

Spin-Rotation Interaction and Anisotropic Chemical Shift in $^{13}\text{CS}_2$ *

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The ^{13}C nuclear spin-lattice relaxation time T_1 was studied in liquid CS_2 from -106°C to $+35^\circ\text{C}$ at resonance frequencies of 14, 30, and 62 MHz. The relaxation is caused by anisotropic chemical shift and spin-rotation interaction. It is shown that for linear molecules the spin-rotation constant C and the anisotropy of the chemical shift $\Delta\sigma$ can be obtained from the relaxation rates without use of adjustable parameters. The analysis yields: $C = -13.8 \pm 1.4$ kHz and $\Delta\sigma = 438 \pm 44$ ppm.

INTRODUCTION

Several mechanisms are known to contribute to the spin-lattice relaxation time T_1 of a nucleus with spin $I = 1/2$ in pure diamagnetic liquids, e.g., dipolar coupling, anisotropic chemical shift, and spin-rotation interaction. In most of the systems studied so far, the relaxation rates are dominated by the dipolar coupling. For the ^{13}C nucleus in CS_2 , on the other hand, the latter two mechanisms are expected to determine T_1 . The possibility that anisotropic chemical shift might be a significant relaxation mechanism in CS_2 was first pointed out by McConnell and Holm (1). As already mentioned by Abragam (2), "the obvious check of changing the resonance frequency ω and investigating whether $1/T_1 \propto \omega^2$ does not seem to have been made." By use of a superconducting magnet, it is now possible to study T_1 at higher frequencies, and data are presented below taken at 14, 30, and 62 MHz. Spin-rotation interaction as a relaxation mechanism for CS_2 has been mentioned recently by Lippmaa *et al.* (3). In a theoretical study of the contributions to T_1 of spin-rotation interaction and anisotropic chemical shift, Deutch and Waugh (4) concluded that both mechanisms would contribute appreciably to T_1 for ^{13}C in simple molecules like CO, CO_2 , and CS_2 .

The spin-rotation interaction is closely related to the paramagnetic part of the chemical shift (5, 6). For linear molecules, this relation may be extended to yield the anisotropy of the chemical shift $\Delta\sigma$ as a function of the spin-rotation constant C to a good approximation. By multiplication of the individual relaxation rates, due to spin-rotation interaction and anisotropic chemical shift, the correlation times for both interactions can be eliminated (4) and a second equation relating $\Delta\sigma$ and C

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is obtained. Thus both quantities can be determined from relaxation studies without use of adjustable parameters. In order to get the necessary information, T_1 has to be studied as a function of ω and the temperature T .

CHEMICAL SHIFT AND SPIN-ROTATION INTERACTION IN LINEAR MOLECULES

The chemical shift tensor $\hat{\sigma}$ is usually represented as the sum of a diamagnetic part $\hat{\sigma}^d$ and a paramagnetic part $\hat{\sigma}^p$:

$$\hat{\sigma} = \hat{\sigma}^d + \hat{\sigma}^p. \quad [1]$$

Taking the C_∞ axis of the diamagnetic linear molecule as the z axis we note that $\sigma_{zz}^p = \sigma_{\parallel}^p$ vanishes, because the angular momentum operator L_z commutes with the total electronic Hamiltonian (2, p. 182). For $\sigma_{xx}^p = \sigma_{yy}^p = \sigma_{\perp}^p$, we have (5, 6)

$$\sigma_{\perp}^p = \frac{e^2}{2mc^2} \left(\frac{\Theta c}{e\mu_0 g_K} C_K - \sum'_{\alpha} \frac{N_{\alpha}}{r_{\alpha}} \right), \quad [2]$$

where $C_K = C_{K\perp}$ is the spin-rotation constant¹ of the K -nucleus. Note that, for a linear molecule, $C_{K\parallel}$ vanishes. Θ is the moment of inertia of the molecule, μ_0 , g_K are the nuclear magneton and the nuclear g -factor of the K -nucleus, respectively, N_{α} is the atomic number of the α -nucleus, and r_{α} the internuclear distance between α - and K -nuclei. The prime on the summation indicates $\alpha \neq K$. For the diamagnetic shift, we use the approximate formula, derived by Flygare and Goodisman (7),

$$\sigma_{av}^d = \sigma_{av}^d(\text{free atom}) + \frac{e^2}{3mc^2} \sum'_{\alpha} \frac{N_{\alpha}}{r_{\alpha}} \quad [3]$$

valid for the isotropic part. These authors also show (7) that similar equations hold for the individual components of $\hat{\sigma}^d$ as well to a good approximation.

$$\sigma_{ii}^d = \sigma_{av}^d(\text{free atom}) + \frac{e^2}{2mc^2} \sum'_{\alpha} \frac{N_{\alpha}}{r_{\alpha}^3} (j_{\alpha}^2 + k_{\alpha}^2); \quad i, j, k = x, y, z; \text{ cyclic.} \quad [3a]$$

From [3a] we get for a linear molecule:

$$\sigma_{zz}^d = \sigma_{\parallel}^d = \sigma_{av}^d(\text{free atom}), \quad [4]$$

$$\sigma_{xx}^d = \sigma_{yy}^d = \sigma_{\perp}^d = \sigma_{av}^d(\text{free atom}) + \frac{e^2}{2mc^2} \sum'_{\alpha} \frac{N_{\alpha}}{r_{\alpha}}. \quad [5]$$

Combining Eqs. [2, 4, and 5] we get for the anisotropic chemical shift of a linear molecule

$$\Delta\sigma = \sigma_{\parallel} - \sigma_{\perp} = - \frac{|e|}{2mc} \frac{\Theta}{\mu_0 g_K} C_K. \quad [6]$$

Equation [6] shows that for linear molecules the anisotropy of the chemical shift is approximately proportional to the product of the moment of inertia and the spin-rotation constant.

An expression for the relaxation rate of a nuclear spin due to spin-rotation interaction has been derived by Hubbard (8). For a *linear* molecule, we get (see also Green

¹ C_K is a component of the spin-rotation tensor defined by $\mathcal{H}_{SR} = \hbar \vec{C}_K \cdot \vec{J}$. Thus spin-rotation constants obtained from molecular beam data have to be multiplied by 2π to be used in our notation.

and Powles (9))

$$\frac{1}{T_1^{\text{SR}}} = \frac{4}{3} \frac{\Theta}{\hbar^2} C_K^2 \tau_{\text{SR}}, \quad [7]$$

where τ_{SR} is the correlation time for angular velocity. Huntress (10) has derived equations for the relaxation rate due to anisotropic chemical shift for a liquid in the diffusion limit. For a linear molecule we have

$$\frac{1}{T_1^{\text{CS}}} = \frac{1}{45} \frac{\omega^2 \Theta}{kT} (\Delta\sigma)^2 \frac{1}{\tau_{\text{SR}}}. \quad [8]$$

In the diffusion limit, the correlation times for angular velocity τ_{SR} and angular reorientation τ_c are related by (10, 8)

$$(\tau_c)_i (\tau_{\text{SR}})_i = \frac{\Theta_i}{6kT}, \quad [9]$$

where the index i refers to the i -th axis. Therefore, Eq. [8] is identical with the expression usually given for $1/T_1^{\text{CS}}$ using τ_c (1, 2)

$$\frac{1}{T_1^{\text{CS}}} = \frac{2}{15} \omega^2 (\Delta\sigma)^2 \tau_c. \quad [8a]$$

It is apparent from Eqs. [7, 8] that τ_{SR} can be eliminated by multiplication of the two relaxation rates (see also Ref. (4)):

$$\frac{1}{T_1^{\text{SR}}} \cdot \frac{1}{T_1^{\text{CS}}} = \frac{4}{135} \frac{\omega^2 \Theta^2}{\hbar^2} C_K^2 (\Delta\sigma)^2. \quad [10]$$

If $1/T_1^{\text{SR}}$ and $1/T_1^{\text{CS}}$ can be determined experimentally, Eqs. [6, 10] can be used to obtain the absolute values of both C_K and $\Delta\sigma$ from relaxation studies. It should be noted that the only molecular parameter besides C_K and $\Delta\sigma$ appearing in Eqs. [6, 10] is the moment of inertia which is well defined for nonassociated liquids.

The close relationship of spin-rotation interaction and anisotropic chemical shift can also be used to calculate the spin-rotation constant for a given linear molecule from a known spin-rotation constant for the same nucleus in a different linear molecule if the relative chemical shift is known. Thus combining Eqs. [2] and [3] and noting that $\sigma_{\parallel}^r = 0$ (see above), we get (7)

$$\sigma_{av} = \sigma_{av}^d (\text{free atom}) + \frac{|e|}{3mc} \cdot \frac{\Theta}{\mu_0 g_K} C_K. \quad [11]$$

Equation [11] is used below to calculate the ^{13}C spin-rotation constants for CS_2 and CO_2 based on the known value for ^{13}CO .

EXPERIMENTAL

The CS_2 sample, isotopically enriched (60%) and oxygen-free, was obtained from Prochem Ltd., Carolyn House, Croydon, England. A Bruker pulse spectrometer was used together with a Siemens superconducting magnet (max. field 80 kG). The NMR probes, operating at 14, 30, and 62 MHz were advanced versions of the single coil probe described earlier (11). All data were directly taken from the oscilloscope.

Details of the method used to measure T_1 have also been described in Ref. (11). The sample temperature was controlled by a regulated nitrogen stream and determined by a thermo-couple near the sample; it was stable within 1°C during the time necessary to determine T_1 (typically ca. 30 min).

RESULTS AND DISCUSSION

In Fig. 1 the experimental spin-lattice relaxation rates $1/T_1$ obtained at 14, 30, and 62 MHz are plotted vs. the temperature. The data were taken over essentially all of the liquid range at atmospheric pressure. So the temperatures are well below the critical temperature. It is apparent from Fig. 1 that relaxation through anisotropic

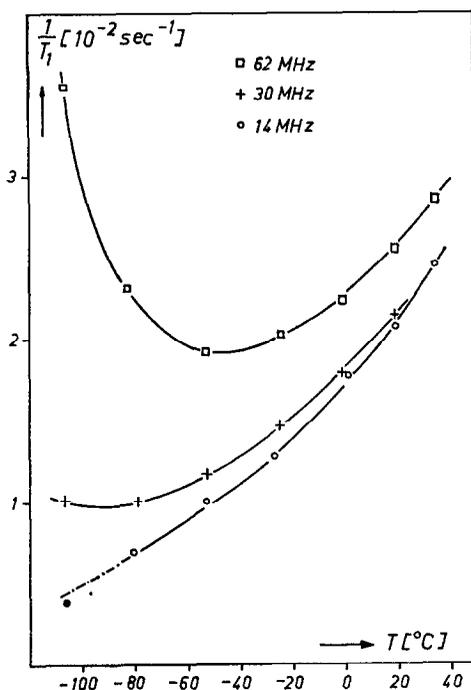


Fig. 1. Experimental relaxation rates in $^{13}\text{CS}_2$ at 14.0369, 30.1026, and 61.9164 MHz.

chemical shift is dominant for 62 MHz at low temperatures. In order to separate the relaxation rate due to anisotropic chemical shift from the magnetic field independent part, in Fig. 2 the relaxation rates for the different temperatures vs. the square of the resonance frequency are plotted. Straight lines are obtained and therefore $1/T_1^{\text{CS}}$ can be determined accurately for 62 MHz.

The field-independent part of the relaxation rate decreases with decreasing temperature, suggesting that it is dominated by the spin-rotation interaction. The dipolar contribution to $1/T_1$ can be estimated (2) to give for the intramolecular interaction $(1/T_1^{\text{d}})_{\text{intra}} < 3 \cdot 10^{-6} \text{ sec}^{-1}$ and for the intermolecular interaction $(1/T_1^{\text{d}})_{\text{inter}} < 2 \cdot 10^{-5} \text{ sec}^{-1}$, which can be neglected compared with the measured relaxation rates

(see Fig. 1). This was also confirmed experimentally by studying T_1 of a CS_2 sample with ^{13}C in natural abundance at 62 MHz. No deviations from the isotopically enriched sample were found.

Thus we are left with only two relaxation mechanisms, and the analysis outlined above can be employed. Figure 3 shows the relaxation rates due to anisotropic chemical shift at 62 MHz and due to spin-rotation interaction, together with the product of the two rates, which is independent of the temperature within experimental

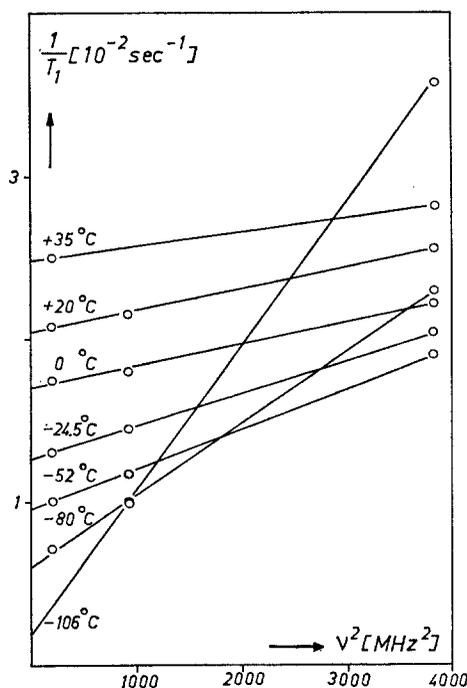


FIG. 2. Frequency dependence of experimental relaxation rates in $^{13}\text{CS}_2$.

error. This result confirms Eq. [10] experimentally, particularly since the individual relaxation rates span about the same range in the temperature interval studied.

In the derivation of Eq. [10] the relation between the correlation times for angular velocity and angular reorientation, Eq. [9] has been used. This relationship has recently been shown (12) to be valid for the quasi-spherical molecule ClO_3F , where both τ_c and τ_{SR} were determined experimentally, at temperatures significantly below the critical temperature. On the other hand, Deverell (13) has noted deviations from Eq. [9]. The values for τ_c he used, however, were calculated from macroscopic diffusion constants using estimated molecular radii. The possibility that the error in τ_c can account for at least part of the discrepancy is mentioned already in Ref. (13). This demonstrates that calculating τ_c using the Stokes-Einstein equation is not sufficient in order to get meaningful values for τ_{SR} using Eq. [9] and thus reliable values for the spin-rotation constants from relaxation rates.

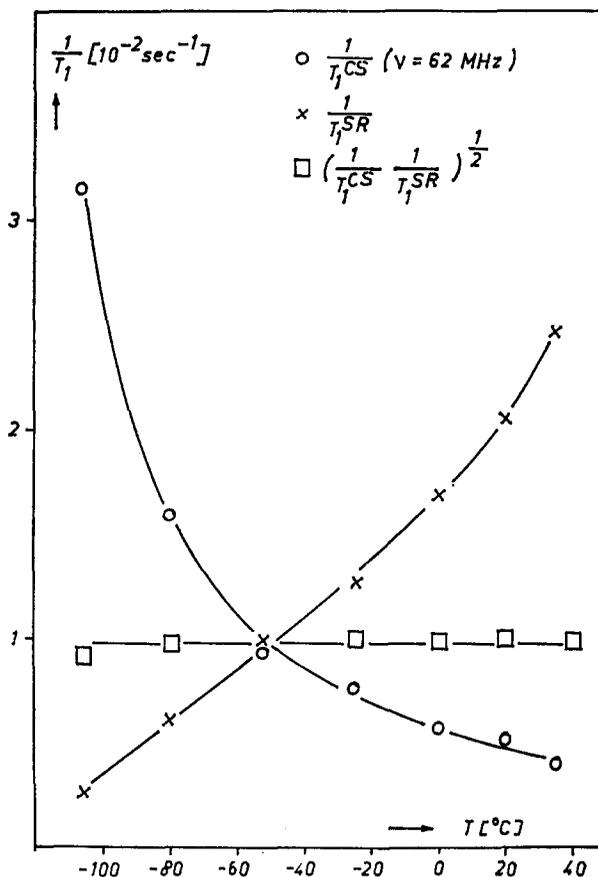


FIG. 3. Relaxation rates due to anisotropic chemical shift at 62 MHz and due to spin-rotation interaction in $^{13}\text{CS}_2$.

Equation [9] is valid only if $\tau_{\text{SR}} \ll \tau_c$ (8, 10). An order of magnitude estimate may be obtained by calculating the correlation times using the macroscopic viscosity (8):

$$\tau_{\text{SR}} = \frac{1}{3} \Theta a^{-3} \pi \eta \quad \text{and} \quad \tau_c = \frac{4\pi}{3} a^3 \eta (kT)^{-1}.$$

As the ratio τ_{SR}/τ_c decreases with decreasing temperature we need only consider the highest temperature (35°C). Using $a = 1.5 \text{ \AA}$, the internuclear distance C-S, we get $\tau_{\text{SR}} \approx 3 \cdot 10^{-14} \text{ sec}$ and $\tau_c \approx 1.3 \cdot 10^{-12} \text{ sec}$ and thus $\tau_{\text{SR}} \ll \tau_c$ is fulfilled. From the experimental data, we then get for ^{13}C in CS_2 , using Eqs. [6, 10]

$$|C| = 13.8 \pm 1.4 \text{ kHz},$$

$$|\Delta\sigma| = 438 \pm 44 \text{ ppm}.$$

As can be seen from Fig. 3, the error in the measured quantity $(1/T_1^{\text{SR}} \cdot 1/T_1^{\text{CS}})^{1/2}$ is only 5%. The error limits are given under the assumption that the diffusion model can be applied and it should be noted that the major error is introduced by using Eq. [6]. The relationship between $\Delta\sigma$ and C expressed in Eq. [6] can be checked for

the similar molecules CO and COS, where both quantities are known from molecular beam data (14, 15). The agreement with experiment is 4% for CO and 10% for COS which is within the experimental error for $\Delta\sigma$.

The correlation times τ_{SR} and τ_c calculated from the relaxation rates with these parameters are given in Table 1. The values for τ_c agree quite well with correlation

TABLE 1
CORRELATION TIMES FOR ANGULAR VELOCITY τ_{SR} AND ANGULAR REORIENTATION τ_c in CS_2

| T (°K) | τ_{SR} (sec) | τ_c (sec) |
|----------|--------------------------|----------------------|
| 308 | $10.0 \cdot 10^{-14}$ | $1.0 \cdot 10^{-12}$ |
| 293 | $8.7 \cdot 10^{-14}$ | $1.4 \cdot 10^{-12}$ |
| 273 | $7.6 \cdot 10^{-14}$ | $1.5 \cdot 10^{-12}$ |
| 248.5 | $6.3 \cdot 10^{-14}$ | $2.1 \cdot 10^{-12}$ |
| 221 | $5.4 \cdot 10^{-14}$ | $2.4 \cdot 10^{-12}$ |
| 193 | $3.9 \cdot 10^{-14}$ | $4.1 \cdot 10^{-12}$ |
| 167 | $1.8 \cdot 10^{-14}$ | $8.4 \cdot 10^{-12}$ |

times estimated from light scattering data (16). It is seen that the diffusion model should be a good description at the lowest temperatures. For the highest temperature studied, on the other hand, the difference in the correlation times is only one order of magnitude. The experimental fact that $1/T_1^{\text{SR}} \cdot 1/T_1^{\text{CS}}$ is constant over the entire temperature range (Fig. 3) indicates, however, that even for the highest temperatures the deviations from the diffusion limit are within experimental error.

Spin-rotation constants are extremely difficult to compute from first principles as such a calculation depends on the knowledge of all the excited energy levels of the molecule. By use of Eq. [11], however, semiempirical values for the spin-rotation constant can be calculated with which our data may be compared. For linear molecules the anisotropy of the chemical shift can also be calculated from a known value of $\Delta\sigma$ for a different linear molecule and the relative isotropic chemical shift, using the assumption that σ_{\parallel} is constant for linear molecules as outlined above.

A molecule suitable for comparison is ^{13}CO : The spin-rotation constant of ^{13}C is known from molecular beam data with high accuracy (14), the anisotropy of the chemical shift is also given in Ref. (14), and the relative isotropic chemical shifts of ^{13}CO , $^{13}\text{CO}_2$, and $^{13}\text{CS}_2$ have been determined (17). In Table 2 values calculated in this way for C and $\Delta\sigma$ are given for CO_2 and CS_2 together with the experimental data for COS (15). The experimental and calculated absolute values of C and $\Delta\sigma$ for CS_2 agree within experimental error. Furthermore, the comparison of the calculated values for CO_2 and CS_2 with the parameters measured for COS shows that the signs are also correct as obtained from the calculation. We, therefore, conclude that in CS_2 the ^{13}C spin-rotation constant is negative and the anisotropy of the chemical shift is positive, as shown in Table 2. The close agreement of calculated and observed

TABLE 2
CALCULATED AND OBSERVED VALUES FOR SPIN-ROTATION CONSTANTS AND ANISOTROPIC
CHEMICAL SHIFTS IN $^{13}\text{CO}_2$, ^{13}COS , and $^{13}\text{CS}_2$

| | C (kHz) | | $\Delta\sigma$ ppm | |
|----------------------------|-----------------|-------|--------------------|-------|
| | exptl | calcd | exptl | calcd |
| CO_2 ^a | | -33.9 | | 315 |
| COS ^b | -19.5 ± 1.3 | | 372 ± 42 | |
| CS_2 ^a | -13.8 ± 1.4 | -12.6 | 438 ± 44 | 428 |

^a This work. ^b See Ref. (15).

values demonstrates that our method of eliminating the correlation times allows the determination of spin-rotation constants and shift anisotropies from relaxation studies. This is particularly important for symmetric molecules like CS_2 , which are unsuitable for microwave or electric molecular beam resonance experiments.

Note added in proof: Recently $\Delta\sigma$ has been determined (18) in solid $^{13}\text{CS}_2$. The value obtained, $\Delta\sigma = 425 \pm 15$ ppm, compares remarkably well with our result.

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