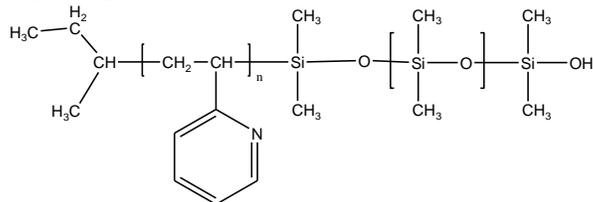


Sample Name: Poly(2-vinyl pyridine-b-dimethylsiloxane)

Sample #: P18682-2VPDMS

By controlled radical process

Structure:



Composition:

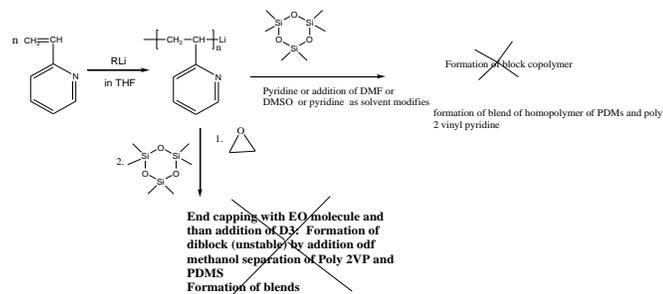
$M_n \times 10^3$ 2VP-b-DMS	Mw/Mn
20.0-b-1.0	1.22

Synthesis Procedure:

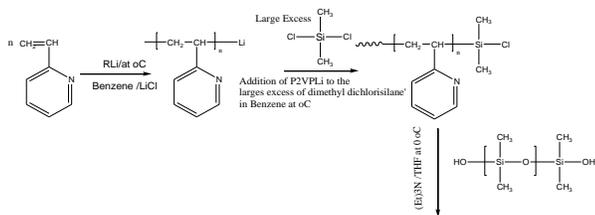
Poly(2-vinyl pyridine-b-dimethylsiloxane) is synthesized by one of the following routes.

Different routes for the synthesis of poly 2 vinyl pyridine with polydimethyl siloxane:

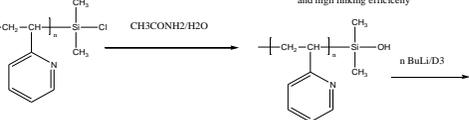
1. Direct Anionic Polymerization by sequential addition of 2VP followed by D3 monomer



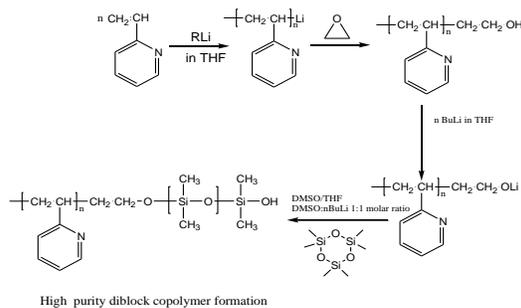
2. From the linking reaction of end functionalized polymer: For the synthesis of Block copolymer > Mn 10,000



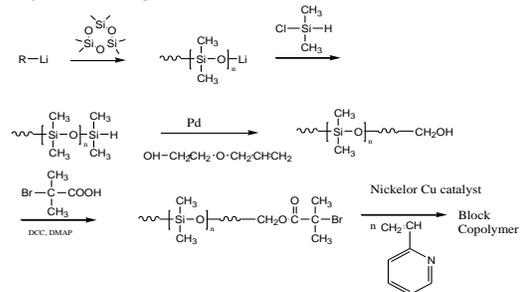
3. Block copolymer formation Mn > 10,000 is excellent and than difficult to adjust the stoichiometry and high linking efficiency



3. Formation of first Poly 2vinyl pyridine OH terminated polymer than reacting the isolated P2VPOH polymer with a BuLi followed by addition of D3 in the presence of DMSO equimolar amount with nBuLi



4. By Controlled radical process:



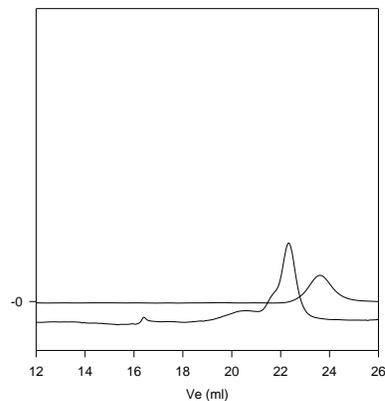
Characterization:

Polymers were analyzed by size exclusion chromatography (SEC) and ¹H-NMR spectroscopy by comparing the peak area of the 2-vinyl pyridine proton at about 8.2 ppm with the dimethyl siloxane protons at 0.08 ppm. Copolymer PDI is determined by SEC.

Solubility:

Poly(2-vinyl pyridine-b-dimethyl siloxane) is soluble in THF, CHCl₃ and toluene

P18682-2VPDMS



Size exclusion chromatography of

— Poly(2VP), M_n=20,000 Mw/Mn 1.18
 — Block Copolymer P2VP(20,000)-b-PDMS(1000), PI= 1.22
 Composition for ¹H NMR

¹H NMR for the polymer:

