Studies on Succinic Acids

IV.* Evidence for the Existence of Intramolecular Hydrogen Bonding in Certain Highly Alkylated Succinic Acids by Infra-Red Spectra

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The infra-red spectra of racem-\(a,\alpha\)'-diisopropyl-, racem-\(a,\alpha\)'-dicyclohexyl- and tetraethylsuccinic acid and their monopotassium salts have been found to be consistent with an intramolecularly hydrogen-bonded structure. All these acids have very large \(K_1/K_4\) ratios. The infra-red spectra of meso-\(a,\alpha\)'-diisopropyl-, meso-\(a,\alpha\)'-di-(tert.-butyl)-, and meso-\(a,\alpha\)'-dicyclohexylsuccinic acid and their monopotassium salts are compatible with a structure containing one carboxyl group and one carboxylate ion which are essentially independent of each other.

There is chemical and physico-chemical evidence for the existence of an intramolecular hydrogen bond in certain dibasic acids and their monoanions \(^1-^4\). These include highly branched malonic acids (I), racem-\(a,\alpha\)'-dialkylsuccinic acids (II) with bulky alkyl groups, tetraalkylsuccinic acids (III), \(\beta,\beta\)'-dialkylglutaric acids (IV), \(\text{cis-1,2-cyclopropane dicarboxylic acids (V), cis-1,2-dicarboxyethylenes (VI), and others.}

\[
\begin{align*}
\text{I} & \quad \text{II} & \quad \text{III} \\
\text{IV} & \quad \text{V} & \quad \text{VI}
\end{align*}
\]

In all these cases the steric relationships probably play an important role. For example, it is well known that the accommodation of alkyl groups in a carbon chain capable of ring closure will increase the ease of formation and stability of the corresponding cyclic compound. In an alkylated dibasic acid the formation of a cyclic structure can be accomplished by the formation of an intramolecular hydrogen bond, provided the arrangement of alkyl groups will be energetically favoured. In the other types of dibasic acids the carboxyl groups are rigidly held at distances which make hydrogen bonding possible. The most

*Fig. 1. Infra-red spectra of racem-α,α'-disopropylsuccinic acid (A) and its monopotassium salt (S).*

important arguments for the existence of an intramolecular hydrogen bond in dibasic acids of these kinds are:

1. They have anomalously high ratios between their first and second dissociation constants, which cannot be explained on the basis of electrostatic effects alone \(^1\),\(^3\).

2. Measurements of the equilibrium constants of the reaction meso-acid ⇄ racem-acid for \(\alpha,\alpha'\)-dialkylsuccinic acids indicate that the racemic form is the

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*Fig. 2.* Infra-red spectra of *meso*-\(\alpha,\alpha'\)-disopropylsuccinic acid (A) and its monopotassium salt (S).

more stable isomer \(^2\), i.e., the stability order is the reverse of that predicted from conformational analytical considerations.

3. They are very easily anhydridized, some of them even in aqueous solution or in the solid state \(^4\).

Shahat \(^6\) has shown by X-ray analysis that in crystalline maleic acid there exists an unsymmetrical hydrogen bond. Cardwell \textit{et al.}\(^7\) have investigated maleic acid and its monopotassium salt in the solid state by means of infra-red

\textbf{Fig. 3.} Infra-red spectra of racem-\(\alpha,\alpha'-\text{di-(tert.-butyl)}\)-succinic acid (A) and its monopotassium salt (S).

\textit{Acta Chem. Scand.} 13 (1959) No. 2
spectra and found evidence for an intramolecular hydrogen bond in both acid and salt. In the latter case the absence of the carbonyl band indicated that the hydrogen bond might be symmetric.

The purpose of the present investigation was to study the infra-red spectra under various conditions for a series of alkylsubstituted succinic acids and their monopotassium salts. Four acids, racem-a,a'-diisopropyl-, racem-a,a'-di-(tert.-butyl)-, racem-a,a'-dicyclohexyl-, and tetraethylsuccinic acid, which exhibit very high $K_1/K_2$-ratios ($4 \times 10^8$--$3.5 \times 10^8$ in 50 % aqueous ethanol) and three acids, the meso forms of the above acids, with low $K_1/K_2$-ratios (about 100) have been selected, and the infra-red spectra of the acids and their monopotassium salts have been examined in the solid state and in methanolic solution. Some of the salts were also studied in deuterium oxide solution. Methanol is the only solvent which meets the requirements of being transparent in the double bond stretching frequency region and of dissolving both

![Infra-red spectra](image)

*Fig. 4.* Infra-red spectra of meso-a,a'-di-(tert.-butyl)-succinic acid (A) and its monopotassium salt (S).

acids and salts in sufficient concentrations. The discussion will be limited to a
consideration of the carbonyl stretching frequency and the carboxylate asym-
metric stretching frequency, as the other bands associated with these groups
either are difficult to recognize or fall in a region where the solvent has a large
absorption. The results are shown in Figs 1—7 and in Table 1, which gives the
wave-numbers of the bands in the double bond region together with the
approximate molar extinction coefficients in methanolic solution.

At a first glance it is seen that the spectra of the low-ratio acids and their
salts are consistent with the structure VII for the acids and VIII for the
monoanions.

Thus, in the solid state the carbonyl stretching frequency of the acids has a
single peak at about 1695 cm\textsuperscript{-1}, and no significant shift occurs in the spectra
of the corresponding salts. In methanolic solution the situation is similar;

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Fig. 5. Infra-red spectra of racem-\textit{a},\textit{a}'-dicycloyhexylsuccinic acid (A) and its mono-

potassium salt (S).

\textit{Acta Chem. Scand.} 13 (1959) No. 2
however, the carbonyl band has split into two peaks at about 1720 and 1700 cm\(^{-1}\). These peaks are not shifted in the spectra of the salts in methanol, but the molar extinction coefficients are reduced to about half their values in the acids. The carboxylate asymmetric stretching frequency occurs in the solid state at 1550—1560 cm\(^{-1}\) and in methanol it is shifted to about 1570 cm\(^{-1}\). The only meso acid salt which has been studied in deuterium oxide solution

*Fig. 6. Infra-red spectra of meso-\(\alpha,\alpha’\)-dicyclohexylsuccinic acid (A) and its monopotassium salt (S).*

(the others are too slightly soluble) shows a single carbonyl peak at 1 680 cm\(^{-1}\) and the carboxylate band at 1 550 cm\(^{-1}\), i.e., both bands have shifted about 20 cm\(^{-1}\) towards lower wave-numbers as compared with the spectra in methanol. Summarizing, the double bond region of a low-ratio acid salt can be interpreted as a superposition of the carboxyl group and carboxylate ion spectra, and consequently the structure of the monoanions is best represented by VIII without interactions between the carboxyl groups.

Fig. 7. Infra-red spectra of tetraethy succinic acid (A) and its monopotassium salt (S). 

Acta Chem. Scand. 13 (1959) No. 2
Table 1. Carboxyl frequencies of alkylsubstituted succinic acids

<table>
<thead>
<tr>
<th>Succinic Acid</th>
<th>Carbonyl Stretching Frequency</th>
<th>In Methanol</th>
<th>Molar Extinction Coefficient ε</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>In KBr, wave number in cm⁻¹</td>
<td>wave number in cm⁻¹</td>
<td>In methanol</td>
</tr>
<tr>
<td>Racem-a,α'-diisopropyl-</td>
<td>1695 vs</td>
<td>1718 sh</td>
<td>1704</td>
</tr>
<tr>
<td>monopotassium salt</td>
<td>1680 sh</td>
<td>1690</td>
<td>1650 sh</td>
</tr>
<tr>
<td>Meso-a,α'-diisopropyl-</td>
<td>1692 vs</td>
<td>1723</td>
<td>1701</td>
</tr>
<tr>
<td>monopotassium salt</td>
<td>1687 sb</td>
<td>1720 sh</td>
<td>1700</td>
</tr>
<tr>
<td>Racem-a,α'-di-(tert.-butyl)-</td>
<td>1695 vs, 1615 w</td>
<td>1716 sh, 1640</td>
<td>1708</td>
</tr>
<tr>
<td>monopotassium salt</td>
<td>1675 mb</td>
<td>1682</td>
<td>1640</td>
</tr>
<tr>
<td>Meso-a,α'-di-(tert.-butyl)-</td>
<td>1690 vs, 1635 s</td>
<td>1720, 1700</td>
<td>1708</td>
</tr>
<tr>
<td>monopotassium salt</td>
<td>1687 sb</td>
<td>1718 sh, 1695</td>
<td>1690</td>
</tr>
<tr>
<td>Racem-a,α'-dicyclohexyl-</td>
<td>1700 s, 1625 sh</td>
<td>1720</td>
<td>1705</td>
</tr>
<tr>
<td>monopotassium salt</td>
<td>1690 sb</td>
<td>1695</td>
<td>1650</td>
</tr>
<tr>
<td>Meso-a,α'-dicyclohexyl-</td>
<td>1695 s</td>
<td>1717</td>
<td>1697</td>
</tr>
<tr>
<td>monopotassium salt</td>
<td>1697 m</td>
<td>1717</td>
<td>1698</td>
</tr>
<tr>
<td>Tetraethyl-</td>
<td>1683 vs</td>
<td>1716</td>
<td>1691</td>
</tr>
<tr>
<td>monopotassium salt</td>
<td>1682 m</td>
<td>1678</td>
<td>1678</td>
</tr>
</tbody>
</table>

vs = very strong; s = strong; m = medium; w = weak; sh = shoulder; b = broad.

A similar interpretation is not possible for the spectra of the high-ratio acids and their salts. In methanolic solution all the racem-acids but not tetraethylsuccinic acid show a carbonyl band of low intensity at about 1650 cm⁻¹ (racem-a,α’-di-(tert.-butyl)-succinic acid has a weak band at 1615 cm⁻¹ also.

Acta Chem. Scand. 13 (1959) No. 2
succinic acids and their monopotassium salts.

<table>
<thead>
<tr>
<th>In D$_2$O wave number in cm$^{-1}$</th>
<th>Carboxylate ion, asymm. stretching frequency</th>
<th>$\varepsilon_{C=O}$ (acid)</th>
<th>$\varepsilon_{C=O}$ (salt)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>In KBr, wave number in cm$^{-1}$</td>
<td>In methanol wave number in cm$^{-1}$</td>
<td>Molar extinction coefficient $\varepsilon$</td>
</tr>
<tr>
<td>1 645 sh</td>
<td>1 570 vsb</td>
<td>1 568</td>
<td>400</td>
</tr>
<tr>
<td>1 680</td>
<td>1 550 sb</td>
<td>1 568</td>
<td>470</td>
</tr>
<tr>
<td>1 650 sh</td>
<td>1 570 vsb</td>
<td>1 575</td>
<td>290</td>
</tr>
<tr>
<td></td>
<td>1 550 sb</td>
<td>1 568</td>
<td>450</td>
</tr>
<tr>
<td></td>
<td>1 568 vs</td>
<td>1 573</td>
<td>360</td>
</tr>
<tr>
<td></td>
<td>1 558 m</td>
<td>1 568</td>
<td>440</td>
</tr>
<tr>
<td>1 650 sh</td>
<td>1 560 vsb</td>
<td>1 568</td>
<td>340</td>
</tr>
</tbody>
</table>

in the solid state). This band must be due to a carbonyl group taking part in an intramolecular hydrogen bond (IX). A band at 1 630 cm$^{-1}$ in solid maleic acid has been explained in the same way. Accordingly, the peaks at 1 720 and 1 700 cm$^{-1}$ are assigned to the other carbonyl group which may take

*Acta Chem. Scand. 13 (1959) No. 2*
part in an external hydrogen bond. The cause of the splitting around 1 700 cm\(^{-1}\) is probably coupling of the carbonyl frequencies\(^8-10\), and not the existence of monomers and polymers in solution as the relative intensities of the two bands are independent of concentration.

The spectra of the high-ratio acid salts show the following main features. In both the solid state and in solution the salts have a large, almost continuous absorption in the region 1 600—1 400 cm\(^{-1}\). In methanolic solution the carbonyl frequency has shifted by 10—25 cm\(^{-1}\) towards lower wave-numbers as compared with those of the acids, which is indicative of an intramolecular hydrogen bond\(^11\). The carbonyl band has been considerably reduced in intensity and its molar extinction coefficient is only a small fraction of that in the acid. In the hydrogen maleate ion the carbonyl band has completely disappeared\(^7\), which points to the existence of a symmetric hydrogen bond in this ion with a structure intermediate between the extreme structures \(X_a\) and \(X_b\).

\[
\begin{align*}
X_a & \quad CH\equiv CH \\
O\equiv C & \quad C-O^- \\
O-H\cdots O & \\
\quad CH\equiv CH \\
O\equiv C & \quad C-O^- \\
O\cdots H-O & \\
\end{align*}
\]

The fact that there actually is found a carbonyl peak in the spectra of the salts in question, though of appreciably reduced intensity, suggests that a structure for the monoanion with an unsymmetric intramolecular hydrogen bond (XI) is best compatible with the spectral data. In the deuterium oxide solution of a high-ratio acid salt the carbonyl peak has shifted towards lower wave-numbers and the carboxylate peak towards higher wave-numbers. This behaviour lends further support to an intramolecularly hydrogen-bonded structure.

**EXPERIMENTAL**

The acids investigated in this paper were prepared and purified as described previously\(^4\). The monopotassium salts were prepared by dissolving accurately weighed amounts of acids and potassium carbonate (analytical grade reagent) in water or aqueous

ethanol by slight warming, filtering, and then evaporating in vacuo at room temperature. The last traces of solvent were removed by warming at 110° for 3 h. The purity was checked by analysis for carbon and hydrogen. It is by no means certain that a solid salt spectrum actually refers to the hydrogen potassium salt, but these spectra are given for comparison. The discussion above is based on the salt spectra in methanolic solution, where only the ion species HA⁻ is present.

The infra-red spectra were recorded by a Perkin Elmer model 21 spectrophotometer equipped with a sodium chloride prism. The solid samples of the acids and salts were examined in pressed potassium bromide plates. The wave-numbers given in Table 1 are accurate within ±5 cm⁻¹.

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REFERENCES


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