Preparation of Cellulose Esters with Aromatic Carboxylic Acids

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Abstract: The preparation of cellulose esters with aromatic carboxylic (substituted benzoic) acids in the presence of p-toluenesulfonyl chloride (TsCl) in pyridine (Py) was investigated. Cellulose was readily acylated with substituted benzoic acids, such as nitro-, chloro-, methyl- and methoxybenzoic acid, by use of the Py/ TsCl/ acid system, and corresponding cellulose esters with high degrees of substitution (DS) were obtained. The difference among the substituents and among the isomers (o, m- and p-isomers) of the acid did not influence the reaction significantly.

With an amino-substituted benzoic (p-aminobenzoic) acid, the cellulose ester could not be obtained, because acylation of the NH₂ group with the other p-aminobenzoic acid took place in preference to the OH group of cellulose. When the NH₂ group was protected with a methyl group, on the other hand, the corresponding ester was obtained by this method.

1. Introduction

It is well known that organic esters of cellulose can be prepared by the use of carboxylic acid anhydride with an acid catalyst, carboxylic acid chloride in the presence of a base and carboxylic acid in combination with trifluoroacetic anhydride [1, 2]. In any methods, either the acid anhydride or the acid chloride, i.e. an activated carboxylic acid is necessary.

In the field of organic cellulose esters, there has been increasing interest in new methods and routes for the preparation of esters with specific acids to provide new characteristics and functions to cellulose. A method for the preparation of cellulose esters directly with carboxylic acids, without using acid anhydrides or acid chlorides, will lead not only to the simplification of the preparation and isolation of the conventional acid derivatives but also the preparation of cellulose esters with specific carboxylic acids whose derivatives are difficult to obtain by conventional methods.

Brewster and Ciotti [3] have shown a method for the preparation of esters from alcohols and carboxylic acids with a pyridine (Py) solution of aromatic sulfonyl chloride. We have applied this method to the preparation of cellulose esters directly with carboxylic acids, and reported that this is generally applicable to the preparation of esters of higher aliphatic and benzoic acids [4] as well as cellulose acetate [5]. In addition, we have found that carboxylate salts can also be used for the esterification as same as carboxylic acids [6]. In this paper we describe the esterification with substituted benzoic acids by a similar system (Eq. 1).

\[
\text{Cell-OH} + \text{R-COOH} \underset{\text{Py/TsCl}}{\rightarrow} \text{Cell-O-C-OR}
\]

Recently, a variety of cellulose triesters are utilized as functional materials for chromatographic optical resolution [7]. This method seem to have a possibility to prepare new cellulose esters directly with specific acids.

2. Experiment

2.1 Cellulose Sample

Acetate grade pulp (a-cellulose 96.1%) was used as a cellulose sample. The sample was grounded in Wiley mill to less than 40 mesh before the pretreatment described below.

2.2 Pretreatment

Water pretreatment was employed to the cellulose to increase its reactivity. After being immersed in distilled water at room temperature for 1 h, the sample was
squeezed on a glass filter, rinsed completely with Py to remove the water and then squeezed again.

2.3 Acylation

The pretreated sample (1 g) was introduced into a solution of 7 g of p-toluenesulfonyl chloride (TsCl) in 30 g of Py. Aromatic carboxylic acid equimolar to TsCl was subsequently added into the solution. After the reaction for 2-20 h at 50 °C, the reaction mixture was poured into an excess amount of ethanol. The product was collected on a glass filter, washed with ethanol, Soxhlet-extracted with ethanol for 6 h, and dried at room temperature.

Infrared spectra (IR) of the products were recorded on a model A202 infrared spectrophotometer (Japan Spectroscopic Co. Ltd.) by the KBr disk technique. The reaction mixture was analyzed by 1H-NMR measurements (HITACHI R-90 FT NMR) using chloroform-d$_4$ containing tetramethylsilane as an internal reference.

3. Results and Discussion

The property of a substituted benzoic acid depends upon the nature and position of the substituent attached to the benzene ring. Table 1 lists the substituted benzoic acids used in this work and their acidity constants (Ka).

<table>
<thead>
<tr>
<th>Substituent</th>
<th>o-</th>
</tr>
</thead>
<tbody>
<tr>
<td>-CO$_2$H</td>
<td>1.32 × 10$^{-5}$</td>
</tr>
<tr>
<td>-CO$_2$OH</td>
<td>1.32 × 10$^{-5}$</td>
</tr>
</tbody>
</table>

Fig. 1 Rates of esterification with Py/ TsCl/ nitrobenzoic acids (o-, m- and p-isomer).

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Fig. 2 IR spectra of original pulp and cellulose esters prepared with o-, m- and p-nitrobenzoic acids.

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observed in the spectra. The spectrum of the product prepared with p-isomer agreed well with that of the product prepared by the acid chloride method. It is known that cellulose is tosylated and chlorinated by the treatment with TsCl in Py. However, there was no detectable sulfur and chlorine in the present products, and this shows that neither tosylation nor chlorination of the OH groups occurred on cellulose in this reaction. From these results the products were confirmed to be cellulose esters of corresponding nitrobenzoic acids with high DS values.

IR spectra of the products prepared with chloro-, methyl- and methoxy-benzoic acids by the procedure same as for nitrobenzoic acids were shown in Figs. 3, 4, and 5, respectively. These spectra also have absorptions characteristic of the ester and the aromatic ring similarly to those of the products prepared with nitrobenzoic acids. In addition, since the OH absorption became very weak, these products were also cellulose esters of the corresponding acids with high DS values.

The DS values of these products calculated from the elemental analysis data are summarized in Table 2. The small differences in DS values among isomers and among substituents show that cellulose esters with high DS values can be prepared from substituted benzoic acids independent of the substituents and isomers by the Py/TsCl/acid system. Yields of these products corresponded to those calculated from DS values.

The preparation of cellulose ester with p-aminobenzoic acid was attempted next for the introduction of an amino group to cellulose. The trial, however, was unsuccessful. The product obtained was not the ester but a mixture of unreacted pulp and the compound whose IR spectrum was shown in Fig. 6 (B). Since there are absorptions

![Fig. 3 IR spectra of cellulose esters prepared with o-, m- and p-chlorobenzoic acids.](image.png)

![Fig. 4 IR spectra of cellulose esters prepared with o-, m- and p-methylbenzoic (toluic) acids.](image.png)

![Fig. 5 IR spectra of cellulose esters prepared with o-, m- and p-methoxybenzoic acids.](image.png)

<table>
<thead>
<tr>
<th>Substituent</th>
<th>o-</th>
<th>m-</th>
<th>p-</th>
</tr>
</thead>
<tbody>
<tr>
<td>-NO₂</td>
<td>2.85</td>
<td>2.70</td>
<td>2.95</td>
</tr>
<tr>
<td>-Cl</td>
<td>2.90</td>
<td>2.90</td>
<td>2.80</td>
</tr>
<tr>
<td>-CH₃</td>
<td>2.40</td>
<td>2.60</td>
<td>2.50</td>
</tr>
<tr>
<td>-OCH₃</td>
<td>2.60</td>
<td>2.70</td>
<td>2.50</td>
</tr>
</tbody>
</table>

*Reaction Time: 20 h.*
characteristic of N-H stretching (3300 cm⁻¹), amide C=O stretching (1650 cm⁻¹; amide I absorption) and N-H bending (1500 cm⁻¹; amide II absorption) in the spectrum, the compound seems to be an amide compound that is formed by condensation of p-aminobenzoic acid. This indicates that the acylation of the NH₂ groups took place in preference to the OH groups of cellulose in the Py/ TsCl/ p-aminobenzoic acid system. Fig. 6 (C) shows the IR spectrum of the product that was prepared with p-(N,N-dimethylamino) benzoic acid instead of p-aminobenzoic acid in the system. The absorptions due to ester group, C=O(1720 cm⁻¹) and C-O (1260 cm⁻¹) stretchings, show that the product is a cellulose ester. The DS values calculated from the elemental analysis data was 1.10, lower than those of the esters described previously. These findings show that when the NH₂ group of an amino benzoic acid was protected, the corresponding cellulose ester could be obtained with the Py/ TsCl/ acid system.

Brewster and Ciotti (3) have shown that carboxylic acids react rapidly with arylsulfonyl chloride (ArSO₂Cl) in a Py solution to form symmetrical anhydrides, and that the anhydrides serve as acylating reagents of alcohols (Eq. 2).

\[
\text{Py/ ArSO₂Cl} \rightarrow \text{R-CO-O-CO-R} \quad \text{(2)}
\]

Relating to this, ¹H-NMR spectra were taken to examine the composition of the present system. Fig. 7 show ¹H-NMR spectra (2-3 ppm) of Py/ benzenesulfonyl chloride (BsCl)/ p-methylbenzoic (toluic) acid system which did not include the cellulose sample. BsCl was used as a ArSO₂Cl instead of TsCl because the methyl proton peak of TsCl overlapped with that of toluic acid in the spectra. The peak at 2.38 was assigned to the methyl protons of p-toluic acid based on its ¹H-NMR spectrum in Py solution. We have confirmed the formation of acetic anhydride in the Py/ TsCl/ AcOH system (9) by the ¹H-NMR measurement. Brewster and Ciotti (3) have prepared benzoic anhydride by the Py/ BsCl/ benzoic acid. The peak at 2.42 ppm, therefore, is considered to be due to the methyl protons of p-toluic anhydride. The anhydride was formed just after the mixing of Py, BsCl and the acid (Fig. 7 (A)), and increased with time while the acid decreased. After standing for 1 h at room temperature, the acid was mostly converted to the anhydride (Fig. 7 (C)), which were stable in the system thereafter. The anhydride thus formed in situ will react with cellulose, leading to the ester.

Solubility of the esters obtaind in this work was in-
Table 3  Solubility of Cellulose Esters\(^{a)}\) in Various Organic Solvents

<table>
<thead>
<tr>
<th>Solvent</th>
<th>(-\text{NO}_2)</th>
<th>(-\text{Cl})</th>
<th>(-\text{CH}_3)</th>
<th>(-\text{OCH}_3)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>o- m- p-</td>
<td>o- m- p-</td>
<td>o- m- p-</td>
<td>o- m- p-</td>
</tr>
<tr>
<td>DMSO</td>
<td>(\triangle)</td>
<td>(\triangle)</td>
<td>(\times)</td>
<td>(\times)</td>
</tr>
<tr>
<td>DMF</td>
<td>(\triangle)</td>
<td>(\triangle)</td>
<td>(\times)</td>
<td>(\times)</td>
</tr>
<tr>
<td>Acetone</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\times)</td>
</tr>
<tr>
<td>Ethyl acetate</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\times)</td>
</tr>
<tr>
<td>Dichloroethane</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\triangle)</td>
<td>(\triangle)</td>
</tr>
<tr>
<td>THF</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\triangle)</td>
<td>(\triangle)</td>
</tr>
<tr>
<td>Chloroform</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\triangle)</td>
</tr>
<tr>
<td>Dichloromethane</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\triangle)</td>
<td>(\triangle)</td>
</tr>
<tr>
<td>Ethyl ether</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\times)</td>
</tr>
<tr>
<td>Pentane</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\times)</td>
</tr>
<tr>
<td>Benzene</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\times)</td>
<td>(\times)</td>
</tr>
</tbody>
</table>

\(^{a)}\)  DS values are shown in Table 2.

\(^{b)}\)  \(\triangle\), Soluble; \(\times\), swelling and/or partly soluble; \(\times\), insoluble.

acknowledgments

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References