Supporting Information for:

Linear Viscoelasticity of Weakly Hydrogen-Bonded Polymers Near and Below Sol-Gel Transition

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1. Experimental Section

1.1. Quantities of the compounds used for RAFT polymerization

Table S1. Amounts of reagents used for the synthesis of the (co)polymers.

Sample ID	AIBN (mg)	DBTTC (mg)	DMF (mL)	nBA (g)	THY (g)
PnBA	0.82	14.5	5.0	1.28	0
PTHY10	0.82	14.5	5.0	1.15	0.32
PTHY30	0.82	14.5	5.0	0.64	1.62
PTHY100	0.82	14.5	5.0	0	3.24

2. Characterization

2.1. ^{1}H NMR of PTHYi

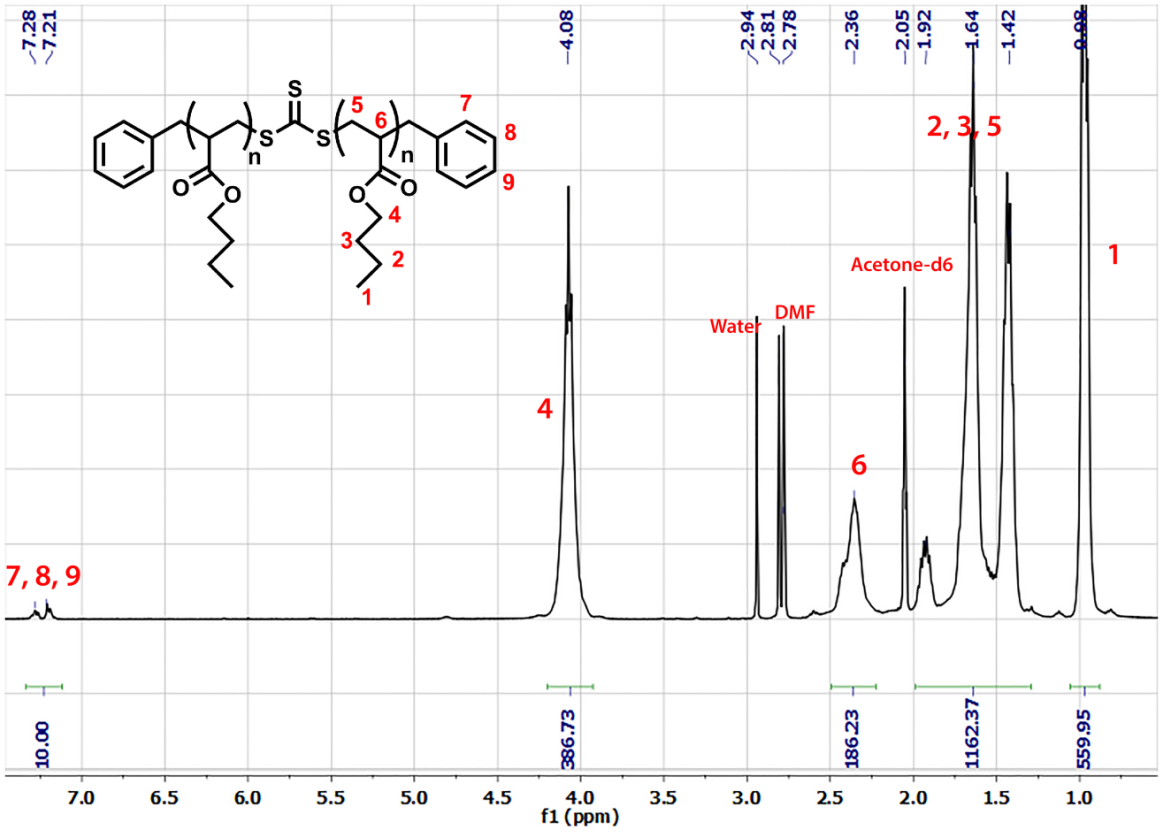

Figure S1. ^{1}H NMR spectrum of PnBA.

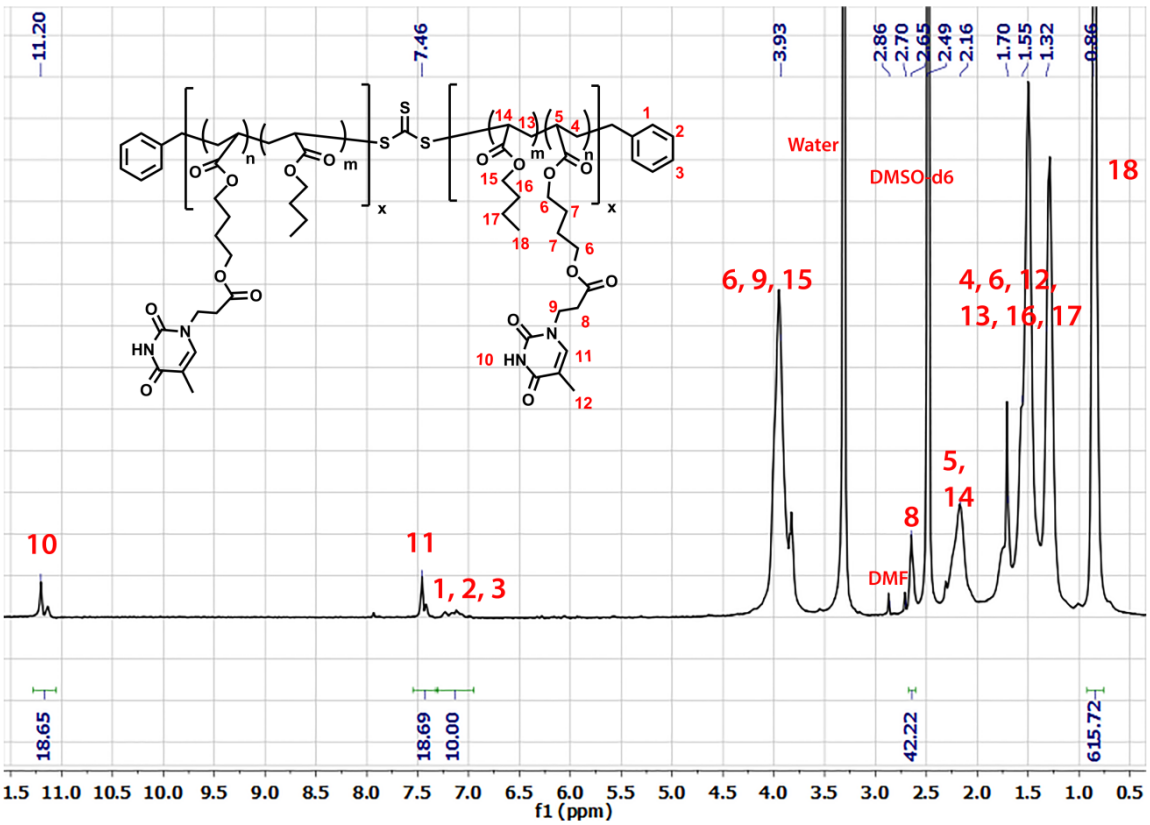

Figure S2. ^{1}H NMR spectrum of PTHY10.

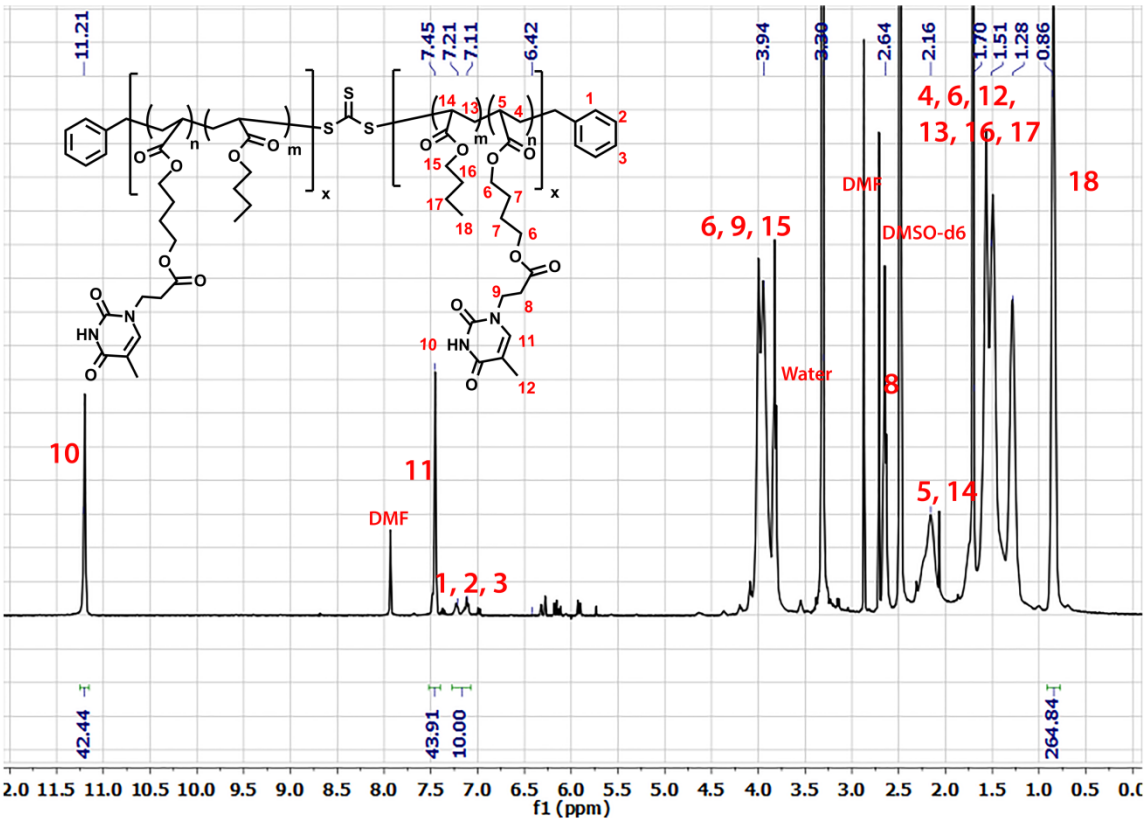

Figure S3. ^{1}H NMR spectrum of PTHY30.

