

Transition	$\alpha = \mu_a(\text{HDSe}) / \mu_a(\text{HDO})$
$2_{20} - 2_{21}$	$0,68 \pm 0,02$
$4_{31} - 4_{32}$	$0,67 \pm 0,02$
$7_{43} - 7_{44}$	$0,70 \pm 0,02$
$9_{54} - 9_{55}$	$0,67 \pm 0,02$
	$\alpha_{av} = 0,68 \pm 0,02$

for these measurements was 2kv/cm. The resulting value of  $\alpha$  is the ratio of the dipole components along the  $a$  axis for the molecules HDSe and HDO; values are given in the table. The total dipole moment of HDSe is associated with  $\alpha$  by the simple expression

$$\mu(\text{HDSe}) = \alpha \mu(\text{HDO}) \sin \delta_2 / \sin \delta_1,$$

where  $\mu(\text{HDO})$  is the dipole moment of HDO and  $\delta_1$  and  $\delta_2$  are the angles between the  $a$  axis and the direction of the dipole moment for HDSe and HDO respectively.  $\delta_1$  is approximately  $45^\circ$ , and if we take  $\delta_2 = 20^\circ 30'$  and  $\mu(\text{HDO}) = 1.84$  Debye units<sup>4</sup> we obtain 0.62 Debye units for the dipole moment of HDSe, which differs extremely from the value of 0.24 Debye units obtained in Ref 1.

<sup>1</sup>Jache, Moser and Gordy, J. Chem Phys. **25**, 209 (1956).

<sup>2</sup>V. G. Veselago and A. M. Prokhorov, J. Exptl. Theoret. Phys. (U.S.S.R.) **31**, 731 (1956); Soviet Phys. JETP **4**, 750 (1957).

<sup>3</sup>Weisbaum, Beers and Herrmann, J. Chem. Phys. **23**, 1601 (1955).

<sup>4</sup>M. W. P. Strandberg, J. Chem. Phys. **17**, 901 (1949).

Translated by I. Emin

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## Second Relaxation in a Spin System at Room Temperature

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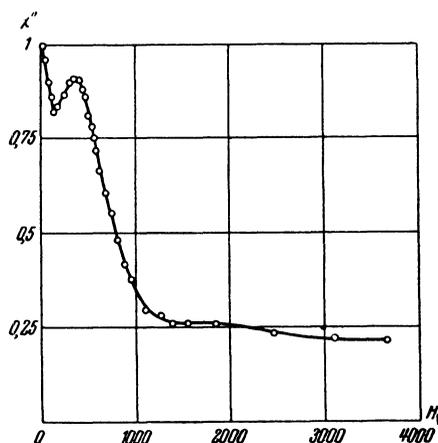
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THE investigation of paramagnetic absorption in  $\text{Cr}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$  in parallel fields at  $300^\circ\text{K}$ ,

employing Zavoiskii's grid current method<sup>1</sup>, revealed an effect which is unusual at room temperature. The absorption  $\chi''(H_{\parallel})$  plotted in the figure for a 160 Mc oscillating field has quite a narrow peak. The right-hand half width was of the order of 300 oersteds. As  $\nu$  increases the absorption peak is shifted in the direction of higher constant magnetic field strengths; this is shown in the following set of data:

$\nu$ in Mc =	48	93	131	160
Magnetic field in oersteds at peak =	250	290	330	360



The intensity of the absorption peak compared with absorption at  $H=0$  diminishes with increasing frequency from 10 to 160 Mc. At frequencies of the order of 660 Mc, the shape of the paramagnetic absorption curve differs very little from the usual  $\chi''(H_{\parallel})$  curve which is described by Shaposhnikov's formula<sup>2</sup>  $\chi'' = (1-F)^2 \rho_s \nu$ .

Below 10 Mc, spin-lattice relaxation influences the absorption curve so strongly that a peak is hardly discernible.

This phenomenon is apparently associated with the new form of spin-spin relaxation discovered by DeVrijer and Gorter<sup>3,4</sup> in potassium chrome alum at the temperature of liquid hydrogen. Gorter and his associates<sup>5</sup> later detected this type of relaxation in a number of other materials but again only at very low temperatures. So far as we know the effect has not previously been observed at room temperature.

At the present time the author is using the grid current method in similar investigations of other chromium salts and salts of  $\text{Mn}^{++}$ ,  $\text{Cu}^{++}$  and  $\text{Fe}^{+++}$ .

In conclusion I take this opportunity to thank B. M. Kozyrev for guidance and constant assistance.

