

Supporting information for

Hydrophobically Associating Polymers Dissolved in Seawater for Enhanced Oil Recovery of Bohai Offshore Oilfields

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Table S1. The influence of monomer and initiator dosage on dissolution time and solution viscosity

No.	AM (g)	AA (g)	AMPS (g)	POM (g)	VA-044 (g)	NaHSO ₃ (g)	(NH ₄) ₂ S ₂ O ₈ (g)	Hydrolysis Degree (%)	Dissolution Time ^a (min)	Solution Viscosity ^a (mPa·s)
N1	56	0	15	7	0.01	0.01	0.01	0	20	10.3
N2	184	35	15	7	0.01	0.01	0.01	13	30	21.5
N3	184	70	15	7	0.01	0.01	0.01	25	70	44.5
N4	184	70	30	7	0.01	0.01	0.01	26	65	47.3
N5	184	70	45	7	0.01	0.01	0.01	23	60	54.5
N6	184	70	60	7	0.01	0.01	0.01	25	60	55.2
N7	184	70	60	7	0.01	0.01	0.02	25	55	49.1
N8	184	70	60	7	0.01	0.01	0.03	25	60	47.5
N9	184	70	60	7	0.01	0.01	0.04	25	65	34.2
N10	184	70	60	7	0.01	0.01	0.05	26	65	23.2
N11	184	70	60	7	0.01	0.01	0.06	25	70	15.6
N12	184	70	60	7	0.01	0.01	0	25	65	10.3
N13	184	70	60	10	0.01	0.01	0.01	25	125	99.1
N14	184	70	60	9	0.01	0.01	0.01	25	105	89.6
N15	184	70	60	8	0.01	0.01	0.01	25	95	73.4
N16	184	70	60	6	0.01	0.01	0.01	25	55	37.6
N17	184	70	60	5	0.01	0.01	0.01	25	50	28.5
N18	184	70	60	4	0.01	0.01	0.01	25	40	21.6
N19	184	70	60	3	0.01	0.01	0.01	25	30	17.7
N20	184	70	60	2	0.01	0.01	0.01	25	25	9.8

^a Polymer concentration, 0.175%; particle size of polymer powder, 40-120 mesh; salinity, 3.26×10^5 mg/L; shear rate, 7.34 s^{-1}

Table S2. The influence of reaction condition (the amount of cosolvent and chain transfer agent; pH value) on dissolution time and solution viscosity

No.	AM (g)	AA (g)	AMPS (g)	POM (g)	VA- 044 (g)	NaHSO ₃ (g)	(NH ₄) ₂ S ₂ O ₈ (g)	pH	Hydrolysis Degree (%)	Dissolution Time ^a (min)	Solution Viscosity ^a (mPa·s)
N21	184	70	60	7	0.015	0.01	0.01	8	24	50	57.3
N22	184	70	60	7	0.02	0.01	0.01	8	25	50	55.1
N23	184	70	60	7	0.025	0.01	0.01	8	25	45	52.1
N24	184	70	60	7	0.003	0.01	0.01	8	25	40	49.0
N25	184	70	60	7	0.035	0.01	0.01	8	24	35	46.3
N26	184	70	60	7	0.04	0.01	0.01	8	25	35	42.1
N27	184	70	60	7	0.045	0.01	0.01	8	23	30	39.2
N28	184	70	60	7	0.05	0.02	0.01	8	24	25	35.1
N29	184	70	60	7	0.01	0.03	0.01	8	25	55	56.2
N30	184	70	60	7	0.01	0.05	0.01	8	24	50	61.5
N31	184	70	60	7	0.01	0.06	0.01	8	25	45	58.7
N32	184	70	60	7	0.01	0.01	0.01	8	23	35	56.5
N33	184	70	60	7	0.01	0.02	0.01	8	24	35	54.2
N34	184	70	60	7	0.01	0.03	0.01	8	25	30	49.1
N35	184	70	60	7	0.01	0.10	0.01	8	25	25	41.0
N36	184	70	60	7	0.01	0.01	0.01	2	23	N/A ^b	N/A ^b
N37	184	70	60	7	0.01	0.01	0.01	4	23	N/A ^b	N/A ^b
N38	184	70	60	7	0.01	0.01	0.01	6	23	55	60.0
N39	184	70	60	7	0.01	0.01	0.01	8	23	40	56.5
N40	184	70	60	7	0.01	0.01	0.01	10	23	40	49.2
N41	184	70	60	7	0.01	0.01	0.01	12	23	35	37.1

^a Polymer concentration, 0.175%; particle size of polymer powder, 40-120 mesh; salinity, 3.26×10^5 mg/L; shear rate, 7.34 s^{-1}

^b The polymers do not dissolve, so their aqueous solution viscosity and dissolution time cannot be measured.

Table S3. The effect of reaction time and temperature on dissolution time and solution viscosity

No.	AM (g)	AA (g)	AMPS (g)	POM (g)	VA-044 (g)	NaHSO ₃ (g)	(NH ₄) ₂ S ₂ O ₈ (g)	Reaction Time (h)	Hydrolysis Degree (%)	Reaction Temperature (°C)	Dissolution Time ^a (min)	Solution Viscosity ^a (mPa·s)
N42	184	70	60	7	0.01	0.01	0.01	3	24	0	10	6.4
N43	184	70	60	7	0.01	0.01	0.01	6	25	0	10	17.9
N44	184	70	60	7	0.01	0.01	0.01	9	25	0	25	36.1
N45	184	70	60	7	0.01	0.01	0.01	12	25	0	27	54.2
N46	184	70	60	7	0.01	0.01	0.01	15	24	0	32	55.6
N47	184	70	60	7	0.01	0.01	0.01	12	25	0	40	56.2
N48	184	70	60	7	0.01	0.01	0.01	12	23	10	35	46.1
N49	184	70	60	7	0.01	0.01	0.01	12	24	20	30	32.6
N50	184	70	60	7	0.01	0.01	0.01	12	25	30	30	19.5
N51	184	70	60	7	0.01	0.01	0.01	12	24	40	30	11.3
N52	184	70	60	7	0.01	0.01	0.01	12	25	0	30	55.2

^a Polymer concentration, 0.175%; particle size of polymer powder, 40-120 mesh; salinity, 3.26×10^5 mg/L; shear rate, 7.34 s^{-1}