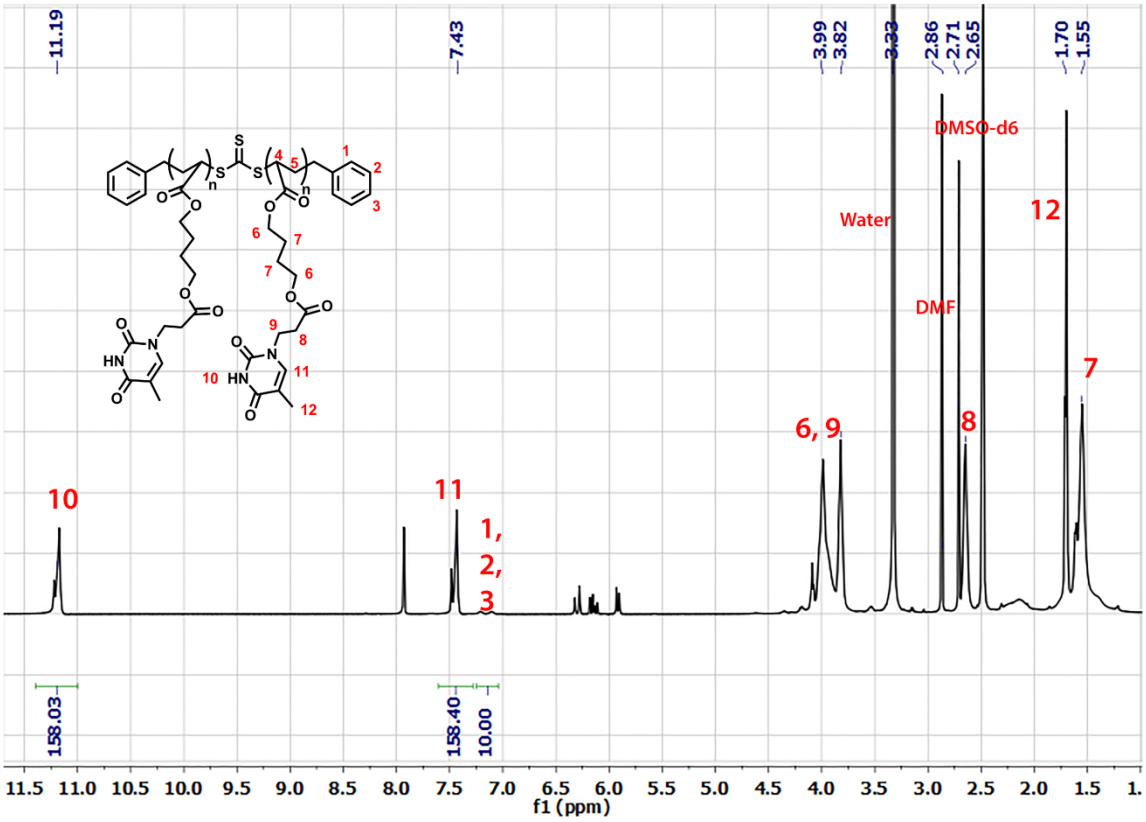

Figure S4. ^{1}H NMR spectrum of PTHY100.

2.2. Comparison of low field chemical shifts for the copolymers

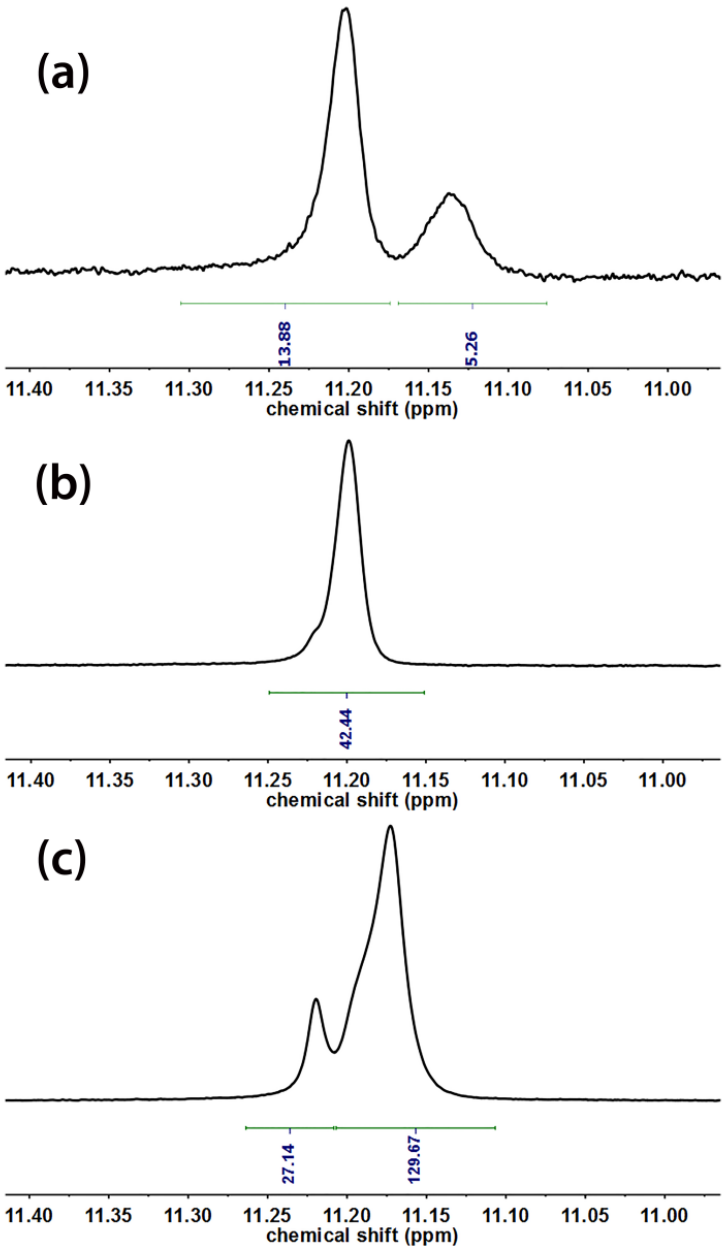

Figure S5. ^{1}H NMR spectrum of (a) PTHY10, (b) PTHY30, and (c) PTHY100, enlarged at high chemical shifts belonging to the hydrogen bondings of thymine.

3. Shift factors and the Arrhenius fitting

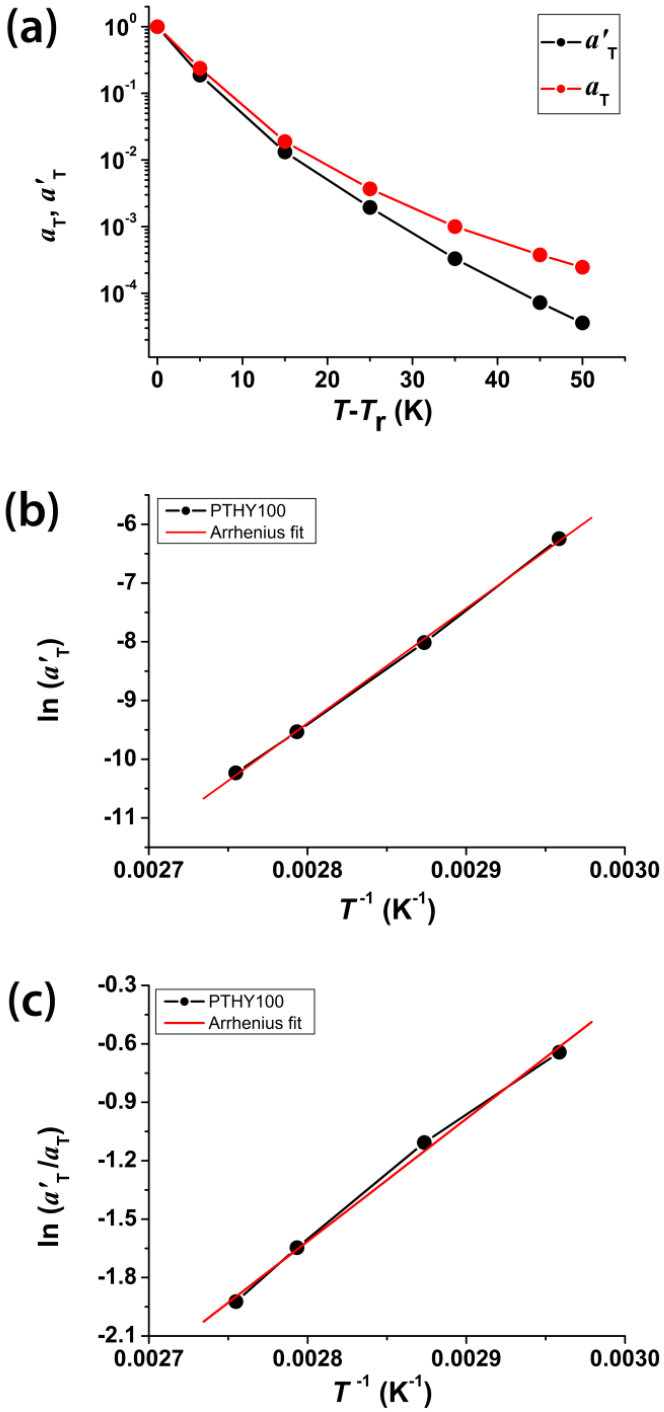

Figure S6. Shift factors a_T and a'_T vs. (a) T – T_r and (b) 1/T for PTHY100 (c) ratio of shift factors obtained from the two methods a'_T/a_T vs. 1/T.