

Supercritical Fluid Chromatographic Separation of Polyethylene Glycol Polymer

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The use of compressed (dense) gases and supercritical fluids as chromatographic mobile phases in conjunction with liquid chromatographic (LC)-type packed columns was first reported by Klesper *et al.* in 1962.¹ During its relatively short history, supercritical fluid chromatography (SFC) has become an attractive alternative to GC and LC in certain industrially important applications. SFC gives the advantage of high efficiency and allows the analysis of non-volatile or thermally labile mixtures.

The density of the mobile phase in SFC is about 200-500 times that in gas chromatography. The effect of shorter intermolecular distances and the resulting increase in molecular interactions is an enhanced solubilizing capability of the solvent towards various solutes. Compounds with much higher molecular weights than gas chromatography normally allows can therefore be chromatographed. However, the most commonly used mobile phases in SFC are all relatively non-polar fluids. Carbon dioxide, the most widely used fluid, is no more polar than hexane,² even at high densities. Solute polarity should be between that of the stationary phase and the mobile phase in order to have a well behaved separation. Few real samples contain only non-polar solutes, so a major objective of research in SFC has been directed toward increasing the range of solute polarity that can be handled by the technique. To bring the SFC technique into routine use, mobile phases that are more polar than the commonly used carbon dioxide are necessary.

The solvent strength of supercritical CO₂, even at high density, is not sufficient for the elution of polar solutes. Polar mobile phase such as NH₃⁴ exhibits useful properties, but a more practical way to extend the range of compounds separable by SFC is to use a mixed mobile phase. The solubility of the solute in the supercritical phase can be influenced considerably by adding modifiers to the mobile phase. The use of modifiers has been reported by Jentoft and Gouw³ and by Novotny *et al.*⁴ The latter group showed that adding 0.1% 2-propanol to pentane as the mobile phase decreases the observed partition coefficient (K) values for many polynuclear aromatic hydrocarbons by 20-35%. Thus, the addition of modifiers (generally organic solvents) to a supercritical mobile phase changes the polarity of the mobile phase and also leads to deactivation of the column packing material. In capillary SFC, most separations are made with pure CO₂ because of its compatibility with an flame ionization detection (FID); except for formic acid and water, the addition of any common modifier precludes the use of

FID.⁵ Modifiers are essential in packed-column SFC for the elution of polar compounds⁶ and are extensively used.

Several workers⁶⁻⁸ have reported the influence of modifiers on peak shape, selectivity and retention time in capillary and packed-column SFC. A simple and effective way to add modifiers to a supercritical fluid mobile phase is to use a saturator column,^{9,10} which is usually a silica column saturated with polar solvents.

In this work, water was used as a polar modifier and a μ -Porasil column as a saturator column. The μ -Porasil column was inserted between the pump outlet and the injection valve. During the passage of the supercritical fluid mobile phase through the silica column, a polar modifier (water) can be dissolved in the pressurized supercritical fluid. Polyethylene glycol polymer is relatively polar polymer since it has a polar functional group (-OH). Polyethylene glycol polymer has never been separated using water modified mobile phase. In this paper, using a μ -Porasil column as a saturator column, excellent supercritical fluid chromatograms of polyethylene glycol oligomers were obtained.

Experiments

A CCS (Computer Chemical System, Avondale, PA) Model 5000 supercritical fluid chromatograph was used with a 100 mm \times 2 mm i.d. packed column (Nucleosil diol). This system was equipped with a C14W loop injector (Valco) and a flame ionization detector. SFC-grade carbon dioxide (Scott Specialty Gases) was used as a basic mobile phase. Experimental conditions for at 150 °C, pressure programmed from 1500 to 3500 psi at 25 psi min⁻¹, detector at 300 °C and 5 mL min⁻¹ restrictor flow-rate at 1500 psi. For the addition of modifiers to supercritical CO₂, at μ -Porasil column (250 mm \times 4.6 mm i.d.) which is manufactured for normal-phase HPLC by Waters was used. Its functional group is a silanol (SiOH) group. The μ -Porasil column was saturated with modifiers using a Model 600 syringe pump (Lee Scientific) and placed between the pump and injector. Figure 1 shows the schematic diagram of supercritical fluid chromatography system used for the separation of polyethylene glycol polymer.

Results and Discussion

When modifiers are used with supercritical CO₂ in order to chromatograph more polar substances, the binary mixture

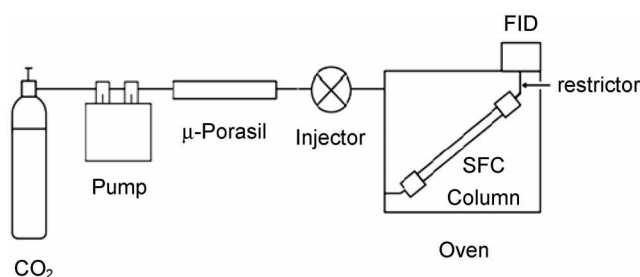


Figure 1. Schematic diagram of supercritical fluid chromatography system used for the separation of polyethylene glycol polymer.

of eluents can contaminate the instrument. Especially when water or formic acid is used as a modifier, the modifier remaining in a pump may cause corrosion of the pump, and when methanol is used as a modifier, methanol remaining in the pump can be eluted slowly during the next run. This may affect the time required to achieve chemical equilibrium for the subsequent separations, and also many modifiers can evaporate and contaminate the air in the laboratory. A good way to overcome these problems is to use a saturator column^{9,10} to add polar modifiers to supercritical CO₂. For the experiment, water was used as a modifier and a μ -Porasil column was used as a saturator column with system design similar to that described by previous work.⁹ With this design, a polar modifier (water) can be added to pressurized CO₂ after the pump, and thus no modifier remains in the pump. Supercritical CO₂ is delivered from the pump to the μ -Porasil column which is saturated with water. When supercritical CO₂ passes through the μ -Porasil column, water molecules held on the hydroxy groups of the μ -Porasil by hydrogen bonding can dissolve in the pressurized supercritical fluid. Thus non-polar supercritical CO₂ can have the characteristics of a polar mobile phase because it can absorb a polar solvent, *i.e.*, water. Therefore, after passing through the μ -Porasil column, supercritical CO₂ is changed into a new mobile phase with different polarity, and it is possible to separate polar samples using this modified mobile phase. Polyethylene glycol oligomers were analysed with this modified mobile phase (supercritical carbon dioxide-water phase) using a μ -Porasil column as saturator

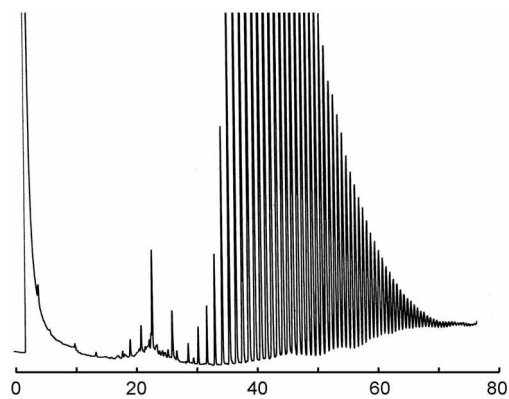


Figure 2. Chromatogram of oligomers of polyethylene glycol (No. 1) using a μ -Porasil column.

column. The chromatograms are shown in Figure 2 and Figure 3. Excellent separations were obtained. These kinds of separations are almost impossible with other chromatographic techniques. Figure 4 and Figure 5 show the separation of polyethylene glycol oligomers with pure carbon dioxide as the mobile phase. As expected, when a μ -Porasil column was used, more peaks were separated. These results agree well with those reported in the case of dimethylpolysiloxane.⁹ In the separations of dimethylpolysiloxane, when water was added on non-polar mobile phase, more

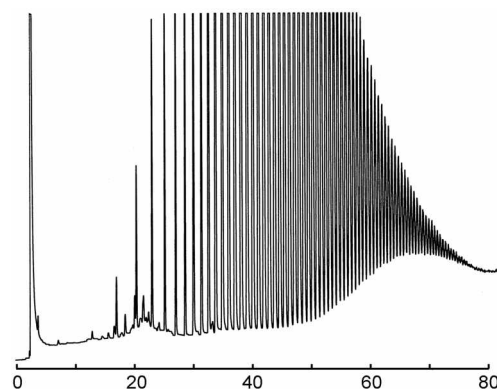


Figure 3. Chromatogram of oligomers of polyethylene glycol (No. 2) using a μ -Porasil column.

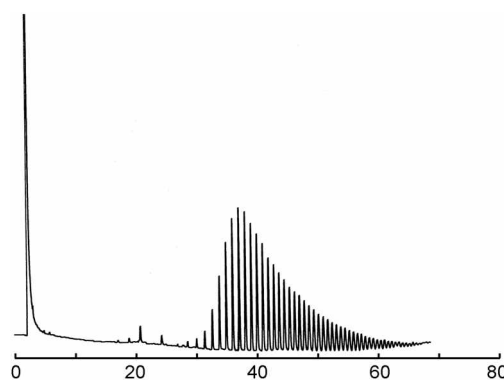


Figure 4. Chromatogram of oligomers of polyethylene glycol (No. 1) without using a μ -Porasil column.

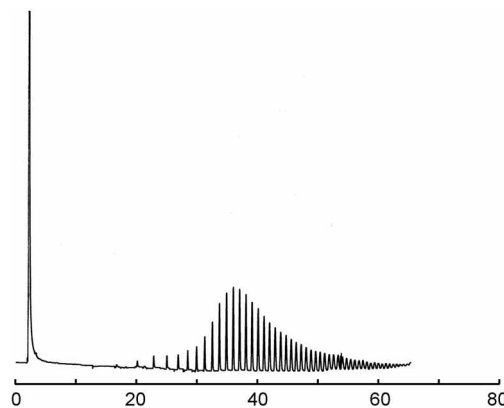


Figure 5. Chromatogram of oligomers of polyethylene glycol (No. 2) without using a μ -Porasil column.

oligomers were eluted. This may be due to the polarity of polyethylene glycol polymer.

Conclusion

The excellent supercritical fluid chromatograms were obtained for the separations of polyethylene glycol polymers. When small amount of water was added on supercritical carbon dioxide fluid, more oligomer peaks were observed. After passing through the μ -Porasil column, supercritical carbon dioxide is changed into a new mobile phase with different polarity, and it was possible to elute more oligomers using this modified mobile phase.

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