The following full text is a publisher's version.

For additional information about this publication click this link.
http://hdl.handle.net/2066/143018

Please be advised that this information was generated on 2019-12-22 and may be subject to change.
Studies of Solvation Shell Changes from the Nuclear Magnetic Resonance Linewidth of $^{23}$Na

By A. M. Groren and J. Smid

(Chemistry Department, State University of New York, College of Forestry, Syracuse, New York 13210)

and E. de Boer

(Department of Physical Chemistry, University of Nijmegen, Nijmegen, The Netherlands)

Reprinted from

Chemical Communications 1971

The Chemical Society, Burlington House, London W1V OBN
Studies of Solvation Shell Changes from the Nuclear Magnetic Resonance
Linewidth of $^{23}\text{Na}$

By A. M. Groten and J. Smid*
(Chemistry Department, State University of New York, College of Forestry, Syracuse, New York 13210)
and E. de Boer
(Department of Physical Chemistry, University of Nijmegen, Nijmegen, The Netherlands)

Summary
Addition of polyglycol dimethyl ethers or macrocyclic polyethers to a solution of sodium tetraphenylboron ion-pairs in tetrahydrofuran strongly broadens the $^{23}\text{Na}$ n.m.r. line, indicating that the tetrahydrofuran solvation shell is replaced by the complexing agent.

The solvation state of ions or ion-pairs frequently affects their physical properties e.g. their optical and e.s.r. spectra, and reactivity. In mixtures of solvating agents, preferential complexation with the ionic species may occur, and the composition of the solvation shell becomes an important variable. $^1$H N.m.r. spectra are useful in the identification of the solvent species complexed to an ion or ion-pair, especially when the ionic entity is either paramagnetic or exhibits strong diamagnetic anisotropy. However, in many cases the proton shifts of the ion solvation shell molecules are small or masked by the solvent.

We report that the addition of small quantities of alkali complexing reagents to solutions of salts such as sodium tetraphenylboride markedly affects the linewidth of the $^{23}\text{Na}$ n.m.r. signal, indicating a change in the solvation shell around the Na$^+$ ion. For nuclei with spin quantum number $I > \frac{1}{2}$, particularly for Na which has a comparatively large quadrupole moment, the linewidth $(1/T_{2\text{re}})$ is largely determined by quadrupole relaxation [equation $(1)$],

$$\left(\frac{1}{T_{2\text{re}}}ight) = \frac{3}{4\pi} \frac{2I + 3}{I^2(I - 1)} \left(\frac{\alpha Q}{h}\right)^2 \left(1 + \frac{k^2}{3}\right) \tau_r$$

(1)

where $\tau_r$ refers to the rotational correlation time characterizing the tumbling motion of the ion or ion-pair, $\alpha Q$ is the nuclear electric quadrupole moment, $\eta$ denotes the largest value of the electric field gradient, and $\kappa$ is the asymmetry parameter.

By combining the expression $\tau_r = 4\pi \eta a^2/3RT$ ($\eta$ is the solution viscosity and $a$ is the radius of the ionic species approximated by a sphere) with equation $(1)$, equation $(2)$ ($\kappa = 0$) is obtained.

$$\left(\frac{1}{T_{2\text{re}}}\right) \approx \left(\frac{\alpha Q}{h}\right)^2 \frac{a^2 \eta}{T}$$

(2)

We measured the linewidth of the $^{23}\text{Na}$ n.m.r. signal of a
0.2M-tetrahydrofuran (THF) solution of NaBPh₄ as a function of temperature, then added 0.2M-hexaethyleneglycol dimethyl ether (MeO(CH₂CH₂O)₆Me or glyme 7) (I) or 0.2M-dimethylidibenz-18-crown-6 (II) and repeated the measurements. The $^{23}$Na resonance signal was obtained on a Varian DP 60 EL spectrometer at a frequency of 15.1 MHz, and operating in the high resolution mode. With the crown compound present the spectrometer was operated in the wide line mode. The chemical shifts were measured with an AEG gaussmeter, using a 1M-aqueous NaCl solution as standard.

Changes in the $^{23}$Na chemical shift from 5.7 p.p.m. (NaBPh₄-THF) to 9.2 p.p.m. [NaBPh₄-(I)-THF] to 14.9 p.p.m. [NaBPh₄-(II)-THF] were observed, all shifts being upfield from NaCl and measured at 40 °C. The shifts are comparable to those found for NaBPh₄ in pure tetraglyme and indicate a change in the solvation shell around the Na⁺ ion. Such alkali n.m.r. shifts have also been reported for sodium and lithium salts on changing the solvent medium and can be correlated with the solvating ability of the respective solvents. The Figure shows changes in the linewidths, as $1/T_2$ vs. $\eta/T$ plots. The viscosities, $\eta$, (measured with an Ubbelohde viscometer) are those of the 0.2 M-salt solutions in the absence or presence of the complexing agent. The large change in the $^{23}$Na linewidth of the NaBPh₄ salt on addition of an equimolar quantity of (I), and even larger change on addition of (II) is evidence that in the solvent separated NaBPh₄ ion-pair the THF solvation shell around the Na⁺ ion is replaced by (I) or by (II). Studies with small quantities of tetraglyme show similar behaviour and addition of more than an equimolar quantity of (I), after correction for the increased viscosity, does not lead to further line broadening, indicating the formation of 1:1 complexes between the crown compound or glymes and the NaBPh₄.

Two factors may be responsible for the increased linewidth, viz., a change in the rotational correlation time $\tau_r$ of the ion-pair on changing the solvation shell around the Na⁺ ion, or a change in the electric field gradient at the Na nucleus. We suggest that the latter factor is the more important one. The overall size of the solvated ion-pair does not change significantly when the THF solvation shell is replaced by either (I) or (II). Viscosities of the 0.2M-NaBPh₄ solutions are nearly identical in the absence and presence of the complexing agent. However, in the THF-separated ion-pair the oxygen atoms are nearly spherically distributed around the Na⁺ ion. This is probably not the case for compound (I) and certainly not for (II) which derives its strong alkali complexing property largely from the fact that the polyether ring can assume a nearly planar conformation, with the Na⁺ ion situated in the hole of the polyether ring. This non-spherical type of solvation would strongly increase the electric field gradient contributed by the lone electron pairs of the oxygen atoms at the Na nucleus. The small curvature in the linewidth plots may indicate that slight changes take place in the solvated ion-pair structure as the temperature is lowered, e.g., an increased ordering around the Na⁺ ion.

This method is useful in the study of ion and ion-pair solvation, especially in cases where $^1$H n.m.r. studies do not provide conclusive evidence. Also, the complexation of certain crown compounds with sodium ions or ion-pairs may be slow enough for a kinetic study to be made from the change in the linewidth of the $^{23}$Na n.m.r. signal.

We thank the National Institutes of Health and the University of Nijmegen, for financial support.

(Received, April 19th, 1971; Com. 896.)

---