<u>Sample Name</u>: Poly(N-Vinyl Pyrrolidinone) Sample #: P7583-NVP (dialysed)

Structure:

$$CH3 \longrightarrow O \longrightarrow C \longrightarrow C \longrightarrow C \longrightarrow C \longrightarrow CH_2 \longrightarrow CH_3$$

$$CH3 \longrightarrow O \longrightarrow CH_2 \longrightarrow CH_3$$

$$CH3 \longrightarrow O \longrightarrow CH_2 \longrightarrow CH_3$$

Composition:

Mn x 10 ³	Mw/Mn (PDI)
0.7	1.2

Synthesis Procedure:

Polymer is synthesized by RAFT polymerization of N-vinyl 2-pyrrolidinone using 4,4'-azo(4-cyanopentanoic acid) as initiator and xanthate as chain transfer agent.. Polymerization was carried out for 24h reaction time.

Purification of the Polymer: Polymer was purified by column packed with silica and the eluent was de-ionized water. The obtained polymer solution was dialysed against deionized water for 24h.

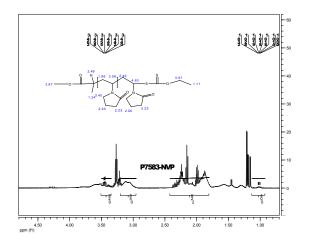
Characterization:

Polymer analyzed by size exclusion chromatography (SEC) to obtain the molecular weight and polydispersity index (PDI) using water containing $0.1M\ NaNO_3$ and $0.01M\ NaH_2PO_4$ as eluent. The Mn was then also calculated from $^1H\text{-}NMR$ spectroscopy by comparing the peak area of the terminal end groups protons at about 3.6 ppm and

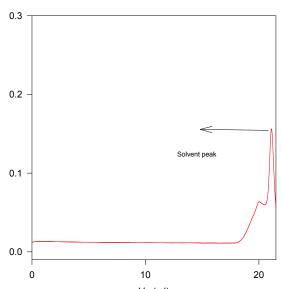
1.2 ppm and of pyrrolidinone units at 3.2 ppm to 2.0 ppm polymer molecular weight distribution is determined by SEC.

Solubility: Polymer is soluble in water.

¹H-NMR Spectrum of polymer bearing terminal Xanthate group:



SEC of Sample of the block copolymer: P7583-NVP



Ve (ml)
Size Exclusion Chromatography of Poly(N-vinylpyrrolidone)
M_o=700, M_w=850, PI=1.2