Allyltrimethoxysilane as the Reagent of Double Bond Introduction to Porous Silica for Preparation of a Chiral Stationary Phase in Comparison with Allyltriethoxysilane

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Chiral separation has been extensively used not only to produce pure enantiomers but also to analyze mixtures of enantiomers. Many different types of chiral stationary phases have been developed such as brush or ligand-exchange type phases, cyclodextrin phases, protein phases, polysaccharide phases, etc. Optically active polymers compose a great portion of the chiral phases and have been thoroughly investigated in a recent review.1 Among them, polysaccharide chiral phases have been known to be very useful for their versatility in chiral separation.2 Derivatives of polysaccharide chiral phases are now very popular chiral stationary phases and are commercially available, but they are not compatible with some common LC solvents such as tetrahydrofuran and chloroform for their solubility in the solvents.3 Such a problem can be solved by securing chemical bonding between the silica surface and the polysaccharide ligand.4,5 Such phases (Diacel) have been commercially available only very recently. Introduction of double bonds to both the silica surface and the polysaccharide ligand followed by cross polymerization between the silica and the ligand has been the method of securing chemical bonding. The surface silanol groups of silica are used in the reaction of introducing double bonds, and two different reagents have been used to incorporate double bonds: 3-(trimethoxysilyl) propyl methacrylate6 and allyltrimethoxysilane.6,7 Improvements in double bond introduction to the silica surface and consequent cross polymerization seem to be important for commercial polysaccharide chiral phases with chemically bonded polysaccharide ligands.

In this study, we have found that an improvement of the process can be achieved by simply employing allyltrimethoxysilane instead of allyltriethoxysilane (traditional reagent) as the reagent of double bond introduction.

Experimental Section

Chemicals. Heptane and 2-propanol were of HPLC grade and purchased from Fisher (Pittsburgh, PA, USA) and used without purification. Allyltrimethoxysilane, allyltriethoxysilane, trans-stilbene oxide, benzoin, warfarin, 10-undecenoyl chloride, 3,5-dimethylphenylisocyanate and HMDS (1,1,1,3,3,3-hexamethyldisilazane) were purchased from Aldrich (Milwaukee, IL, USA). AIBN (2,2'-azobisisobutyro-

Figure 1. The overall schematic of synthesis of the chiral stationary phase by cross polymerization of allylsilica with a cellulose derivative with double bonds.
nitrile) was a gift from a neighboring laboratory. Microcrystalline cellulose was purchased from Merck (Whitehouse Station, NJ, USA). Silica gel (Adsorbosphere, 5 μ, 80 Å) was obtained from Alltech (Deerfield, IL, USA).

**HPLC.** Two Shimadzu (Tokyo, Japan) 10AD pumps, a Shimadzu DGU-14A membrane degasser, a Rheodyne (Cotati, CA, USA) 7520 injector with a 0.5 μL injection loop, an Isco (Lincoln, NE, USA) CV4 capillary window detector, and the home-made 0.5 mm ID glass-lined micro-column were combined to compose the HPLC system. The chromatographic data were obtained by a PC system, and a software MultiChro 2000 from Youlin-Gisul (Sungnam, Korea) was used to acquire and process the data. An Alltech (Deerfield, USA) slurry packer was used to pack the micro-columns used.

**Synthesis of chiral stationary phase.** The chiral stationary phases were prepared by cross polymerization of allylsilica with a cellulose derivative having double bonds (Figure 1).

- The cellulose derivative was obtained by the process in the literature with some modifications. Microcrystalline cellulose 1 g, 10-undecenoyl chloride 1 g were dissolved in pyridine and stirred for 1 hr at 100 °C, then 1 mL 3,5-dimethylphenyl isocyanate was added and stirred for 24 hr at 100 °C. Upon addition of methanol to the reaction mixture, the crude product was precipitated. The precipitate was dissolved in chloroform and recrystallized by methanol to give pure 10-undecenoyl-cellulose(3,5-dimethylcarbamate).

- Porous allylsilica was prepared as follows. The porous silica (5 μ, 80 Å) was baked at 300 °C for 3 hr to remove adsorbed water completely. The porous silica 1 g was dispersed in toluene and reacted with 0.4 mL HMDS was added for endcapping and the reaction mixture was stirred for 1 hr at the same temperature. After completion of reaction, the silica was filtered, thoroughly washed with toluene, THF, 50/50 (v/v %) methanol/water, and methanol, and dried. Porous allylsilica was also prepared by the same procedure described above but using allyltrimethoxysilane instead of allyltri-methoxy-silane for comparison purpose.

- The allylsilica 1 g was dispersed in THF, and 0.3 g cellulose derivative and 0.015 g AIBN were added. The reaction mixture was evaporated in a rotavapor at room temperature. Cross polymerization was carried out at 100 °C for 4 hr while the bottle was rotated in the rotavapor. The final product was thoroughly washed with chloroform under reflux and acetone in series and dried to yield the chiral stationary phase.

- The stationary phase (made by allyltrimethoxysilane vs. allyltriethoxysilane) was dispersed in methanol and packed in the microcolumn.

**Results and Discussion**

The results of chiral separation of R- and S-enantiomers of trans-stilbene oxide, benzoin, and warfarin by the column (0.5 × 300 mm) packed with the chiral phase made by the conventional reagent (allyltrimethoxysilane) are compared with the results obtained by the column packed with the chiral phase made by allyltrimethoxysilane in Figure 2 (for 90/10 v/v % heptane/2-propanol), Figure 3 (for 98/2 v/v % heptane/2-propanol), and Table 1. The retention data of warfarin in 98/2 (v/v %) heptane/2-propanol were not obtained since warfarin enantiomers retained too long in that eluent. The test samples were prepared by dissolving the solutes in pure 2-propanol since solubilities of the solutes in the eluents are limited. This discrepancy between the sample solvent and the eluents caused some strange looking system peaks in front of the main peaks of R- and S-enantiomers, and they were confirmed by injection of 2-propanol and there was no confusion in identifying the main peaks.

We can easily note that the selectivities and resolutions obtained with the chiral phase made by allyltrimethoxy-
Silane were better than those obtained with the chiral phase made by allyltriethoxy silane although the extent of improvement was variant depending upon the solute types and elution conditions. This improvement in separation efficiency of allyltrimethoxy silane over allyltriethoxy silane was even more evident when a very short column was made for both cases and compared (Figure 4). We had to use a rather fast flow rate (50 \( \mu \)L/min) for the microcolumn to observe the peaks of warfarin enantiomers since the retention times of warfarin isomers were too long even with the short column (5 cm) in 98/2 (v/v %) heptane/2-propanol. Anyway, in all cases, only one peak was observed for the chiral phase made by allyltrimethoxy silane while two peaks (R- and S-enantiomers) were observed for the chiral phase made by allyltriethoxy silane.

Table 1. Comparison of the selectivities (\( \alpha \)) and resolutions (Rs) in separation of R- and S-enantiomers of trans-stilbene oxide, benzoin, and warfarin by the chiral column (0.5 \( \times \) 300 mm) a) and b) The column packed with the chiral phase made by the conventional reagent (allyltriethoxy silane). b) The column packed with the chiral phase made by allyltrimethoxy silane

<table>
<thead>
<tr>
<th>Elution condition</th>
<th>trans-stilbene oxide</th>
<th>benzoin</th>
<th>warfarin</th>
</tr>
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<tbody>
<tr>
<td>90/10 (v/v %)</td>
<td>Column a) 1.44  0.75</td>
<td>1.30  1.18</td>
<td>1.97  2.63</td>
</tr>
<tr>
<td>heptane/2-propanol Column b) 1.52  1.85</td>
<td>1.36  1.99</td>
<td>2.40  3.72</td>
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</tr>
<tr>
<td>98/2 (v/v %)</td>
<td>Column a) 1.66  2.06</td>
<td>1.78  2.64</td>
<td>– –</td>
</tr>
<tr>
<td>heptane/2-propanol Column b) 2.21  3.35</td>
<td>1.79  2.72</td>
<td>– –</td>
<td></td>
</tr>
</tbody>
</table>

Figure 3. Comparison of Separation of R- and S-enantiomers of trans-stilbene oxide and benzoin in 98/2 (v/v %) heptane/2-propanol at a flow rate of 10 \( \mu \)L/min by the column (0.5 \( \times \) 300 mm) packed with a) the chiral phase made by the conventional reagent (allyltriethoxy silane) and b) the chiral phase made by allyltrimethoxy silane. The detection wavelength was 254 nm.

Figure 4. Comparison of Separation of R- and S-enantiomers of trans-stilbene oxide, benzoin, and warfarin in 98/2 (v/v %) heptane/2-propanol by the short column (0.5 \( \times \) 30 mm) packed with a) the chiral phase made by the conventional reagent (allyltriethoxy silane) and b) the chiral phase made by allyltrimethoxy silane. The flow rate was 10 \( \mu \)L/min for trans-stilbene oxide and benzoin, and 50 \( \mu \)L/min for warfarin. The detection wavelength was 254 nm.
allyltrimethoxysilane.

Such superiority of the phase made with allyltrimethoxysilane over that made with allyltriethoxysilane was clear when the results of elemental analysis of both phases were compared (Table 2). We can note that more allyl groups were chemically attached to the silica by allyltrimethoxysilane than by allyltriethoxysilane and that consequently more polysaccharide ligands were chemically bound to the silica by cross polymerization. The retention times and separation efficiencies of a pair of enantiomers tend to improve as the content of the chiral ligand (carbon load) bound to the porous silica surface increases unless the ligand amount is larger than the limit over which the separation efficiency rather degrades because of too slow mass transfer rate. Actually, we observed longer retention times and better separation efficiencies with the chiral phase made by allyltrimethoxysilane than with the chiral phase made by allyltriethoxysilane (Figures 2-4).

The reason why an improved carbon load was observed when allyltrimethoxysilane is used instead of allyltriethoxysilane can be explained as follows. Methanol is supposed to be evolved when allyltrimethoxysilane reacts with a silanol group while ethanol should be evolved when allyltriethoxysilane is used. The reaction will proceed more readily if the reaction product (methanol or ethanol) is removed more easily. The boiling point of methanol is lower than that of ethanol and should be removed from the reaction mixture more readily. Furthermore, more bulky ethoxy groups will impose higher steric hindrance when an allyltriethoxysilane molecule approaches to a silanol group, which causes less reactivity of allyltriethoxysilane than allyltrimethoxysilane.

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References