High Resolution and Solid State NMR Investigations of Subvalent Gallium Compounds*

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The $^{71}$Ga NMR spectra of Ga[ GaX₄] melts and of solutions in benzene and other hydrocarbons show discrete sharp Ga¹ and broad Ga²⁺ resonances. In the light of recent structure determinations, the solution Ga¹ signals must be attributed to bis(arene)Ga⁺ complexes in which the gallium atom is $σ^2$-bonded to the hydrocarbons. The low line widths and strong high field shifts are attributed to an almost spherical shielding of the metal nucleus by the $4s^2$ electrons. Solid state $^{69}$Ga and $^{71}$Ga NMR spectra of Ga[ GaCl₄] crystalline powder show only Ga¹ resonances. While the $^{71}$Ga¹ line is rather narrow, the $^{69}$Ga¹ line has a complex fine structure. Consistent with the crystal structure of Ga[ GaCl₄], the Ga¹ ion is calculated to have a very low quadrupole coupling constant $\eta = 0.44$. Experimental and simulated line shapes (using literature models) are in satisfactory agreement, implying that the $^{69}$Ga signal splitting is due to second order quadrupolar effects for the central $m_I = +1/2 \Rightarrow -1/2$ transition. The analogous splitting of the $^{71}$Ga NMR line is too small to be detected.

Introduction

Subvalent gallium compounds with the metal in the non-classical oxidation states +1 or +2 have attracted considerable attention ever since their discovery in the first decades of this century [1]. Many physical methods have been used to elucidate various features of structure, bonding, and reactivity, but it was not until 1965 that NMR spectroscopy could be employed for these purposes [2, 3]. Very recently it appears that $^{71}$Ga NMR could become almost a routine technique in studies of reaction mechanisms and products [4, 5].

In the course of our own investigations on novel arene complexes of gallium and its homologues [1, 5–10] it was also necessary to reconsider the structures of the arene-free Ga[ GaX₄] species. The original X-ray structure determination of the chloride, Ga[ GaCl₄], was not very accurate on modern standards [11]. Interestingly enough, a recent study of In[ InCl₄] confirmed the structural concept of the Ga/Cl analogue [12], while the compound Ga[ GaI₄] was shown to have an entirely different structure [13].

In order to contribute to the questions of the environmental symmetry of the gallium(1) cations in the crystal, in the melt, and in non-aqueous solutions, Ga NMR spectroscopy was used for studies of Ga[ GaCl₄] in all three phases. Ga[ GaBr₄] was included into the solution studies.

Experimental

Samples: Ga[ GaCl₄] and Ga[ GaBr₄] were prepared according to literature methods [1] and their identity checked by melting points and analytical data. Solutions were obtained from high purity aromatic hydrocarbons, which were dried and saturated with nitrogen.

Spectra: For both the solid state and the solution measurements a Bruker CXP 200 was used. For the $^{69}$Ga powder pattern the Larmor frequency was set at $v_L = 47.989$ MHz, for $^{71}$Ga at $v_L = 60.790$ MHz. A pulse width of 4 $\mu$s was used ($DL = 50$ ms, $NS = 2000$, $F = \pm 200$ Hz). HCl-acidic aqueous GaCl₃ solutions served as reference ($\delta = 0$ ppm).

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For the $^{71}$Ga solution spectra the spectrometer frequency $v = 60.990$ MHz was employed. HNO$_3$-acidic aqueous Ga(NO$_3$)$_3$ served as an external standard.

For the calculations of the spectrum shown in Fig. 3 the literature procedure was followed [14].

**Results**

I. $^{71}$Ga and $^{69}$Ga NMR Studies of Polycrystalline Ga$[GaCl_4]$ 

The $^{69}$Ga NMR spectrum of gallium(1) tetrachlorogallate(III) as measured at 303 K exhibits a resonance with a complex fine structure as shown in Figure 1. The central line appears at $\delta = -844$ ppm rel. to the aqueous Ga$^{3+}$ standard. The $^{71}$Ga NMR spectrum consists of only one, relatively sharp resonance (Fig. 2) with the same chemical shift.

Considering the relations $Q(^{69}$Ga) > $Q(^{71}$Ga) and $\mu(^{69}$Ga) < $\mu(^{71}$Ga) [14], it follows that the fine structure of the $^{69}$Ga signal is not due to shift anisotropy effects, but originates from second order quadrupole effects associated with the central $m = \pm 1/2 \rightarrow -1/2$ transition. According to theoretical calculations, using asymmetry parameters $\eta > 1/3$, these effects lead to signal shapes as shown in Figure 3. The agreement with the experimental spectrum is quite satisfactory for values $e^2qQ/h(^{69}$Ga) = 1.7 ± 0.1 MHz and $\eta = 0.44 ± 0.05$ (see Experimental Part).

For the corresponding $^{71}$Ga quadrupole coupling constant a value $e^2qQ/h(^{71}$Ga) = 1.1 ± 0.1 MHz is obtained from the above data. This value should give rise to a maximum splitting of the main frequency edges of $\Delta v = 1500$ Hz, well below the experimental line width of $\Delta B = 3070$ Hz for the $^{71}$Ga NMR resonance. Quadrupolar splittings are therefore not to be expected in this $^{71}$Ga NMR experiment.
The observed $^{69}$Ga, $^{71}$Ga NMR signals are assigned to the Ga$^1$ ions present in the Ga[GaCl$_4$] sample. The severe distortion of the tetrahedral chloride environment of the Ga$^{III}$ centers [11] is probably responsible for the absence of Ga$^{III}$ NMR resonances. For the quadrupole coupling constants of the Ga$^{III}$ nuclei the condition $e^2 q Q / h (\text{Ga}^{III}) \ll \mu B / h$ is not valid, and their effects exceed the experimental limit of this study.

The low quadrupole coupling constants determined for $^{69}$Ga$^1$ and $^{71}$Ga$^1$ in crystalline Ga[GaCl$_4$] indicate that the dodecahedral coordination sphere of the univalent ions is distorted only slightly from the maximum $D_{2d}$ symmetry. The orthorhombic space group is in agreement with this result [11]. The value for $\eta = 0.44$ also shows that the symmetry of the electrical field gradient tensor at the Ga$^1$ cation is reduced according to the orthorhombic crystal symmetry.

As compared to Ga$^{III}$ species, these effects are relatively small. Clearly, the presence of a $4s^2$ lone pair of electrons at Ga$^1$ leads to a high and symmetrical shielding of the gallium nuclei.

II. $^{71}$Ga NMR Spectrum of Molten Ga[GaCl$_4$].

The $^{71}$Ga NMR spectrum of molten Ga[GaCl$_4$] (at 200 °C) has already been reported by Greenwood and his collaborators [2]. Two resonances were found, which were assigned to Ga$^1$ and Ga$^{III}$, respectively. The Ga$^1$ signal was surprisingly narrow, while the GaX$_4^-$ line showed the usual broadening typical of many gallium resonances.

These results could be confirmed in more recent experiments employing modern equipment. There are similarities between the spectra of aromatic hydrocarbon solutions and the spectrum of the melt, especially regarding the chemical shifts and line widths of the GaX$_4^-$ resonances. This result suggests not only that assignments are correct, but also that the anionic parts of the samples are strongly coordinated to the Ga$^1$ centers thus rendering the GaX$_4^-$ tetrahedra strongly distorted. The environment of the Ga$^1$ centers appears to be of only slightly reduced symmetry as judged from the low line widths of the $^{71}$Ga$^1$ signals. A quasi-dodecahedral halogen cage seems to be an acceptable model for the coordination array around Ga$^1$ in the melt.


In recent structural studies it could be demonstrated that Ga[GaX$_4$] salts are taken up by aromatic solvents as 1:1 or 1:2 arene complexes [1, 5–10]. These arene adducts can be crystallized as oligo-
meric species, in which the Ga$^+$ center is hexahapto coordinated ($\eta^6$) to the arene ring, i.e. perpendicularly above the rings. In the 1:2 complexes the two $\eta^6$-bonded rings form angles of 45–60°.

The high resolution $^{69}$Ga NMR spectra of the solutions of these complexes in excess hydrocarbon are all characterized by a broad GaCl$_4$ or GaBr$_4$ resonance and a sharp (arene)$_2$Ga$^+$ signal (Figure 4). The spectrum of a benzene solution of Ga[GaCl$_4$] has been reported previously [2, 3]. The drastic shift difference of the low-field signals obtained for chloride and bromide, together with a much smaller shift difference of the sharp-field signals, is again convincing evidence for the assignment to Ga$^{III}$ and Ga$^+$, respectively.

The chemical shifts of the gallium(I) resonances depend on the nature of the arene employed as a solvent. This effect is of course due to the direct arene-gallium(I) complexation. For mixtures of arenes, the gallium(I) shift observed corresponds to the value for the complex of the most electron-rich hydrocarbon (hexamethylbenzene > mesitylene > xyylene > toluene > benzene). It is concluded that the metal ion is coordinated to the methyl-richer component.

Mono- or bis(arene) complexes cannot be distinguished by Ga NMR in solution at room temperature. There is rapid arene exchange of coordinated benzene and free benzene, and also between mono- and bis-complex ions. Even cooling of toluene solutions below −50° gave no signal splitting. The benzene exchange process therefore is associated with only very low energies of activation, probably lower than 9 kcal/mole.

Separate signals are registered for mixtures of Ga$_2$Cl$_4$ and Ga$_2$Br$_4$ with arenes consisting of two liquid phases. As soon as the phase border disappears upon further dilution with excess solvent, the second signal is no longer visible.

The narrow lines of the gallium(I) resonances and their extreme high field shifts are indicative of a strong symmetrical shielding of the Ga$^+$ nuclei in the arene complexes. This effect is again attributed to the presence of the 4$s^2$ lone pair of electrons which provides a first “coordination sphere” of high shielding efficiency followed by the arene and halogen ligands at much larger distances.

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