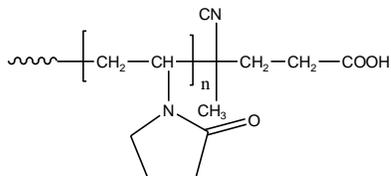


Sample Name:

Poly(N-vinyl pyrrolidone), α -carboxy-terminated

Sample #: **P7110-6-NVPCOOH**

Structure:

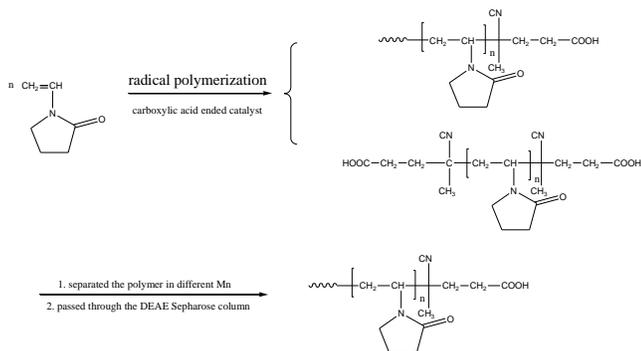


Composition:

$M_n \times 10^3$	PDI
5.7	1.7

Synthesis Procedure:

Monocarboxy terminated poly(N-vinylpyrrolidone) was prepared by radical polymerization of N-vinylpyrrolidone using 4,4'-azobis (4-cyanovaleric acid) as a catalyst. The obtained polymer was fractionated and from each fraction the mono carboxylic acid fraction was separated from its α - ω di carboxylic acid by passing the polymer solution in ethanol through a column packed with DEAE Sepharose resin. The polymer is obtained by precipitation from cold diethyl ether. The scheme of the reaction is illustrated below:



Characterization:

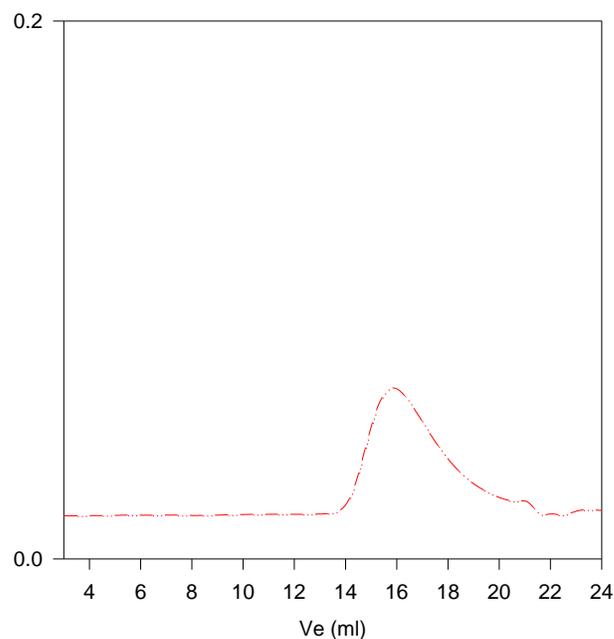
The molecular weight of the polymer was determined by acid base titration and polydispersity was determined by size exclusion chromatography (SEC) using a Varian liquid chromatograph equipped with a UV and refractive index detector in DMF containing 0.01M LiBr salt.

Solubility:

Polymer is soluble in chloroform, THF, DMF, ethanol and water, and precipitate out from hexanes and ether.

SEC profile of the Sample:

P7110-6 NVPCOOH



Size exclusion chromatography in DMF at 40 °C:
Eluent containing 0.01 M LiBr

— Monocarboxylic acid ended poly(N-vinylpyrrolidone),
 $M_n=5700$, $M_w=9700$, $PI=1.7$. (M_n obtained by titration)