NEW DEVELOPMENTS IN CATIONIC AND ANIONIC ALDEHYDE POLYMERIZATION

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Abstract—Aldehydes polymerize readily with cationic and anionic initiators to polyacetals. Higher aliphatic aldehydes are conveniently polymerized with anionic initiators at low temperatures, frequently to isotactic polyaldehydes. The range of known polyaldehydes has recently been extended to poly(n-dodecaldehyde). Polymers with side chains longer than C₃ possess the characteristics of the crystallization of the aliphatiç side chain separate and in addition to the crystallization of the main chain.

An NMR investigation of the rate of polymerization and copolymerization of chloral showed that the polymerization becomes diffusion controlled above approximately 40% conversion. At that conversion small molecules are entrapped in the polymer matrix and individual parts of these molecules have different mobility as may be detected by NMR.

Bromal has recently been polymerized at low temperatures where its crystallization was avoided by judicious selection of solvent and temperature of polymerization. The tribromomethyl group of bromal is the largest side group of any monomer which has ever been polymerized. Dichlorobromoacetaldehyde and chlorodibromoacetaldehyde were synthesized and polymerized to what are completely insoluble and apparently isotactic polymers. Fluorodichloroacetaldehyde and difluorochloroacetaldehyde were also synthesized and polymerized. Fluorodichloroacetaldehyde gives only insoluble and apparently isotactic polymers. Difluorochloroacetaldehyde gave soluble amorphous polymers in addition to crystalline polymers.

Future trends in aldehyde and carbonyl polymerization are discussed and attempts have been made to identify unsolved problems in this area. The successful preparation of tetrachloroethylene oxide, in good yield, has been disclosed together with the failure of all attempts to homopolymerize this compound or copolymerize it with potential comonomers.

I. PRESENT STATE OF ALDEHYDE POLYMERIZATION

Aldehydes are polymerized with ionic initiators to polyacetals but with a much lower H°/mol and consequently also a lower ceiling temperature than α-olefins of comparable structure are polymerized to polyolefins. In ethylene(CH₂CH₂) which is compared with (CH₂O) formaldehyde, the carbonyl oxygen is formally replaced by a terminal CH₂ group. Consequently, propylene (CH₃CHCH₂) should also be compared with acetaldehyde (CH₃CHO).

By far the most important polyaldehyde is polyoxy-methylene (polyformaldehyde). It is now commercially produced by anionic polymerization of purified formaldehyde. The initiation of this polymerization is often carried out by an acylate and the propagation of this polymerization is via the growing alkoxide anion. Since the acylate is not a very efficient initiator for formaldehyde polymerization, much of the polymerization is dominated by chain transfer to chain transfer agents present in the monomer, for example, water and alcohols.

\[ R^\ominus + CH₂O \rightarrow R-CH₂-O^\ominus \]
\[ R-CH₂-O^\ominus + n.CH₂O \rightarrow R-(CH₂-O)n-CH₂-O^\ominus \] (1)

Cationic polymerization of formaldehyde may also be carried out but polymers of relatively high molecular weight (polyoxymethylene) can only be prepared with cationic initiators which have very stable anions because of the ease of chain termination by recombination with anion fragments. Cationic initiation, propagation and termination of formaldehyde polymerization can be described in a very similar manner to anionic polymerization.

\[ R^\ominus + O=CH₂ \rightarrow R-O^\ominus=CH₂ \]
\[ R-O^\ominus=CH₂ + n.OCH₂ \rightarrow R-(O-CH₂-O)n=CH₂ \] (2)

Anionic copolymerizations of formaldehyde could be very conveniently carried out with monomers which could undergo anionic homopolymerization such as isocyanates and ketenes, while cationic copolymerization of formaldehyde has been reported with compounds which normally undergo cationic homopolymerization such as vinyl ethers and cyclic ethers.

The discussion of the polymerization of substituted aldehydes is most conveniently divided into the polymerization of higher aliphatic aldehydes, including the cyclopolymerization of aliphatic aldehydes, and the polymerization of haloaldehydes. Higher aliphatic aldehydes may be polymerized with anionic and cationic initiators but require strong nucleophiles such as alkoxides while relatively weak electrophiles are effective. For the initiation of the polymerization of haloaldehydes, weak nucleophiles may be, but strong electrophiles must be used, to be effective. It should also be pointed out that in essentially all these polymerizations, the polymer precipitates during the polymerization as a crystalline polymer which adds an additional driving force for the polymerization.

The most extensively studied aldehyde other than formaldehyde is acetaldehyde. It was an important...
discovery that amorphous polyacetaldehyde must be made at low temperature and provided a sound experimental basis for the principal of ceiling temperature. Subsequent work showed that the low temperature polymerization of acetaldehyde gave amorphous, atactic or crystalline isotactic forms of polyacetaldehyde with classical cationic initiators, Lewis acids, such as BF₃, or anionic initiators, such as alkali metal alkoxides respectively. It was found at the same time that aluminum alkyls, or aluminum alkyls modified by reaction with alcohols, amines and other organic compounds, could give highly effective initiators for acetaldehyde polymerization and produced initiators which were capable of polymerizing acetaldehyde to highly isotactic polymers. Modified aluminum alkyls and zinc alkyls are not only good initiators for the polymerization of acetaldehyde and higher aliphatic aldehydes but are very effective for the polymerization of ethylene oxide, propylene oxide and other cyclic ethers, thio ethers, etc. lactones and lactams.

Aliphatic aldehyde polymers became increasingly regular as the length of the side group increased from acetaldehyde to n-butyraldehyde. The phenomenon of a dual melting point was noticed starting with the crystalline n-valeraldehyde polymer, but is particularly pronounced in the case of poly(n-heptaldehyde). A further development in aldehyde polymerization was the systematic study of the polymerization of aliphatic aldehydes with polar side groups which is still proceeding. These monomers were predominantly addition products of acrolein, for example, the nitriles, ethers, thio ethers and esters of beta substituted propionaldehydes.

Dialdehydes with the proper spacing of 2–3 carbon atoms between the aldehyde groups could be cyclopolymerized, as, for example, in the case of succinaldehydes, glutaraldehydes, but also in the case of unsaturated and aromatic dialdehydes, for example, malealdehyde and phthalaldehyde. This ability to polymerize aromatic and unsaturated dialdehydes is in contrast to the fact that, aromatic monoaaldehydes, such as benzaldehydes, where the aldehyde group is directly attached to the aromatic ring, cannot be polymerized.

Polymers of haloaldehydes have also been studied in the past, particularly the polymerization of chloral. A significantly new development of recent years is the discovery of the cryotachensic polymerization of chloral which allows the preparation of films and sheets, the fabrication of the infusible and insoluble chloral polymer, by monomer casting.

Most haloaldehyde polymers are polymers of substituted acetaldehydes which are halogenated to various extents because compounds with fully halogenated longer aliphatic chains cannot be prepared with the exception of perfluoro carbonyl compounds. Examples of perfluorodialdehydes are known and their polymerization has been described. Fluoral has been polymerized to crystalline, presumably isotactic, polyfluoro and amorhous, presumably atactic, fluorocarbon soluble polyfluorol. The kinetics and the thermodynamic behavior of fluoropolymerization have also been studied.

At the present time aldehyde polymerization is studied in the following areas: (a) the polymerization and copolymerization of aldehydes with normal aliphatic side chains, (b) the preparation of aliphatic aldehyde polymers with various functional groups, (c) more extended studies of chloral polymerization and (d) the preparation and polymerization of perhalogenated acetaldehydes including acetaldehydes with F, Cl and Br substitution.

II. POLYMERS OF HIGHER ALIPHATIC ALDEHYDES
A. Structural characteristics of polymers of high aliphatic aldehydes

We had earlier recognized that crystalline polymers of n-valeraldehyde, n-heptaldehyde and n-octaldehyde showed the behavior of a "double melting point." It appeared reasonable to suggest that the higher melting point (near 150°C) was associated with the melting of the polyoxymethylene backbone chain (equilibrium melting point of polyoxymethylene is 178°C), and the lower melting point was associated with the melting of a separately crystallized phase of the waxy aliphatic side chains.

We have now investigated more thoroughly the preparation and the melting behavior of the C₅ to C₈ aldehyde polymers, with particular emphasis on the behavior of crystallization and melting of the aliphatic side chain. Crystalline poly(n-butyraldehyde) shows only one melting point near 245°C. Poly(n-valeraldehyde) did not show distinctly the behavior of side chain crystallization. Much of our work was done with poly(n-heptaldehyde) because this polymer shows this behavior most clearly (Fig. 1).
ane at −78°C with lithium tertiary butoxide as the initiator.

The polymer was acetate capped with a mixture of acetic anhydride/pyridine and was of moderate molecular weight (~10,000) as judged by osmometric determination. Poly(n-heptaldehyde) (PHA) was highly crystalline and according to 300 MHz PMR spectroscopy, it was isotactic.

\[
\text{C}_6\text{H}_{13} \quad \text{(CH}_3\text{)}_3\text{CO}+\text{n C}=\text{O}
\]

isotactic.

Poly(n-heptaldehyde) (PHA) was highly crystalline and weight (−40,000) as judged by osmometric determination. The polymer was acetate capped with a mixture of acetic anhydride/pyridine and was of moderate molecular weight (~10,000) as judged by osmometric determination. Poly(n-heptaldehyde) (PHA) was highly crystalline and according to 300 MHz PMR spectroscopy, it was isotactic.

Transition regions which could be attributed to side chain crystallization were in the range of 75–105°C. At least two transitions can be identified, one at 77–79°C and another one at 86–87°C. Figure 1 shows this region in PHA “as obtained” and after recrystallization. It can be seen that the low temperature transition at 76°C increased remarkably at the expense of the 103°C transition after slow crystallization.

Additional information was obtained by studying the transition region by extrusion just above the “melting” of the 103°C transition after slow crystallization. The crystallinity and the apparent size of crystallites are related to the temperature of the DSC transitions and their contribution to the total heat of fusion, \(\Delta H_f\).

### B. Poly(n-dodecylaldehyde)

Higher aliphatic aldehydes with side chains longer than \(n\)-octaldehyde have not yet been successfully polymerized. The failure for their polymerization could be traced to the relatively higher crystallization rate of the aldehyde as compared with its rate of polymerization. Several reports from the literature indicate that aliphatic aldehydes with longer aliphatic side chains show peculiar melting point behavior. It is apparent that association of the aldehyde molecules occurs in solution and perhaps even a liquid crystal type solution is formed under certain conditions. This might be the reason why crystallization rather than polymerization of these aldehydes occurs even in the presence of the initiator. Where polymerization temperature and temperature of crystallization are similar, very accurate temperature conditions must be maintained in order to accomplish polymerization rather than the crystallization of the aliphatic aldehyde.

When optimum conditions were chosen with lithium tertiary butoxide as the initiator in methylcyclohexane, \(n\)-dodecylaldehyde, polymerized to crystalline poly(n-dodecylaldehyde) in reasonable yields. The polymer could be endcapped and showed the following analysis. Elemental analysis: Calcd. for \(\text{C}_{12}\text{H}_{23}\text{O}\), C, 78.28; H, 12.80.

\[
\begin{align*}
\text{Poly(n-dodecylaldehyde) was allowed to react with an aqueous/alcoholic solution of 2,4-dinitrophenyl hydrazine in the presence of sulfuric acid and gave a high yield of the 2,4-dinitrophenylhydrazone of n-dodecylaldehyde with a melting point of } 104–107°C. & \\
\text{Poly(n-dodecylaldehyde) was soluble in chloroform and the i.r. spectrum could be measured in chloroform solution as well as for the neat compound. The polymer was cast onto a sodium chloride plate from CDC13 solution and the solvent evaporated. It can be seen from this spectrum (Fig. 2) that the carbonyl band of the initial monomer, n-dodecanal, is not present and only a small amount of carbonyl absorption attributable to the acetate end group is noticeable. In addition, the carbon hydrogen bands between 3000 and 2800 cm\(^{-1}\) are strong but the C–O–C absorptions in the 1200–1050 cm\(^{-1}\) are relatively weak as one would expect from the structure of this poly(n-dodecylaldehyde).} &
\end{align*}
\]

The transitions of this polymer were measured in a DSC-1B Perkin–Elmer instrument at a heating rate of 20°C/min. The results are summarized in the following table.

### Table 1. Side-chain crystallinity of oriented and unoriented poly(n-heptaldehyde)

<table>
<thead>
<tr>
<th>Polymer</th>
<th>Thermal history</th>
<th>Temperature of DSC peaks (°C)</th>
<th>(\Delta H_f) (cal/g)</th>
<th>Side chain crystallinity (%)</th>
<th>Contribution of the last DSC peak for (\Delta H_f) (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>PHA</td>
<td>Unoriented</td>
<td>86 99</td>
<td>8.4</td>
<td>21</td>
<td>67</td>
</tr>
<tr>
<td>PHA-EA</td>
<td>Extruded</td>
<td>77 104</td>
<td>3.3</td>
<td>8</td>
<td>16</td>
</tr>
<tr>
<td>PHA-EA</td>
<td>Annealed at 93°, 4 hr</td>
<td>75 103</td>
<td>6.3</td>
<td>16</td>
<td>20</td>
</tr>
<tr>
<td>PHA-EA</td>
<td>Annealed at 93°, 10 hr</td>
<td>76 104</td>
<td>7.3</td>
<td>18</td>
<td>33</td>
</tr>
<tr>
<td>PHA</td>
<td>Reference</td>
<td>87 101</td>
<td>7.3</td>
<td>18</td>
<td>61</td>
</tr>
<tr>
<td>PHA-A</td>
<td>Annealed at 90°, 2 hr</td>
<td>78 103</td>
<td>7.1</td>
<td>18</td>
<td>44</td>
</tr>
<tr>
<td>PHA-A</td>
<td>Annealed at 90°, 10 hr</td>
<td>79 105</td>
<td>7.4</td>
<td>19</td>
<td>59</td>
</tr>
</tbody>
</table>

1Poly(n-heptaldehyde) (PHA) was oriented by extrusion in INSTRON, PHA-E, at 100°C and 5 cm/min, through a capillary of \(d = 0.5\) mm and \(l = 25.4\) mm.
Fig. 2. Poly(n-dodecylaldehyde): I.R. spectrum, neat on NaCl.

20°C/min under nitrogen (Fig. 3). In the first scan two transition peaks in the region of 50–70°C were observed. The first peak had a maximum at 59°C and the second peak at which was much more prominent 69°C. After slowly cooling below the transition region and rescanning, only one peak was observed at 60°C. This is apparently the first peak which was observed in our first scanning. Although these results are of a preliminary nature, it shows clearly that the transition behavior of the side chain crystallization is qualitatively similar to that of the behavior of poly(n-heptaldehyde) where the transition region consists of at least three responses. Peak heights and transition temperatures depend on the thermal history of the poly(n-dodecylaldehyde) sample.

The thermal degradation of poly(n-dodecylaldehyde) was also measured using a TGS-1 Perkin–Elmer thermal balance; the maximum degradation temperature of this sample was 274°C at a heating rate of 20°C/min.

III. HALOALDEHYDE POLYMERS

A. Polychloral

Chloral can be polymerized most effectively and rapidly with anionic initiators. When the polymerization was carried out under cryotachensic conditions, by initiation of bulk monomer above the threshold temperature of polymerization, uniform pieces of polychloral are obtained in about 85% yield. The yield limitation is apparently not caused by the attainment of equilibrium between polymer and monomer but rather because the monomer is unable to diffuse to the reaction site. It has been found that polymerization of chloral in the presence of about 20% of an inert solvent, such as hexane, increased the mobility of the monomer and yields of more than 96% of polychloral have been obtained.

**Rate of polymerization.** We carried out rather extensive studies of the progress of the homopolymerization of chloral and its copolymerization with phenyl isocyanate by PMR spectroscopy in which the disappearance of chloral in the case of homopolymerization and the disappearance of chloral and comonomer in the case of copolymerization was determined. The rate of polymerization depends primarily on the type and amount of initiator used and is faster with increasing amounts of initiator. Since polychloral precipitated as an apparently homogeneous gel, as polymerization proceeded, the rate of polymerization decreased as the conversion increased.

The effectiveness of anionic initiators falls primarily in two main categories, those that react immediately and to a high degree with one mole of chloral to form the initiating alkoxide,

\[
\text{CCl}_3 + \text{CHO} \xrightarrow{\text{catalyst}} \text{CCI}_3\text{CHO} \]

and those whose initiation equilibrium does not favor the addition of the nucleophile to the electrophilic carbonyl carbon atom of chloral.

\[
\text{H}^+ + \text{CHO} \xrightarrow{\text{catalyst}} \text{H} - \text{O}^=\text{CH} \]

\[
\text{H}^+ + \text{CCI}_3\text{CHO} \xrightarrow{\text{catalyst}} \text{H} - \text{O}^=\text{CH} \]

Lithium tertiary butoxide reacts essentially quantitatively with chloral to form an alkoxide which can be
readily detected by proton resonance of the acetalic hydrogen at approximately 5 ppm. The other group of initiators which includes tertiary aromatic amines showed no significant reaction with one mole of chloral and no proton resonance was found at 5 ppm.

Triphenylphosphine was the most extensively studied initiator in our rate study of chloral polymerization. Triphenylphosphine reacted almost instantaneously with one mole of chloral to form triphenyl dichlorovinyl oxy phosphonium chloride. As a consequence, the actual initiator for chloral polymerization is the chloride ion. It can be seen from Fig. 4 that the polymerization of chloral with triphenylphosphine goes to completion in 5–15 min depending on the initiator concentration. At 0.1 mol% initiator concentration the polymerization is slow and proceeds at an initial rate of about 14%/min while at 0.3 mol% the initial rate is about 33%/min.

With lithium tertiary butoxide as the initiator, the polymerization is much faster and even at 0.1 mol% initiator concentration the polymerization goes to completion as fast as the PMR measurements can be made.

Other compounds which polymerize chloral belong to the group of slower initiators. As compared with triphenylphosphine, pyridine is a slightly faster initiator and tetrabutyl ammonium chloride is somewhat slower but γ-collidine is even slower than tributylammonium chloride for the polymerization of chloral using the same amount of initiator.

Interestingly enough, the rate of chloral polymerization is practically unaffected by the kind of diluent used for the polymerization. Aromatic and aliphatic hydrocarbons in up to 10–15% of diluent show similar rates. This is not really surprising because chloral monomer is much more polar than the inert solvents and is probably strongly associated and solvates the growing polymer. The relative rate of disappearance of chloral and comonomer during the copolymerization of chloral and isocyanates depends greatly on the type of isocyanates used for the copolymerization. Aromatic isocyanates are more readily incorporated into the copolymer as seen in Fig. 5.

Mobility of small molecules in polychloral matrix. In the course of our rate study of chloral polymerization by PMR spectroscopy it was found that broadening of the line width of the PMR signals of chloral as well as that of various solvents was observed as the conversion of chloral to polychloral increased.19

The line width of the solvents increased depending on the type of groups attached to the proton which is responsible for the line broadening. The chloral signal showed only a slight increase in the signal broadening during polymerization even at high conversion. In the case of toluene, the line broadening was very substantial especially in the aromatic proton region as compared with the signal broadening of the protons of aliphatic substituents. The line broadening is believed to be a spin–spin relaxation process in our system arising from time correlation functions of magnetic dipole interactions among all hydrogen atoms. Since the signals were close to Laurentzian, the spin–spin relaxation time, $T_2$, could be obtained from the spectra line width. However, various complications, particularly the inhomogeneity of the magnetic field and possible inhomogeneity of the polychloral gel, did not allow us to make accurate estimates of $T_2$. As a consequence, the line width ($\Delta H/H_0$) was used as the measure of the mobility of the small molecules in the polychloral matrix. When the ratio between the line width of chloral and also of other small molecules used as solvents, is plotted against the conversion of chloral to polychloral, curves were obtained which showed that aromatic groups are more readily broadened than some aliphatic groups especially if they are situated at the end of the molecule as is the case of toluene (Fig. 6).

It is believed that both chloral and the solvents are entrapped in the polychloral matrix. However, aromatic hydrocarbons are apparently more strongly entrapped than chloral. This behavior seems to indicate that there is a different degree of interaction between the trichloromethyl groups of the polychloral which are protruding from the polymer backbone chain and individual parts of small molecules, and cause an interaction of various degrees with small molecules. With aromatic compounds, there is a possible charge transfer type interaction between the aromatic ring and the trichloromethyl group which
polychloral: PMR line broadening of toluene and chloral signals.

prevents full mobility of the aromatic ring at high conversions; the -CHO group of chloral is essentially free.

Stabilization of chloral homopolymer by chain transfer/termination reactions. Chloral homopolymer prepared by anionic polymerization is relatively unstable. It cannot be acetylated with acetic anhydride under the normal conditions apparently because of the inability of the acetylating reagents, acetic anhydride, to penetrate to the reaction site.2° The end groups of chloral homopolymers have still not been identified because they have no characteristic absorption in the infrared spectrum. Since polychloral is insoluble and infusible, only a limited amount of characterization, most of which relies on some solution techniques, is possible. The determination of the thermal degradation spectrum is one of the major means of characterizing chloral polymers and distinguishing them from one another. Polychloral samples prepared with different initiators under different reaction conditions give distinct and characteristic spectra of polymer stability. In general, polychloral samples prepared by anionic polymerization, are less stable than those prepared by cationic polymerization. From comparative studies of anionic and cationic polymers and polymers which have been stabilized in various ways (for example, by treatment with PCl₅). The chemical nature of the stabilization is not always known and a characterization of individual polymer samples can be made by the measurement of its thermal degradation spectrum (Fig. 7).

It was believed possible to improve the stability of polychloral samples and possibly prepare polychloral of ultimate thermal stability by carrying out the polymerization in the presence of acetylation agents. We have indeed found that anionic chloral polymerization can be carried out in the presence of acetyl chloride or benzoyl chloride if the polymerization is initiated by triphenylphosphine. Up to 2 mol% of acetyl chloride could be added to the polymerization mixture without interfering with the polymerization and the conversion of monomer to polymer is similar with or without acetylating agents. Lithium tertiary butoxide could not be used as the initiator because it reacted immediately with the acetylating agent which prevented the polymerization.

During this polymerization there is an apparent competition between the addition of chloral and acetyl chloride to the growing polymeric alkoxide.

However, very little acetyl chloride reacted during the polymerization and only after the completion of the polymerization, in a much slower reaction, acetylation of the completed polymer was observed. The progress of the acetylation can be readily followed by the determination of the i.r. spectrum of the final polymer

\[
\begin{align*}
\text{Initiator concentration} & \quad 0.1 \text{ mol \%} \\
& \quad 0.2 \text{ mol \%} \\
& \quad 0.3 \text{ mol \%}
\end{align*}
\]

\[\Delta H/2 \quad \text{Hz} \]

\[\begin{array}{c}
\text{Ph} \\
\text{Me} \\
\text{Cl}
\end{array}\]

\[\begin{array}{c}
\text{Conversion, \%} \\
0 \quad 20 \quad 40 \quad 60 \quad 80
\end{array}\]

Fig. 6. Polychloral: PMR line broadening of toluene and chloral signals.

![Graph showing PMR line broadening of toluene and chloral signals.](image)

![Graph showing conversion of monomer to polymer.](image)

![Graph showing thermal stability (DTG) of various samples.](image)

![Chemical structures and equations for acetylation of polymers.](image)

Fig. 7. Polychloral: Thermal stability (DTG) of various samples.

\[
\begin{align*}
+ \text{CCl₃CHO} & \rightarrow P₁ - \text{C-O-C\text{=}O} \\
& \text{CCl₃} \quad \text{CCl₃} \\
+ \text{CH₃C=O} & \rightarrow P₁ - \text{C-O-C\text{=}O} \\
& \text{Cl} \quad \text{H} \\
& \text{Cl} \quad \text{Cl}
\end{align*}
\]
(Fig. 8) and further confirmed by a change in the thermal degradation spectrum, the appearance and increase of a peak of the DTG curve at higher temperatures, indicating the production of a more stable polychloral fraction (Fig. 9).

This result seems to indicate that chloral polymerization initiated by triphenylphosphine (triphenyl divinylloxoy phosphonium chloride) in the presence of limited amounts of acylating agents proceeds first in a fast reaction to a polychloral with alkoxide end groups that become occluded in the later part of the reaction. This has two consequences: (1) The monomer cannot penetrate to the reactive site which leads to a cessation of the polymerization at 85% conversion. (2) When acylation is attempted in a second reaction, the reagent apparently cannot diffuse to the reaction site and the acylation of the alkoxide does not succeed. On heating this unstable polymer, it will, however, degrade rather than being acetylated. If the acylating agent is, however, part of the polymerization mixture, it remains in close vicinity to the occluded but still reactive alkoxide and can then undergo the desirable acylation reaction (Fig. 10). The results of our investigation showed that one-half of the reactive sites could be acylated under our present reaction conditions. Additional work, perhaps the use of an acylating or an alkylating group as part of the counter cation, would be a more efficient way to carry out this reaction.

B. Cationic polymerization

Early work on the polymerization of chloral has shown that aluminum chloride and sulfuric acid can act as initiators. We carried out additional work in attempts to determine the scope of the cationic polymerization of chloral. Many commonly used Lewis acids, cationic salts and other cationic initiators have, however, proved to be ineffective. We have confirmed that aluminum chloride is indeed an adequate initiator for chloral polymerization but an equally good initiator was trifluoromethyl sulfonic acid. In both cases, the polymerization did not proceed in the high rate which was established for the case of anionic polymerization and the morphology of the polymer observed was different from the gel-like character of anionic polymers. SbCl₅ or SbCl₃, with 5 mol% of acetyl chloride also gave polychloral in modest yields after 3 days of reaction time. This latter combination and the establishment of trifluoromethyl sulfonic acid as a good initiator established firmly the cationic polymerization of chloral. Other initiators such as ferric chloride and titanium tetrachloride gave only a few per cent yield of polymer after several days of reaction while most other Lewis acids were ineffective.

C. Other perhaloaldehyde polymers

Chloral and fluoral polymers are the only well investigated perhaloaldehyde polymers. Fluoral may be prepared as a crystalline and presumably isotactic polymer or as an amorphous and soluble, presumably atactic polymer. Polychloral has only been prepared in one form, the crystalline insoluble, isotactic polymer. The only other perhaloacetaldehyde polymer which was described briefly in a patent was an insoluble polymer from difluorochloroacetaldehyde.

We have undertaken a more thorough study of the preparation of the unknown perhaloacetaldehydes with more than one halogen substituted in the perhalomethyl group in an attempt to study their polymerization or at least to determine their polymerizability. When polymerization was established, it was found desirable to determine whether the polymer could be made in stereoregular and/or in atactic form, in order to find out if size and shape of the side chain had any influence upon the stereoregulation of polymer formation.

Unlike olefins and vinyl ethers which require transition metal initiators whose metal atom must have the proper ligand structure and stability of the coordination complex in order to produce stereoregular polymers, stereoregularity in aldehyde polymerization can be achieved with simple initiators. Especially in the case of aldehydes with long side chains or bulky side groups such as the trichloromethyl group in chloral, stereoregular, isotactic polymers are readily formed. In the case of chloral relative weak nucleophilic anions such as a chloride ion of tetraalkyl ammonium or tetraalkyl phosphonium salts are good initiators to form isotactic polymer.

Perhaloacetaldehydes could be prepared with a wide range of size and shape of the side groups and their
polymerization could be studied under a variety of relatively simple reaction conditions.

D. Bromoaldehyde polymers

Polybromal. Several attempts have been made in the past by us to polymerize bromal but without success. We have recently developed an improved method for the purification of bromal which could then be polymerized with pyridine as the initiator at —78°.

Earlier work on the polymerization of bromal using lithium tertiary butoxide and triphenylphosphine was unsuccessful. Traditionally, bromal as received from Eastman Kodak Company was carefully distilled under reduced pressure in a spinning band column. As the distillation proceeded, a faint brown color developed even in the "purest" samples. Triphenylphosphine reacts with bromal to form the triphenyl dibromovinyloxy phosphonium bromide. As a consequence, the initiator for the bromal polymerization was believed to be bromide, an anion which was known not to be very effective in chloral polymerization.

We have now developed conditions for a more sensitive gas chromotographic analysis of bromal and have found that three impurities are present in commercial bromal samples one or more of these impurities was the cause of the inability of the aldehyde to polymerize even when the sample was treated with a substantial amount of initiator. In connection with a different aspect of our work on the preparation of haloaldehydes, we found that treatment of bromal with SbF₃ gives bromal which after distillation is and remains colorless. Gas chromatographic analysis showed that two of the impurities have been completely removed by this treatment and that the third peak of the three impurities was reduced to less than 0.03%.

When the sample of purified bromal was treated with about 2 mol% of pyridine at —78° for 72 hr, a gel was obtained which was treated with methanol and gave a slightly yellow powder in 46% yield (Table 2).

\[
\text{CBr}_n\text{CHO} \rightarrow \left(\frac{\text{CBr}_n}{\text{CH} - \text{O}}\right)_{n-1}
\]

The polymer gave an i.r. spectrum which is very similar to the spectra of chloral and does not show any absorption in the carbonyl region (Fig. 11). Strong absorption in the regions of 800—1200 cm⁻¹ indicate the presence of ether linkages. Elemental analysis also indicate that polybromal was obtained. Calcd for \((\text{C}_2\text{H}_3\text{OBr}_3)\), Br, 85.38. Found: Br, 85.11. Pyridine is the only initiator which gave reliably polybromal with our new polymerization grade bromal. Other anionic and cationic initiators have not been effective until now.

Bromal copolymerized with phenyl isocyanates with pyridine as the initiator at —78°. The copolymer yields varied depending on actual copolymer composition but copolymers with up to 50 mol% of phenylisocyanate have been isolated (Table 4).

Polydibromochloroacetaldehyde. Dibromochloroacetaldehyde was prepared in a sequence of steps starting with monochloroacetaldehyde diethyl acetal. The compound was brominated and treated with sulfuric acid whereby crude dibromochloroacetaldehyde separated as a heavy brownish oil. After additional treatment with sulfuric acid the compound was distilled from P₂O₅. Very careful redistillation gave polymer grade dibromochloroacetaldehyde which according to GC analysis contained about 0.1% of impurities.

When initiated with pyridine or SbCl₅ at —45° a polymer was obtained which was worked up after 72 hr and yielded 50% of a slightly yellow insoluble powder (Table 2).

<table>
<thead>
<tr>
<th>Perhaloaldehyde</th>
<th>Initiator type</th>
<th>Polymerization Temperature (°C)</th>
<th>Polymer yield (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>BrCCl₂CHO</td>
<td>pyridine</td>
<td>—30</td>
<td>72</td>
</tr>
<tr>
<td>LiOC(CH₃)₃CHO</td>
<td>3</td>
<td>—30</td>
<td>80</td>
</tr>
<tr>
<td>SbCl₅</td>
<td>170</td>
<td>—30</td>
<td>58</td>
</tr>
<tr>
<td>CF₃SO₃H</td>
<td>170</td>
<td>—30</td>
<td>40</td>
</tr>
<tr>
<td>Br₂CCICHO</td>
<td>pyridine</td>
<td>—45</td>
<td>52</td>
</tr>
<tr>
<td>SbCl₅</td>
<td>72</td>
<td>—45</td>
<td>24</td>
</tr>
<tr>
<td>Br₃CCHO</td>
<td>pyridine</td>
<td>—78</td>
<td>46</td>
</tr>
</tbody>
</table>

Fig. 11. I.R. spectra of polyhaloaldehydes: (KBr pellets). (a) Polybromal; (b) Polydibromochloroacetaldehyde; (c) Polybromodichloroacetaldehyde.
Cationic and anionic aldehyde polymerization

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Br₂CCl—CHO was prepared from bromal and SbF₃. It was purified and polymerized with pyridine and gave an insoluble polymer whose infrared spectrum is in agreement with the proposed formula

\[
\text{Br}_2\text{C} \backslash \text{CHO} \rightarrow \left( \text{Br}_2\text{C} \backslash \text{CHO} \right)_{\infty}
\]  

Elemental analysis: Calcd for (C₂H₂OBr₂), C, 10.91; H, 0.45. Found: C, 11.06; H, 0.68.

E. Fluoroaldehyde polymers

Polydifluorochloroacetalddehyde. Polydifluorochloroacetalddehyde was prepared earlier and its polymerization described in a patent. We have worked out an effective way of preparing difluorochloroacetalddehyde by a sequence of steps starting with methyl difluoroacetate. Reduction with lithium aluminum hydride in ether at −78° gave the hydrate of difluorochloroacetaldhyde which was dehydrated with sulfuric acid and P₂O₅ and then polymerized.

\[
\text{CICF}_2\text{COOCH}_3 \overset{\text{LiAlH}_4}{\longrightarrow} \text{CICF}_2\text{CH(OH)}_2
\]  

\[
\text{CICF}_2\text{CH(OH)}_2 \overset{\text{H}_2\text{SO}_4}{\longrightarrow} \text{CICF}_2\text{CHO}
\]  

\[
\text{ClCIF}_2 \rightarrow \left( \text{ClCIF}_2 \right)_{\infty}
\]  

A number of anionic and cationic initiators proved to be effective for the polymerization of difluorochloroacetalddehyde, but lithium tertiary butoxide, triethylaluminum and triphenylphosphine were most effective (Table 3). The polymerization of difluorochloroacetalddehyde gave not only insoluble but also soluble polymer. Completely acetone soluble polymer was obtained with SbCl₅ and with triphenylphosphine under certain conditions. In most other cases, polydifluorochloroacetalddehyde is either insoluble or a mixture of insoluble and soluble polymer.

<table>
<thead>
<tr>
<th>Perhaloaldehyde</th>
<th>Initiator type</th>
<th>Time (hr)</th>
<th>Polymerization Temperature (°C)</th>
<th>Polymer yield (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CICClCHO</td>
<td>Ph₃P</td>
<td>1</td>
<td>25</td>
<td>55</td>
</tr>
<tr>
<td></td>
<td>AlEt₃</td>
<td>1</td>
<td>78</td>
<td>50</td>
</tr>
<tr>
<td></td>
<td>LiOC(CH₃)₂</td>
<td>1</td>
<td>78</td>
<td>91</td>
</tr>
<tr>
<td></td>
<td>SbCl₅</td>
<td>1</td>
<td>78</td>
<td>56</td>
</tr>
<tr>
<td>FCICClCHO</td>
<td>Ph₃P</td>
<td>120</td>
<td>−5</td>
<td>80</td>
</tr>
<tr>
<td></td>
<td>LiOC(CH₃)₂</td>
<td>360</td>
<td>25</td>
<td>86</td>
</tr>
<tr>
<td></td>
<td>pyridine</td>
<td>1</td>
<td>−78</td>
<td>12</td>
</tr>
<tr>
<td></td>
<td>SbCl₅</td>
<td>1</td>
<td>−78</td>
<td>79</td>
</tr>
<tr>
<td></td>
<td>H₂SO₄</td>
<td>46</td>
<td>25</td>
<td>82</td>
</tr>
</tbody>
</table>
Polyfluorodichloroacetaldehyde. Polyfluorodichloroacetaldehyde was prepared by a sequence of steps starting with methyl trichloroacetate, which was fluorinated with SbF₃ to methyl fluorodichloroacetate. Lithium starting with methyl trichloroacetate, which was fluorinated with phenyl isocyanate and chloral. Polyfluorodichloroacetaldehyde. A number of cationic initiators with anionic and cationic initiators to give insoluble polymers. A number of cationic initiators such as sulfuric acid, TiCl₅ and SbCl₅, gave good yields of polymer, but lithium tertiary butoxide and aluminum hydride reduction at −78° and hydrolysis of the product gave fluorodichloroacetaldehyde which was dehydrated with concentrated sulfuric acid and finally purified by distillation from P₂O₅.²⁶

Polyfluorodichloroacetaldehyde. Polyfluorodichloroacetaldehyde was prepared by a sequence of steps starting with methyl trichloroacetate, which was fluorinated with SbF₃ to methyl fluorodichloroacetate. Lithium aluminum hydride reduction at −78° and hydrolysis of the product gave fluorodichloroacetaldehyde which was dehydrated with concentrated sulfuric acid and finally purified by distillation from P₂O₅.²⁶

Fluorodichloroacetaldehyde could be copolymerized with phenyl isocyanate and chloral.

IV. FUTURE PROBLEMS IN ALDEHYDE POLYMERIZATION

Several problems in aldehyde polymerization are still awaiting further investigation. The general interest will depend on new developments and particularly on the introduction of additional commercial products. Much of the previous work on aldehyde polymerization was spurred by the commercialization of polyformaldehyde, the use of acetaldehyde for the development of monomer casting technique for chloral for the fabrication of novel nonflammable polymers. (i) Future developments in aldehyde polymerization are foreseen in the success of polymerization of aldehydes with long side chains. This polymerization depends very much on the structure of individual monomers in solution and suggests a possible liquid crystal behavior of such solutions. If the molecular weight of these aldehyde polymers with long side chains could be improved, polymers with novel mechanical properties which might be useful as compatibilization agents in polymer mixtures are envisioned. (ii) Because of their much improved autoxidation resistance over polymers with short side chains, these polyaldehydes might give additional understanding of the diffusion of oxygen in microphase separated systems. (iii) Very little is known about copolymers of aliphatic polyaldehydes and it would be desirable to prepare copolymers, in order to improve molecular weight and other properties. (iv) Cryotachensic polymerization of chloral has led to the development of sequential polymerization of chloral and additional polymers which is the only way to prepare blends and crosslinked networks of polychloral. (v) Direct and complete capping by termination reactions described in this paper could lead to chloral polymers of commercially useful stability. (vi) The crystal structure of all the new haloaldehyde polymers should be investigated and the potential energy calculations should be carried out. When successfully completed, we should have a full understanding of the influence of the size and shape of the side groups on the perhaloaacetaldheyde of the individual. (vii) Aliphatic aldehydes with functional groups will continue to be investigated in order to obtain materials with new properties.

REFERENCES


Table 4. Copolymerization of perhaloaldehydes with phenylisocyanate

<table>
<thead>
<tr>
<th>Perhaloaldehyde</th>
<th>Phenylisocyanate (mol%)</th>
<th>Polymerization</th>
<th>Initiator type</th>
<th>Copolymer yield (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CCl₃CHO</td>
<td>5—20</td>
<td>1</td>
<td>0</td>
<td>many</td>
</tr>
<tr>
<td>CCl₃BrCHO</td>
<td>50</td>
<td>5—20</td>
<td>0</td>
<td>pyridine</td>
</tr>
<tr>
<td>CBr₃CCHO</td>
<td>50</td>
<td>5—20</td>
<td>0</td>
<td>pyridine</td>
</tr>
<tr>
<td>CBr₂C₁CHO</td>
<td>50</td>
<td>5—20</td>
<td>0</td>
<td>pyridine</td>
</tr>
<tr>
<td>CCl₄FCHO</td>
<td>50</td>
<td>46</td>
<td>—5</td>
<td>Ph₃P</td>
</tr>
</tbody>
</table>
Cationic and anionic aldehyde polymerization