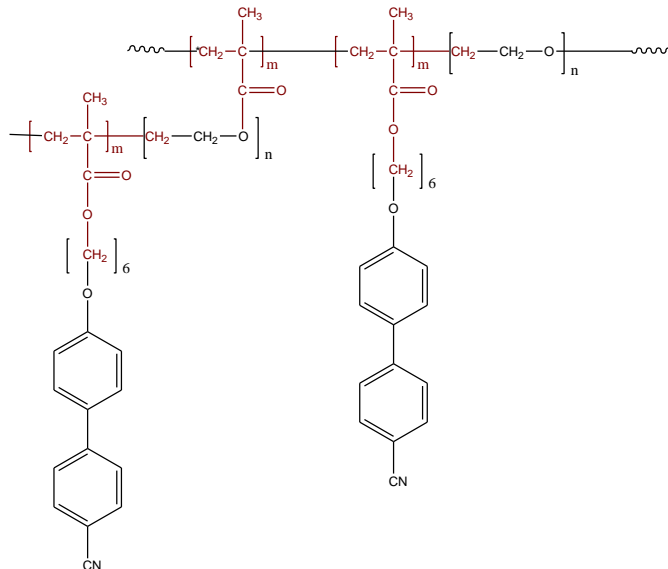


**Sample Name:**

**Poly(6-(4'-cyanobiphenyl-4-yloxy)hexyl methacrylate-block -6-(4'-cyanobiphenyl-4-yloxy)hexyl methacrylate-PEO)**

**Sample #:**

**P9514-4CNBPHMA -b- EO-G- 4CNBPHMAEO**

**Structure:****Composition:**

Mn x 10 <sup>3</sup> 4CNBPHMA-b- EO-G- 4CNBPHMAEO	PDI
10-b-47.0	1.20
Microstructure of 4CNBPHMA block	Syndio;Hetero;iso contents 40:40:20

**Synthesis Procedure:**

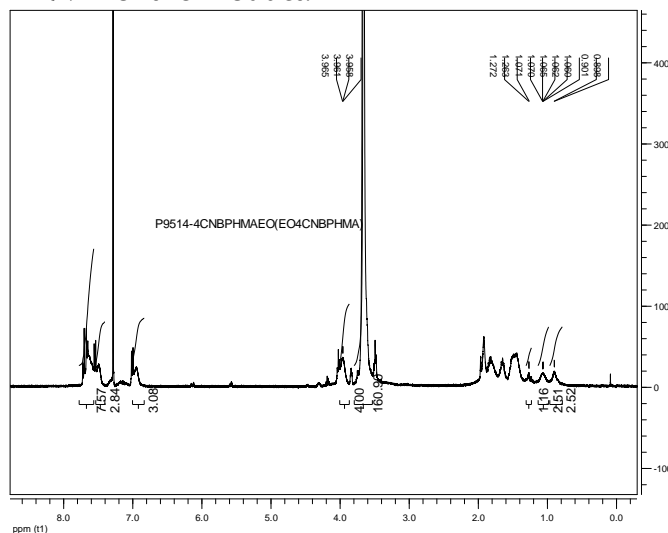
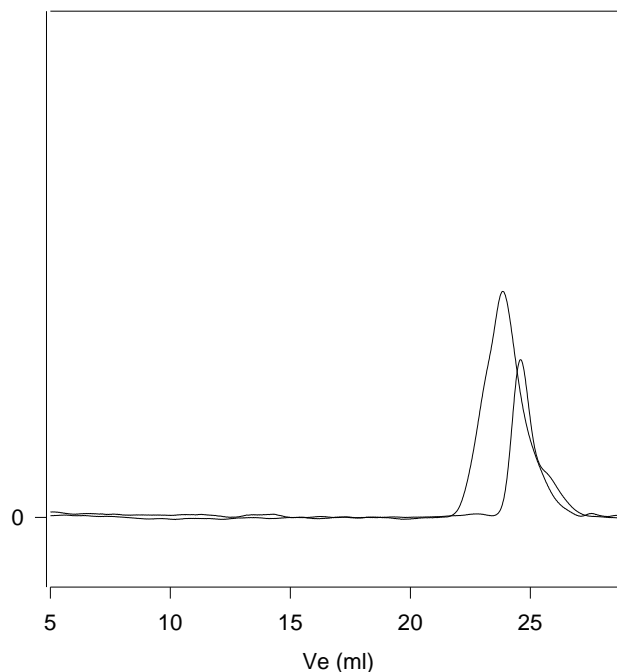
Polymer is synthesized by ionic polymerization process.

**Purification of the polymer:**

The un-reacted PEG can be removed by stirring the polymer in hot water/Methanol. The obtained polymer dissolved in CHCl<sub>3</sub>/toluene and pass through the column packed with silica. The polymer was recovered by precipitation in cold ether/hexane mixture.

**Solubility:**

Polymer is soluble in CHCl<sub>3</sub>, THF and toluene. The polymer precipitated out from hexane.

**HNMR of the Product:****SEC of the block copolymer:****P9514-4CNBPHMA-EO-G-4CNBPHMAEO**

Size exclusion chromatography of the product:

—— Poly(4CNBPHMA), M<sub>n</sub>=10000, M<sub>w</sub>=6000, PI=1.20

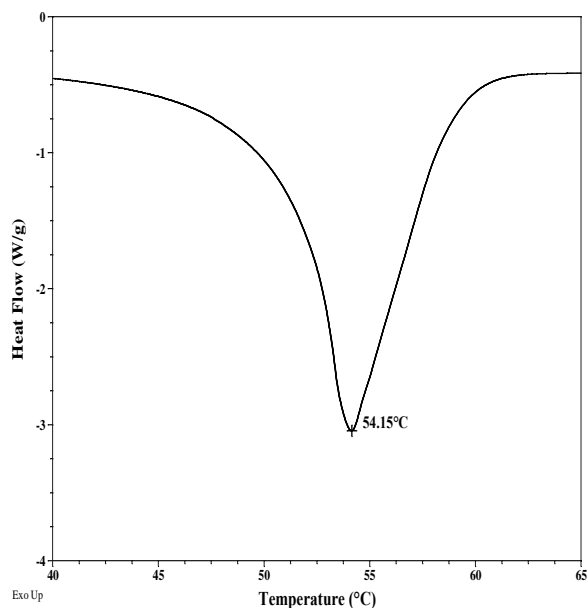
## Thermal analysis of the P9514- 4CNBPHMA EO-G- 4CNBPHMA EO

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.

### Melting curve for PEO block:



### Typical thermal analysis results at a glance:

Sample	$T_m$ (°C)	$T_c$ (°C)	$T_g$ (°C)
EO	54	37	Not distinct
4CNBPHMA	-	-	-

### Crystallization curve for PEO block:

