to liquid nitrogen temperature. The increase in Dq is normal, and has been observed before.⁸ The increasing $-Ds$ value corresponds to a greater T_{2g} splitting, and has also been found in the investigation of the Mössbauer spectra and magnetic susceptibilities (*vide infra*).

The slight increase in Dq^{xy} on going from Fe to Ni, is just as expected from the decreasing ionic radius, and has been observed previously for pyrazole ligands.⁸

From the Dt values, the Dq^z value for N_2H_5^+ can be calculated by the relation: $Dq^z - 7Dt/4 = Dq^x$.

Using $Dq^{xy} = 0.93$ kK and $Dt = -0.1$ kK, this yields: $Dq^z = 1.1$ kK, a value quite normal for nitrogen-donor ligands,⁸ but slightly smaller than 1.15 kK as found for neutral N_2H_4 .¹⁵ This smaller value might be due to the positive charge upon the ligand.

In none of the complexes indications were found for the occurrence of rhombic splittings in the ligand-field bands. As will be shown below, such splittings are rather small for the iron compound, although EPR spectra of the copper compound clearly show the presence of a rhombic component in the ligand-field (*vide infra*).

Magnetic and Mössbauer measurements on $\text{Fe}(\text{N}_2\text{H}_5)_2(\text{SO}_4)_2$. For a more detailed study on the ground-state splitting in the iron(II) compound, we have investigated the Mössbauer spectra and the paramagnetic susceptibility between 80 and 300 °K.

Since the ligand-field spectra showed an increase in $-Ds$ for the present compounds upon cooling, the measurements were carried out in the whole temperature region (Table V), to see if the changes are gradually or not.

As is seen from Table V, all values change rather gradually upon cooling to liquid nitrogen temperature. The increase in the Mössbauer centre shift (C.S.) is just as expected from the second-order Doppler shift.¹⁶ The magnitude of the room-temperature C.S. is somewhere between oxygen- and nitrogen-donor ligands, in agreement with the proposed structure. (Cf. N-donor ligands with C.S. values from 1.3 - 1.4 mm/s, and O-donor ligands with values from 1.5-1.6 mm/s).¹⁷⁻¹⁹

Principally, the analysis of the quadrupole splittings and the magnetic moments as a function of temperature should yield the parameters δ (the ground-state splittings, equal to $5Dt-3Ds$), ϵ (the rhombic splitting), λ (the spin-orbit coupling constant), and k (the orbital reduction factor).

However, as the difference between Q.S. (300 K) and Q.S. (80K) of about 0.7 mm/s is higher than the maximum possible increase of about 0.5 mm/s,²⁰

such a large increase can be explained only by an increasing ground-state splitting upon cooling, i.e. an increasing δ and/or ϵ .

The same appears to be the case for the magnetic moments. For compressed octahedrons we expect a nearly temperature independent behaviour for μ between 300 and 80 K.^{21,22} The observed decrease in our μ values can be due to two mechanisms:

a. An increasing value of the ground-state splitting upon cooling, which would be in agreement with the Mössbauer spectra.

b. Antiferromagnetic exchange in the SO_4 -bridged chains, that has the effect of decreasing μ_{eff} upon cooling. A correction is usually made for lattice compounds by setting χ equal to $C/(T - \theta)$. For comparison in Table V the calculated μ -values with a θ -value of -8 have also been listed.

To obtain detailed information about the exact magnitude of B , measurements as low as liquid Helium are necessary. These are in progress for all these compounds. For the moment we will restrict ourselves to a magnetically diluted system, ignoring any antiferromagnetic exchange.

Therefore, we made use of a full matrix consideration of Fe^{2+} (5D) under the simultaneous action of octahedral tetragonal and rhombic ligand fields and spin-orbit coupling. In this way we obtained μ and Q.S. as a function of T , Dq , Ds , Dt , Dr , K and λ . In the calculation procedures influences of covalency upon the field gradient were ignored, and the lattice contribution was taken to be equal to 0.85 Ds (in kK), expressed in mm/s.³

The calculations were carried out on a IBM 360/65 computer, using programs developed by Mr. A. Vermaas. The main program consists of two parts:

a. A part calculating Q.S. splittings; this is a modified version of the program written by de Vries.²⁰

b. A part calculating the susceptibilities; this program was set up as published by Figgis et al.²¹ and König et al.²², but extended to rhombic splittings and interaction with the higher E-term. Details of the program will be published elsewhere.²⁴

In the analyses of the data, we fixed the values of Dq and of $(4D + 5Dt)$ that were obtained from the ligand-field spectra. Further we used a value of 0.25 barn for the quadrupole moment of ^{57}Fe .^{19,23} The best fit with both measurements was obtained for the following parameters (Table VII):

Table VII. Spectral parameters of $\text{Fe}(\text{N}_2\text{H}_5)_2(\text{SO}_4)_2$.

Temperature	$Dq^{xy,a}$	Ds^a	Dt^a	$Dr^{a,b}$	$-\lambda^c$	k	$\delta^{a,c}$
300K	0.91	-0.4	-0.1	0.3	75	0.7	700
200K	0.92	-0.45	-0.1	0.3	75	0.7	850
100K	0.93	-0.5	-0.1	0.3	75	0.7	1050

^a In kK; ^b upper limit; ^c in cm^{-1}

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The assumption of a θ value of -8° , influences the parameters slightly (*i.e.* λ is increased to about 80 cm^{-1} , k is increased to about 0.8 and δ is decreased by about 10%) and even give a better fit. Summarizing, these results indicate:

a. A compressed tetragonal structure, with a ground-state orbital singlet, having a doublet at about 0.7 - 1.0 kK above it.

b. The rhombic splitting of this doublet - if present - is smaller than 0.3 kK.

c. The values of k and λ indicate some covalency in the Fe-ligand bonds, in agreement with the rather low B-values for the Co and Ni compounds (Table IV).

EPR spectra of some powdered compounds containing Cu^{2+} or Mn^{2+} . To obtain detailed information about the geometry and the kind of bonding in the present compounds. EPR spectra of the Mn^{2+} and Cu^{2+} compounds and impurities of them in the diamagnetic lattices of the Zn and Cd were investigated.

From the results in Table VI it is seen that all copper compounds show a three g-value spectrum, characteristic for rhombic geometry.²⁵ Whether this three-g-value spectrum is derived from axial elongated (*i.e.* $g_3 = 2.0$ and $g_2 = g_1 \approx 2.4$) cannot be directly concluded from the spectra, although the fact that the lowest g-value is about 2.05 points in to the description of elongated octahedral geometry. Anyway, the rhombic distortion is clearly demonstrated by the occurrence of three separate g-values. The calculated g-values from the spectra were obtained from computer produced simulations,²⁶ and for all compounds the results are similar to the observed X and Q band spectra within experimental error.

Copper hyperfine splittings were not observed, except for the dope in the Zn compound (upon g_3). In fact the hyperfine coupling constants may be small due to the nature of the metal-ligand bond (usually Cu^{2+} compounds with oxygen-donor ligands do not show resolved hyperfine splittings),^{25,27} or due to

exchange coupling between the Cu ions. The latter case almost certainly occurs for the pure copper compound.

Exchange coupling must also be responsible for the observed signal of the pure Mn^{2+} compound. The symmetrical signal, with six inflections assigned to hyperfine interactions, at $g = 2.01$ is just as expected for exchange coupled Mn ions. The dopes in the Zn and Cd compounds however, show hyperfine splittings (due to the ^{55}Mn nucleus) as well as fine splittings (due to the zero-field splitting of the $^6\text{A}_1$ ground state). From these the parameters A and D and λ were calculated using the methods of Dowsing and Gibson.²⁸ The slightly larger D value for Mn doped in the Cd compound indicates a somewhat larger distortion. The different D value is in agreement with the observed non-isomorphism of the lattices (Table I).

Conclusions

From the results of the present investigation the following conclusions can be drawn:

1. The complexes $\text{M}(\text{N}_2\text{H}_5)_4(\text{SO}_4)_2$ contain bidentate bridging sulfate anions and monopositive monodentate N_2H_5 ligands.

2. Most spectral and magnetic measurements are in agreement with a coordination polyhedron build up by two nitrogens (*trans*) and described by a tetragonal compression with an additional rhombic splitting. The EPR spectra indicate rhombic symmetry derived from elongated octahedral geometry, on the other hand.

3. The presence of this rhombic distortion could not be proved from infrared, ligand-field and Mössbauer spectra and from magnetic susceptibilities.

4. The ligand N_2H_5^+ behaves as a normal nitrogen donor ligand, with a crystal-field splitting somewhat smaller than neutral hydrazine, which might be explained by the positive charge upon the ligand.

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