reaction (2) is approximately three times greater than that in reaction (1).

\*We make use of this opportunity to express our thanks to B. S. Neganov for help rendered to us in conducting this experiment.

\*\* The geometry of our  $\gamma$ -detector allows detection of all electrons of energy > 10 mev emitted in a cone of opening angle 40°.

\*\*\* On the assumption of an isotropic angular distribution of  $\pi^{0}$ -mesons Eq. (3) tells us that the cross section for formation of  $\pi^{0}$ -mesons from carbon is  $\sigma_{c}^{\pi^{0}}$ = 27.4 ± 6.7 × 10<sup>-27</sup> cm<sup>2</sup>.

<sup>1</sup> V. B. Fliagin, Reports Inst. Nuclear Problems Acad. Sci. USSR 1954

<sup>2</sup> Iu. M. Kazarinov, Reports Inst. Nuclear Problems, Acad. Sci. USSR 1954

<sup>3</sup> G. Moliere, Naturforsch. 3a, 78 (1948).

<sup>4</sup> B. D. Balishov, V. A. Zhukov, B. M. Pontecorvo and G. I. Selivanov. Reports, Inst. Nuclear Problems, Acad. Sci., USSR, 1954

<sup>5</sup> R. W. Hales and B. J. Moyer, Phys. Rev. 89, 1047 (1953).

<sup>6</sup> R. A. Shluter, Phys. Rev. 96, 734 (1954)

<sup>7</sup> A. A. Tiapkin, M. S. Kozodaev and Ia. D. Prokoshkin Dokl. Akad. Nauk. SSSR 100, 689 (1955).

<sup>8</sup> V. P. Dzhelepov, Iu. M. Kazarinov, B. M. Golovin, V. B. Fliagin and V. I. Satarov, Izv. Akad. Nauk; SSSR Ser. Fiz. 19, 5 (1955).

<sup>9</sup> M. G. Meshcheriakov and B. D. Neganov, Dokl. Akad Nauk. SSSR 100, 677 (1955)

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Microwave Spectrum of the C<sub>2</sub>H<sub>5</sub>Cl Molecule

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W E have carried out a preliminary study of the rotational spectrum of the  $C_2H_5Cl$  molecule. A short communication on the spectrum of this molecule was published by Wagner and Dailey<sup>1</sup>, who studied the transitions  $1_{11} - 2_{12}$ ,  $1_{10} - 2_{11}$ ,  $2_{22} - 3_{32}$  and  $2_{20} - 3_{21}$  in the  $C_2H_5Cl^{35}$  molecule, and the  $1_{11} - 2_{21}$  and  $1_{10} - 2_{11}$  transitions in the  $C_2H_5Cl^{37}$  molecule. From these transitions they

obtained the values of the rotational constants B and C, as well as the magnitudes of the quadrupole bonds along the principal axes of inertia.

We have studied additional transitions whose frequencies have the following values, neglecting the effects of quadrupole interactions:

| Molecule                                       | Transition                        | Observed Frequency (mc/sec) |
|--|-----------------------------------|-----------------------------|
| C <sub>2</sub> H <sub>5</sub> Cl <sup>35</sup> | $\overline{0}_{00} - 1_{01}$      | 10 246,20 <u>+</u> 0,05     |
| C <sub>2</sub> H <sub>5</sub> Cl <sup>35</sup> | $1_{01} - 2_{02}$                 | 20 903.80 $\pm$ 0.04        |
| C <sub>2</sub> H <sub>5</sub> Cl <sup>37</sup> | 0 <sub>00</sub> - 1 <sub>01</sub> | 10 456,00 $\pm$ 0,05        |

The frequency of the transition line  $0_{00} - 1_{01}$  coincides with the frequency calculated from the the values for *B* and *C* given in reference 1\*.

The frequency of the transition  $1_{01} - 2_{02}$ depends on the rotational constant A; calculation gave the value  $A = 30,940 \pm 200$  mc/sec. The low accuracy in the determination of A is due to the fact that A enters into the value of the transition frequency  $1_{01} - 2_{02}$  as a small correction of the form  $(B - C)^2/A$ , and the magnitudes of B and Care very similar. Note that, in general, the accuracy of determining A is always small if it is determined from transitions involving a change in the dipole moment  $\mu_a$ . We propose to observe transitions involving the measurement of  $\mu_b$ . Although these transitions give less intense lines, they provide a more accurate determination of A.

Using the transition  $0_{00} - 1_{01}$ , we have determined the dipole moment  $\mu_a$  of the  $C_2 H_5 Cl^{35}$  molecule from the Stark splitting of the line which has the hyperfine structure F = 3/2 - 5/2, at applied fields of 195 v/cm and 292 v/cm. The calculation of  $\mu_a$  was carried out using the "weak field" formula. The value found for  $\mu_a$  was  $1.79 \pm 0.05 D$ .

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<sup>\*</sup> The rotational constants for  $C_2H_5Cl^{37}$  given in reference 1 are incorrect, and were therefore calculated directly from the line frequencies given therein.

<sup>&</sup>lt;sup>1</sup> R. S. Wagner and B. P. Dailey, J. Chem. Phys. 22, 1459 (1954).