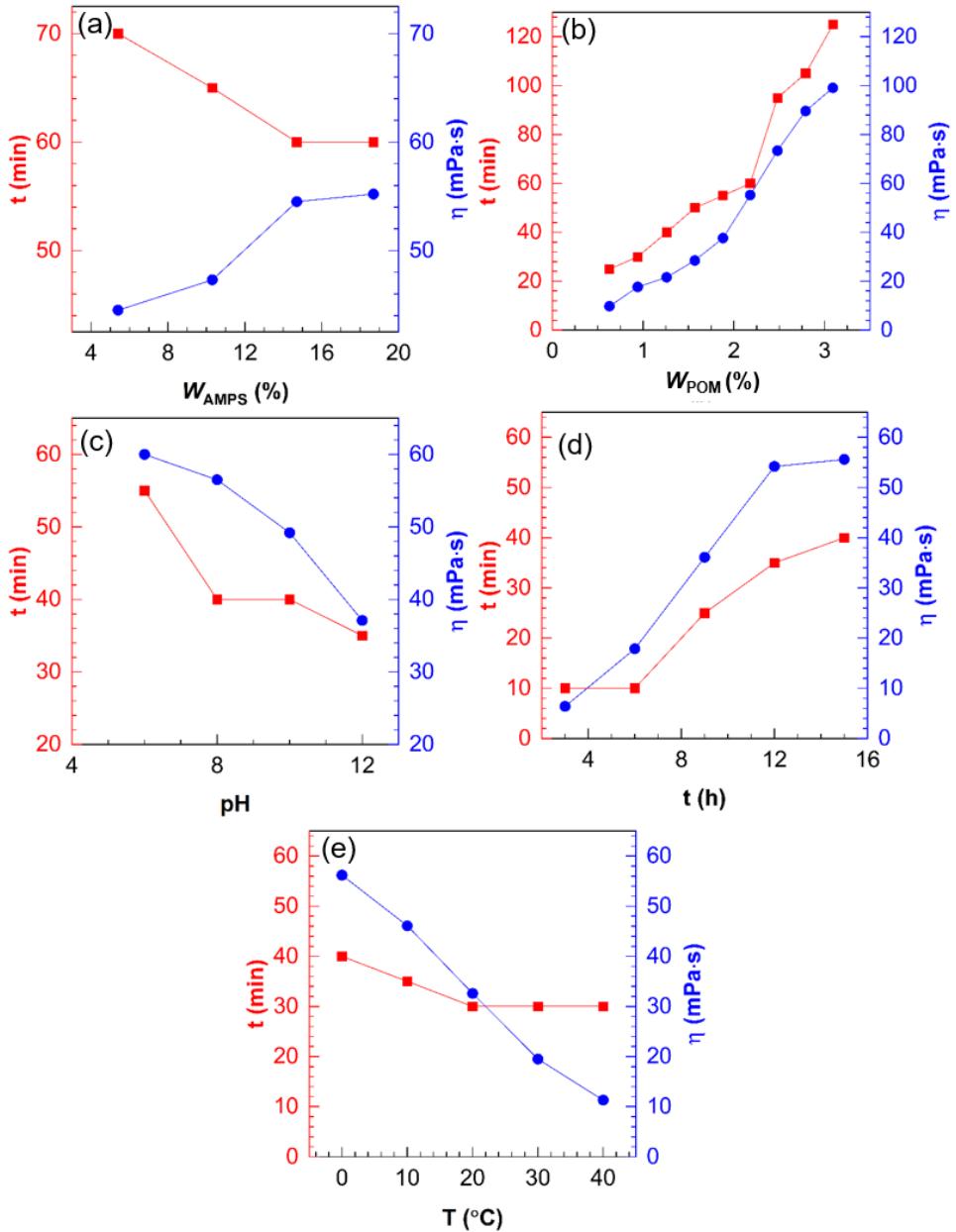


Figure S1. Effect of components and reaction conditions of polymerization under low temperature on dissolution time and viscosity of HAP polymer. (a) Content of AMPS comonomer and the data from N3-N6 in Table S1; (b) Content of POM comonomer and the data from N6, N13-N20 in Table S1; (c) pH value for solution of reactants prior to polymerization and the data from N36-N41 in Table S2; (d) reaction time of polymerization and the data from N42-N46 in Table S3; (e) Temperature of the reaction system at the beginning of polymerization the data from N48-N52 in Table S3.

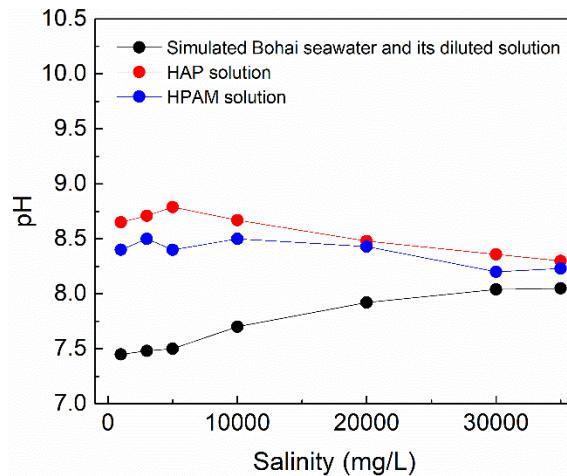


Figure S2. Variation of pH with salinity in HAP, HPAMM solutions and simulated Bohai seawater as well as dilute solutions at 75°C (Polymer concentration, 0.175%; shear rate, 7.34 s⁻¹).

Table S4. Basic parameters of the artificial cores used in this work.

Core	<i>L</i> (cm)	<i>d</i> (cm)	PV (mL)	ϕ (%)	<i>K_w</i> (mD)
No. 1	10.1	2.5	12.5	15.9	331
No. 2	10.0	2.5	12.4	16.1	333
No. 3	10.1	2.5	13.6	16.2	342
No. 4	10.1	2.5	13.2	16.0	339

Table S5. RF and RRF results for solutions of HAP and HPAM (Polymer concentration, 0.175%).

Core	Polymer sample	Permeability	RF	RRF
No. 1	HAP	240 mD	38.7	14.3
No. 2	HPAM	254 mD	25.3	9.5

Table S6. Summary of recovery factors for aqueous solutions of HAP and HPAM.

Core	Samples	E_w (%)	E_p (%)	E_T (%)
No. 3	HAP	22.4	19.7	42.1
No. 4	HPAM	22.5	10.9	33.4

E_w denotes the incremental oil recovery after initial water-flooding.

E_p refers to the incremental oil recovery after polymer injection.

E_T represents the total recovery.