

Separation of statistic MMA-MAA Copolymers using Gradient SEC

Application Note Pharmaceutical Analysis

Author

Dr. Wolfgang Radke contact: <u>WRadke@pss-polymer.com</u>

Isocratic GPC/SEC is a powerful tool to separate macromolecules based on their hydrodynamic volume. In case of homopolymers GPC/SEC allows the fast, precise and easy determination of the complete molar mass distribution.

Unfortunately many modern polymeric materials are copolymers and isocratic GPC/SEC does not provide any information about the chemical composition. Here SEC-gradients or polymer HPLC is the method of choice.

Introduction

Statistic copolymers of methyl methacrylate (MMA) and methacrylic acid (MAA) are widely used in pharmaceutical applications. Besides the molar mass distribution the chemical composition and the amount of comonomers in the copolymer is of importance. Separations by conventional gradient HPLC failed, since the polar eluents required to dissolve polymers of high acid content prevent adsorption onto the stationary phase, resulting in pronounced breakthrough peaks. The problem can be avoided applying SEC-gradients, resulting in the desired separation according to the amount of methacrylic acid. The system can be calibrated using reference materials of known composition. This allows determining the average copolymer composition as well as the compositional heterogeneity.

System Requirements

	Conditions
Pump	PSS SECcurity GPC1260 binary pump • flow rate [mL/min]: 1.0 • mobile phase: Gradient: Chloroform/DMAc
Injection system	PSS SECcurity GPC1260 Autosampler Injection interval: 32 min
Columns	 PSS PROTEEMA precolumn (8*50 mm) PSS PROTEEMA, 3 μm, 100 Å (8x300 mm) Temperature: 60°C
Calibration	PSS MMA-MAA copolymers of different acid content (MAA: 9%, 25%, 31%, 42%, 48%wt)
Loading	Samples dissolved in DMAc 1 mg/mL, 100 µL injection volume
Detector	 PSS SECcurity ELS1000 Gas flow: 1.5 SL/min Nebulizer temperature: 100 °C Evaporator temperature: 200 °C.
Software	PSS WinGPC UniChrom with ChromPilot and Chemical Heterogeneity module





PSS Polymer Standards Service GmbH In der Dalheimer Wiese 5 55120 Mainz | Germany
 Phone
 +49 6131 96239-0

 Fax
 +49 6131 96239-11

 E-Mail
 info@pss-polymer.com

 Web
 www.pss-polymer.com

Polymer Standards Service-USA, Inc. 160 Old Farm Rd, Suite A Amherst | MA 01002 | USA
 Phone
 +1 413 835-0265

 Fax
 +1 413 835-0354

 E-Mail
 usa@pss-polymer.com

 Web
 www.pss-polymer.com



Procedure, Results & Discussion

The experimental procedure of SEC-gradients differs from conventional gradient HPLC. While in conventional HPLC the sample is dissolved in a weak eluent and injected at adsorbing conditions, the sample is dissolved in a strong mobile phase an injected at the end of the SEC-gradient. In the present application the copolymers were dissolved in DMAc.

Gradient details:

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- 0-3 min: 50%DMAc
- 3-8 min: 100%DMAc
- 8-23 min: 100%CHCl₃
- 23-26 min: 5%DMAc
- 26-32 min: linear increase from 5% to 50% DMAc.

Figure 1 shows the separation of 5 statistical MMA-MAA copolymers with MAA contents between 9 and 48%wt using a Chloroform/DMAc gradient from 5% to 50% DMAc. The sample components elute before the injected solvent and the different samples are clearly separated.

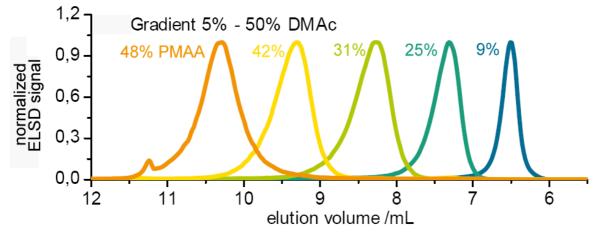
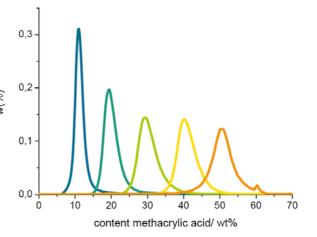
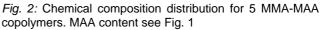


Fig. 1: Separation of 5 MMA-MAA copolymers with different PMMA content using a Chloroform - DMAc gradient on PSS PROTEEMA columns

A nearly linear dependence between MAA content and elution volume is observed. The use of this calibration curve allows the determination of the chemical composition distribution. Figure 2 shows the chemical composition distribution for the 5 samples with the PMAA content between 48% and $\frac{3}{5}$ with the PMAA content between 48% and $\frac{3}{5}$. It is observed that an increase in MAA content results in a significantly broader chemical composition distribution (CCD).







PSS Polymer Standards Service GmbH In der Dalheimer Wiese 5 55120 Mainz | Germany
 Phone
 +49 6131 96239-0

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 www.pss-polymer.com

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 Phone
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