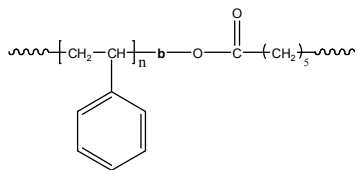


Sample Name: Poly(styrene-b- ϵ -caprolactone)

Sample #: P2037CL

Structure:



Composition:

$M_n \times 10^3$ S-b-CL	M_w/M_n (PDI)
9.50-25.0	1.80

Synthesis Procedure:

Poly(styrene-b- ϵ -caprolactone) is prepared by anionic polymerization with sequence addition of styrene followed by n-butyl methacrylate.

Characterization:

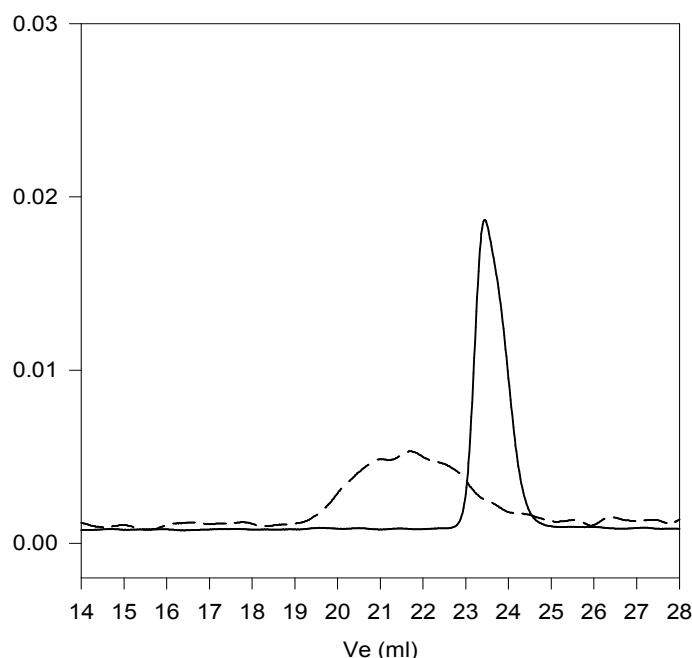
An aliquot of the polystyrene block was terminated before addition of - ϵ -caprolactone and analyzed by size exclusion chromatography (SEC) to obtain the molecular weight and polydispersity index (PDI). The final block copolymer composition was calculated from $^1\text{H-NMR}$ spectroscopy by comparing the peak area of the styrene protons at 6.3-7.2 ppm with the peak area of - ϵ -caprolactone protons at 4.1 ppm. Block copolymer PDI is determined by SEC.

Solubility:

Poly(styrene-b- ϵ -caprolactone) is soluble in THF, Chloroform, DMF, and precipitated in methanol and hexanes.

SEC profile of the block copolymer:

P2037-SCL



— SEC profile of Poly(Styrene-b- ϵ -caprolactone):

— Polystyrene, $M_n=9500$, $M_w=10200$, $PI=1.07$

- - Block Copolymer PS(9,500)-b-P ϵ CL(25,000), $PI=1.8$

Thermal analysis of the sample# P2037SCL

Thermal analysis of the samples was carried out on a TA Q100 differential scanning calorimeter at a heating rate of 10°C/min. The midpoint of the slope change of the heat flow plot of the second heating scan was considered as the glass transition temperature (T_g).

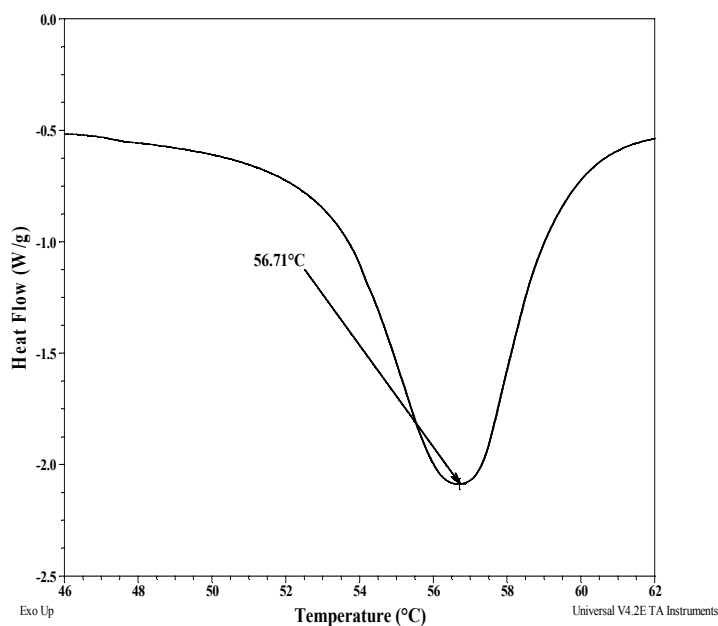
Melting and crystallization curve for the sample

The melting temperature (T_m) was taken as the maximum of the endothermic peak where as the crystallization temperature (T_c) was considered as the minimum of the exothermic peak.

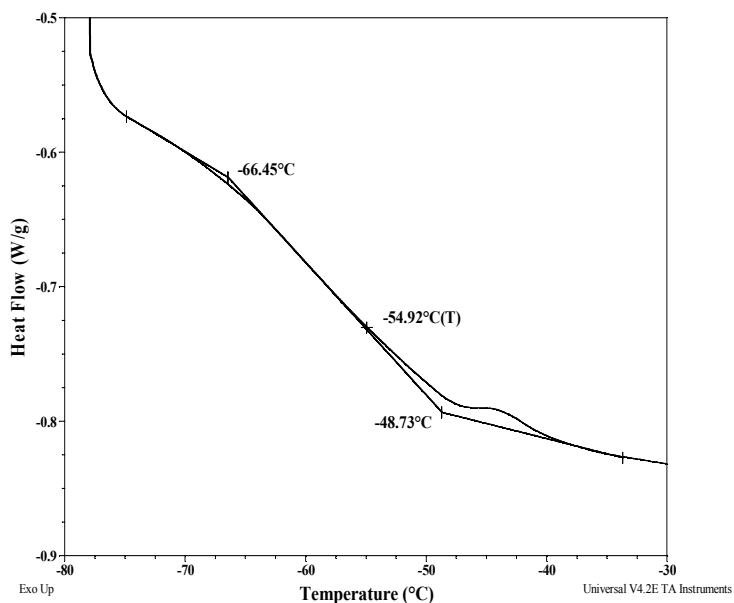
Thermal analysis results at a glance

Sample	T_m (°C)	T_c (°C)	T_g (°C)
ϵ -CL	57	19	-55
PS	-	-	Not distinct

Melting curve for ϵ -caprolactone block:



Thermogram for ϵ -caprolactone block:



Crystallization curve for CL block:

