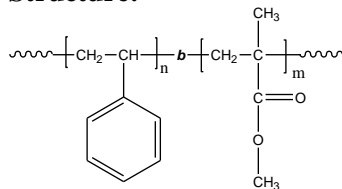


**Sample Name:** Poly(styrene-*b*-methyl methacrylate)  
(*polymethylmethacrylate rich in syndiotactic contents*  
> 78%)

**Sample #:** P9912PP-SMMA

**Structure:**



**Composition:**

Mn x 10 <sup>3</sup> S- <i>b</i> -MMA	PDI
537.0- <i>b</i> -265.0	1.05

T <sub>g</sub> for PS block: 106°C	T <sub>g</sub> for PMMA block: 131°C
Syndio:Hetero:Iso	81:19:0

**Synthesis Procedure:**

Poly(styrene-*b*-methyl methacrylate) is prepared by living anionic polymerization in THF at –78 oC using sec.BuLi initiator in the presence of LiCl. Polystyrene macroanions were end capped with a unit of diphenyl ethylene (DPE) before adding methylmethacrylate (MMA) monomer. For further details please see our published articles.1-5

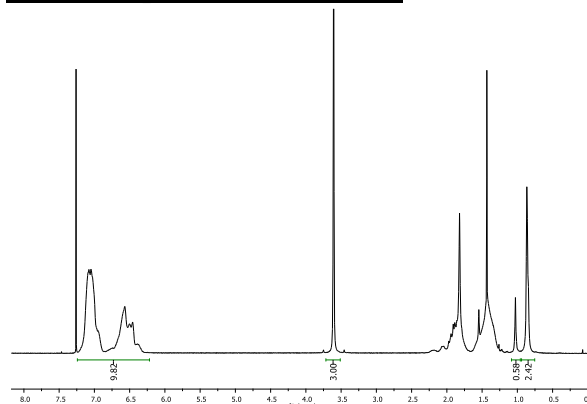
**Characterization:**

The final block copolymer composition was calculated from <sup>1</sup>H-NMR spectroscopy by comparing the peak area of the poly(methyl methacrylate) protons (eg. –OCH<sub>3</sub> at 3.6ppm) with the of aromatic protons of polystyrene at 6.3-7.2 ppm. The molecular weight and polydispersity of copolymer are determined by SEC. Thermal analysis of the samples was carried out using a differential scanning calorimeter (TA Q100) at a heating rate of 10°C/min. The inflection glass transition temperature (T<sub>g</sub>) of the sample has been considered.

**Solubility:**

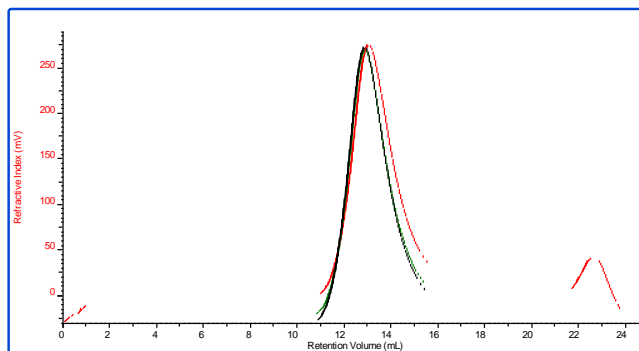
Poly(styrene-*b*-methyl methacrylate) is soluble in THF, toluene, dioxane and CHCl<sub>3</sub>. This polymer readily precipitates from methanol, ethanol, hexanes and water.

**<sup>1</sup>H-NMR Spectrum of Polymer:**



**SEC elugram of the Sample:**  
P9912P

dn/dc	0.1310
Flow Rate	0.7000
Solvent	DMF with LiBr
Method	PSS column-PMMA60K-Jan3-2019-0004.vcm



Sample	Mn	Mw	Mp	Mw/Mn
P9912P_1_2019-06-11	802,196	845,547	819,385	1.054

**Thermogram for the sample**

