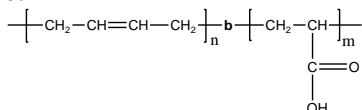


Sample Name:

Poly(1,4-butadiene)-b-poly(acrylic acid)

Sample#: **P43630A-BdAA**

Structure:



Composition:

Mn x 10 ³ PBd-b-AA	PDI
150.0-b-14.0	1.05
90% 1, 4 rich addition	

Synthesis Procedure:

Poly(1,4-butadiene -b- acrylic acid) is prepared by living anionic polymerization with sequence addition of butadiene followed by t-butyl acrylate and hydrolysis of the t-butyl group. The solvents for the polymerization selected to get the polybutadiene with microstructure rich in 1,4 addition or 1,2 addition.

Characterization:

An aliquot of the anionic poly(butadiene) block was terminated before addition of t-butyl acrylate and analyzed by size exclusion chromatography (SEC) to obtain the molecular weight and polydispersity index (PDI). The final block copolymer composition was calculated from ¹H-NMR spectroscopy by comparing the peak area of the vinylic butadiene protons between about 5.0-5.4 ppm with the t-butyl acrylate protons at 1.43 ppm. Block copolymer PDI is determined by SEC. Note: The ¹H-NMR of 1,2-polybutadiene is composed of 1 proton signal at 5.4 ppm and 2 proton signals at 5.0 ppm. Signals due to vinylic 1,4-polybutadiene are present at 5.4 ppm.

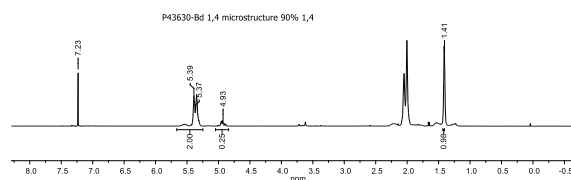
Hydrolysis of the ester was followed by FTIR for the disappearance of ter-butyl ester at 1362cm⁻¹.

Purification of the polymer:

After the Hydrolysis the solvent was removed under vacuum and the obtained polymer was dissolved in THF and neutralized with NaHCO₃ to get pH around 6. The product was filtered, and the filtrate was treated for 2days with Mixed

bed exchange resin, IONA NM -60 H⁺/OH⁻ form (16-50 mesh). The product was filtered and the clear solution was passed through a column packed with basic Al₂O₃. The Filtrate was concentrated under vacuum and dried at room temperature.

¹H-NMR Spectrum of the block copolymer:



SEC elugram of the BdtBuA copolymer:

