

# Effects of Hofmeister Anions on the LCST of PNIPAM as a Function of Molecular Weight<sup>†</sup>

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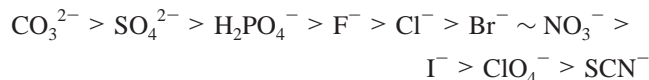
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The effect of a series of sodium salts on the lower critical solution temperature (LCST) of poly(*N*-isopropylacrylamide), PNIPAM, was investigated as a function of molecular weight and polymer concentration with a temperature gradient microfluidic device under a dark-field microscope. In solutions containing sufficient concentrations of kosmotropic anions, the phase transition of PNIPAM was resolved into two separate steps for higher molecular weight samples. The first step of this two-step transition was found to be sensitive to the polymer's molecular weight and solution concentration, while the second step was not. Moreover, the binding of chaotropic anions to the polymer was also influenced by molecular weight. Both sets of results could be explained by the formation of intramolecular and intermolecular hydrogen-bonding between polymer chains. By contrast, the hydrophobic hydration of the isopropyl moieties and polymer backbone was found to be unaffected by either the polymer's molecular weight or solution concentration.

## Introduction

Protein stability, enzyme activity, macromolecule crystallization, as well as protein and polymer folding can be strongly influenced by the type of salts present in the aqueous solution.<sup>1–7</sup> In fact, the ability of ions to influence a broad range of physical properties typically follows a recurring trend called the Hofmeister series.<sup>8–16</sup> The typical order of the series for anions is



The ions on the left are strongly hydrated and referred to as kosmotropes, while the species on the right are more weakly hydrated and called chaotropes. Recently, considerable effort has gone into understanding the ordering of this series and its underlying meaning.<sup>5,14,17–20</sup> Although it was originally suspected that the series had to do with the ordering of bulk water, this turns out not to be the case. Therefore, mechanisms to explain the behavior of specific systems need to take into account the direct interactions of the ions with the macromolecule and its first hydration shell.

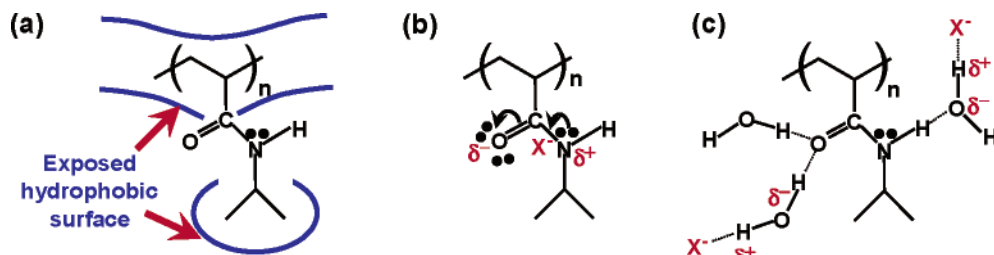
In water, PNIPAM behaves as a thermoresponsive polymer which aggregates and falls out of solution above its lower critical solution temperature (LCST).<sup>21,22</sup> This behavior follows the Hofmeister series.<sup>23–25</sup> Our laboratory has recently elucidated the mechanism of the phase transition for PNIPAM in the presence of salts.<sup>5</sup> This was made possible by the development of a temperature gradient microfluidic apparatus<sup>5,26–30</sup> in combination with dark-field microscopy, which allowed the LCST of PNIPAM to be determined with extremely high-temperature precision. The experiments were conducted on PNIPAM samples with a relatively narrow molecular weight distribution (polydispersity index = 1.40). The number average molecular weight,  $M_n$ , of these samples was 121 000 Da. It was found that three basic interactions were involved depending upon

the specific conditions, a surface tension effect, direct anion binding, and the polarization of water molecules directly hydrating the amide moieties of the polymer (Figure 1). Sodium salts of the chaotropic anions were found to depress the LCST by increasing the surface tension between the hydrophobic portions of the polymer (isopropyl group and hydrocarbon backbone) and adjacent hydration waters. There was also a modest salting-in effect caused by the binding of chaotropic anions. These two mechanisms could account for the behavior of the ions from  $\text{Cl}^-$  to  $\text{SCN}^-$ . On the other hand, the kosmotropes ( $\text{CO}_3^{2-}$ ,  $\text{SO}_4^{2-}$ ,  $\text{H}_2\text{PO}_4^-$ , and  $\text{F}^-$ ) behaved differently. In this case, no salting-in effects were observed. Rather, the anions were found to weaken the hydrogen bonding between the pendent amide groups and the bound hydration waters via a polarization effect. In the presence of a sufficient concentration of a kosmotropic salt, the dehydration of the hydrophilic amide groups would actually occur separately from the removal of hydrophobic hydration waters from the rest of the polymer. In fact, the phase transition was resolved into two distinct steps. The first step was correlated with the dehydration of the amide, while the second involved the loss of water molecules which were hydrophobically hydrating the polymer.

It is known that changing the molecular weight of PNIPAM does not strongly affect its LCST unless the degree of polymerization is low enough that the end group structure becomes significant.<sup>31,32</sup> We wondered, however, if changes in the molecular weight would affect the salt dependence. Four samples were prepared with number average molecular weights ranging from 6980 to 360 000 Da. The results showed that the molecular weight of PNIPAM does influence salt effects. Specifically, it was found that the concentration of kosmotropic ions required to resolve the phase transition into two distinct steps increased with decreasing molecular weight. In fact, it was not possible to decouple the amide and hydrophobic dehydration steps at any salt concentration for the lowest molecular weight sample we investigated. Moreover, molecular weight also influenced the behavior of PNIPAM in the presence of chaotropic anions. In this case, it was found that the binding

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**Figure 1.** Three basic interactions among anions, PNIPAM, and hydration waters are involved in the Hofmeister effect: (a) The hydrophobic hydration of the molecule is associated with surface tension and can be modulated by salt; (b) direct binding of the anion to the amide group of PNIPAM; (c) hydrogen bonding of the amide and its destabilization through polarization by the anion,  $X^-$ .

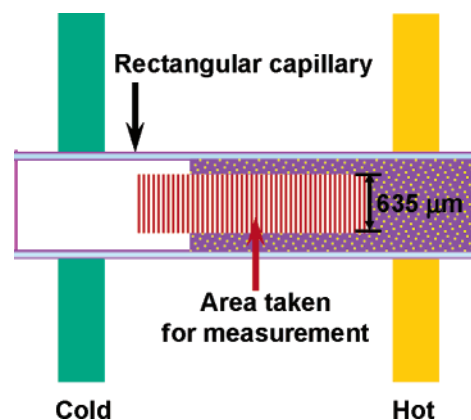
constant of the anions increased with increasing molecular weight and polymer concentration, while the salting-in effect decreased with increasing molecular weight and polymer concentration. By contrast, neither the molecular weight of the polymer nor its solution concentration affected the phase transition when triggered by the loss of water molecules involved in hydrophobic hydration.

### Experimental

**Preparation and Fractionation of PNIPAM.** PNIPAM was obtained by free-radical polymerization of its monomer using 2,2'-azoisobutyronitrile (AIBN) as an initiator in methanol at 70 °C under a positive pressure of  $N_2$ . The purified PNIPAM was fractionated by redissolving it in acetone (10 g/L) and through stepwise precipitation by the addition of hexanes.<sup>31,33</sup> This gave rise to four polymer fractions with weight average molecular weights ( $M_w$ ) of  $4.75 \times 10^5$ ,  $1.70 \times 10^5$ ,  $5.58 \times 10^4$ , and  $1.78 \times 10^4$  g/mol as determined by static light scattering in methanol. Light-scattering experiments were carried out using a Brookhaven Instruments BI-200SM goniometer, a BI-9000AT digital correlator, and a Melles Griot helium neon laser. Molecular weight analysis of light-scattering data was performed using Brookhaven Instruments' Zimm Plot Software. The fractionated samples described above were analyzed by gel permeation chromatography in tetrahydrofuran. This analysis was carried out with a Viscotek I-MBMMW-3078 mixed bed column using a Viscotek gel permeation chromatography instrument. The instrument was equipped with multiple detectors including a Model VE 3580 RI detector and OmniSEC software. The results indicated that the polydispersity index (PDI) of the four fractions used in these studies were 1.32, 1.40, 1.82, and 2.55, respectively, based upon polystyrene reference standards. The  $M_n$  values for these four fractions were determined to be  $3.60 \times 10^5$ ,  $1.21 \times 10^5$ ,  $3.07 \times 10^4$ , and  $6.98 \times 10^3$  Da, respectively. It should be noted that  $M_n = M_w/PDI$ .

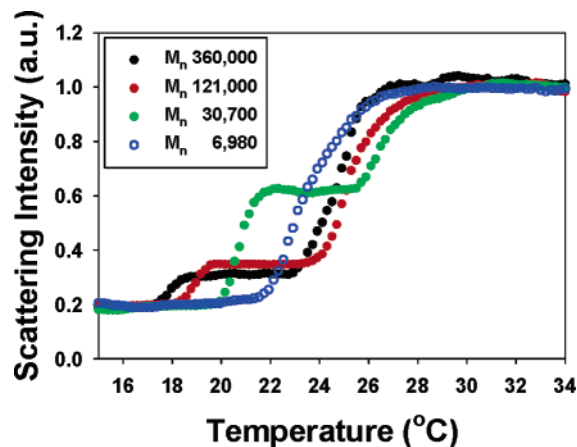
**Preparation of Polymer Solutions.** All inorganic salts used in these experiments were purchased from Aldrich. Low-conductivity  $H_2O$ , produced from a NANOpure Ultrapure Water System (Barnstead, Dubuque, IA) with a minimum resistivity of 18  $M\Omega \cdot cm$ , was used to prepare the polymer and salt solutions. Stock solutions of PNIPAM were prepared by allowing the polymer to dissolve overnight in  $H_2O$  at 4 °C. Aliquots of these solutions were added to water or inorganic salt solutions at the desired concentration. The final polymer concentration for all experiments was 10 mg/mL except for the concentration dependence experiments. In that case, the concentration of the  $M_n$  121 000 sample was set to 2, 5, 10, and 15 mg/mL to study the effect of polymer concentration on the LCST behavior.

**Fabrication of a Temperature Gradient Device and Measurement of LCSTs.** Fabrication of our temperature gradient device has been described previously.<sup>5,27–30</sup> Briefly, a



**Figure 2.** Schematic diagram of the linear temperature gradient used to obtain light-scattering curves. The PNIPAM solution inside the capillary tube shows no light scattering at low temperature (white region inside the tube) but scatters light strongly above the lower critical solution temperature (purple region inside the tube). Data were taken in slices (red vertical lines) from the middle of each tube, and the scattering intensity was averaged over a  $635 \mu m$  line at each temperature.

cover glass acting as both the sample stage and the heat conductor was mounted on top of two parallel brass tubes (1/8 in. wide, K & S Engineering, Chicago, IL), through which hot and cold antifreeze solutions could be individually flowed using standard water bath circulators (Fisher Scientific, Pittsburgh, PA). Rectangular borosilicate capillary tubes (VitroCom, Inc.) with dimensions of 2 cm  $\times$  1 mm  $\times$  100  $\mu m$  (length  $\times$  width  $\times$  height) were used as sample containers and placed parallel to the temperature gradient. Light scattering from polymer solutions was monitored via a CCD camera (Micromax 1024, Princeton Instruments) using dark-field optics under an inverted microscope (Nikon, TE2000-U). Two different polymer solutions with previously measured LCSTs served to calibrate the slope of the gradient in each experiment.<sup>5,27–30</sup> The first control solution was a 10 mg/mL sample of  $M_w$  500 000 Da PNIPAM in  $H_2O$  with a known LCST of  $30.98 \pm 0.05$  °C, and the second was a 10 mg/mL sample of  $M_w$  500 000 Da PNIPAM in 0.7 M KCl with a known LCST of  $22.61 \pm 0.05$  °C.<sup>34</sup> Six rectangular capillary tubes could be fit side-by-side under a  $\times 2$  objective. The temperature along the capillary tubes was determined by counting pixels in a linescan drawn along the direction of the gradient in the corresponding CCD images. The scattering intensity at each temperature was determined by averaging data from a  $635 \mu m$  long slice (50 pixels) perpendicular to the linear temperature gradient (Figure 2). LCST values were taken as the initial break points<sup>23</sup> of these light-scattering intensity versus temperature curves. The measurement of the LCST at each salt concentration was repeated at least eight times, and the mean value was taken. The measurements had a typical standard error of  $\pm 0.05$  °C. It should be noted



**Figure 3.** Light-scattering curves for PNIPAM with different molecular weights in 0.3 M  $\text{Na}_2\text{SO}_4$  solutions. The two-step phase transition can be visualized for PNIPAM with  $M_n$  of 360 000, 121 000, and 30 700 Da.

that the phase transition of PNIPAM was completely reversible under all conditions which were explored.

### Results

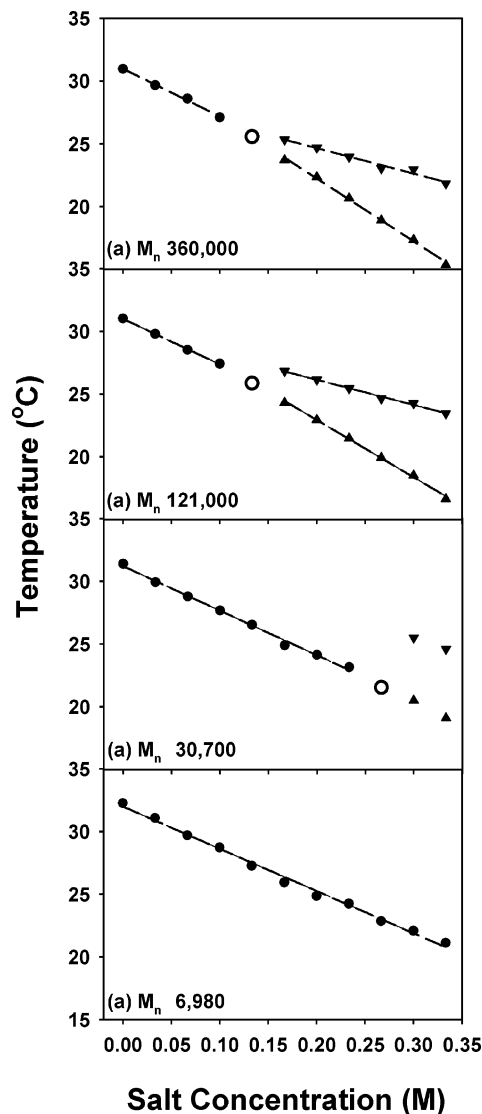
Sodium salts of  $\text{CO}_3^{2-}$ ,  $\text{SO}_4^{2-}$ ,  $\text{H}_2\text{PO}_4^-$ ,  $\text{F}^-$ ,  $\text{Cl}^-$ ,  $\text{Br}^-$ ,  $\text{NO}_3^-$ ,  $\text{I}^-$ ,  $\text{ClO}_4^-$ , and  $\text{SCN}^-$  were employed to investigate specific ion effects on the LCST of PNIPAM because they are representative of the entire Hofmeister series. For  $\text{CO}_3^{2-}$ ,  $\text{SO}_4^{2-}$ ,  $\text{H}_2\text{PO}_4^-$ , and  $\text{F}^-$ , the phase transition of PNIPAM was resolved into two distinct steps in the presence of sufficient concentrations of the respective sodium salts. The first step was the salting-out of the amide group, and the second involved the dehydration of the hydrophobic portions of the polymer.<sup>5</sup> Representative data in Figure 3 at 0.3 M  $\text{Na}_2\text{SO}_4$  clearly reveal that the presence of a two-step LCST is dependent on the chain length of the polymer. In fact, the two-step transition only occurred at a sufficiently high molecular weight of the PNIPAM.

Plots of the temperature for the onset of the lower and higher phase transition steps versus  $\text{Na}_2\text{SO}_4$  concentration are shown in Figure 4 for the four PNIPAM samples. At low salt concentration, only a single LCST was observed and is designated by a filled circle in the plots (●). As the salt concentration was raised, the phase transition was resolved into two separate steps for all but the lowest molecular weight sample. The appearance of the lower step, ▲, and upper step, ▼, of the two-step transition required higher salt concentrations as the molecular weight was decreased. An open circle, ○, is used to denote the concentration where the transition first begins to resolve into two distinct steps. At the lowest molecular weight employed, 6980 Da, only a single phase transition could be found over the entire concentration range investigated.

The LCST versus salt concentration data for each step of the two-step transition can be fit by a simple linear equation

$$T = T_0 + c[M] \quad (1)$$

where  $T_0$  is the LCST of PNIPAM in the absence of salt and  $[M]$  is the molar concentration of the kosmotropic salt. The constant,  $c$ , has units of temperature/molarity, and the individual steps are associated with distinct slopes. In all cases, it was found that the lower step, ▲, of a given two-step transition displayed steeper salt concentration dependence than its corresponding single-step transition, ●, seen at lower  $\text{Na}_2\text{SO}_4$  concentrations. On the other hand, the higher temperature phase transition step, ▼, showed shallower salt dependence than the

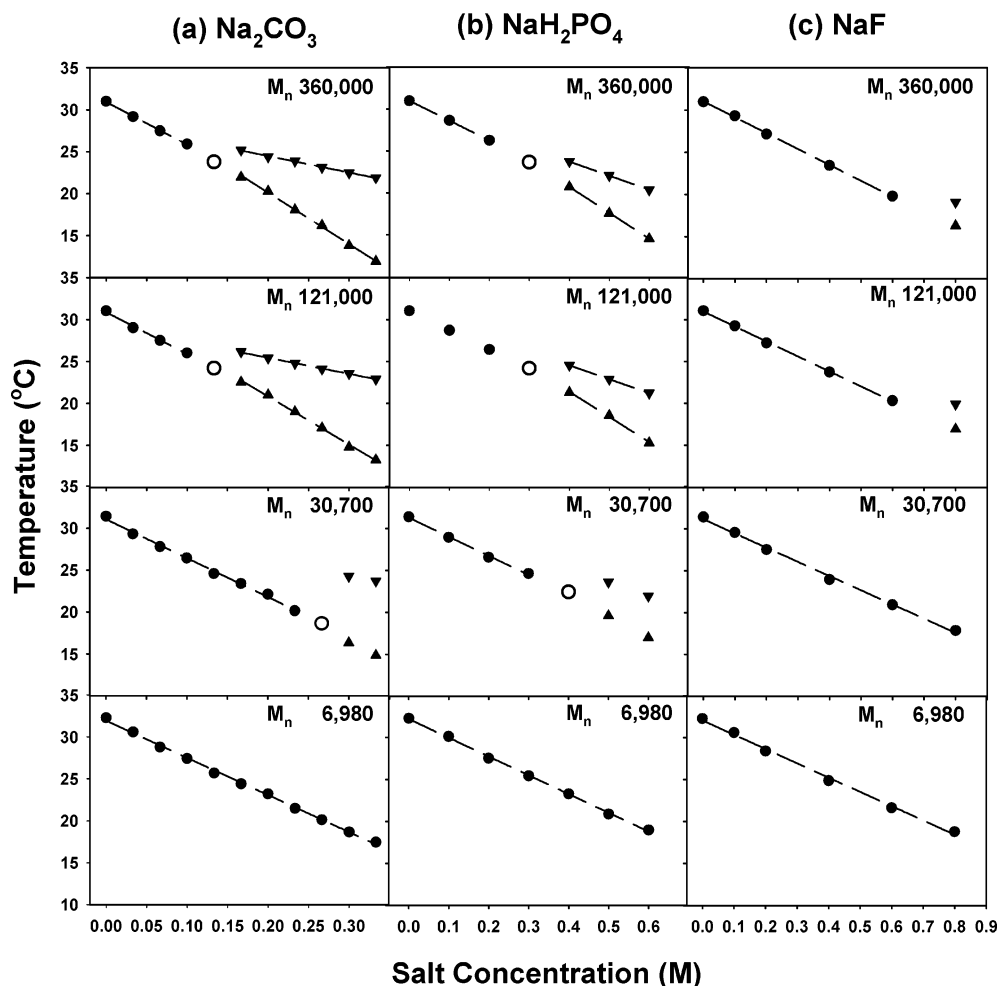


**Figure 4.** LCSTs of 10 mg/mL PNIPAM with different molecular weights in  $\text{Na}_2\text{SO}_4$ : (a)  $M_n$  360 000, (b) 121 000, (c) 30 700, and (d) 6980 Da. The dots (●) represent the initial slope, the upward pointing triangles (▲) designate the lower temperature step in the two-step phase transition, and the downward pointing triangles (▼) are for the higher temperature transition. The unfilled larger circles are used for the point where the phase transition widens without explicitly splitting into two separate steps. This designates the crossover of the transition from a single-step to a two-step process.

initial transition. The other kosmotropic anions,  $\text{CO}_3^{2-}$ ,  $\text{H}_2\text{PO}_4^-$  and  $\text{F}^-$ , behaved similarly with molecular weight. Their LCST values as a function of salt concentration are provided in Figure 5. The abstracted  $c$  values for all of the kosmotropes are given in Table 1.

To investigate the polymer concentration effect in the presence of kosmotropic anions, the 121 000 Da sample was investigated at 2, 5, 10, and 15 mg/mL in the presence of  $\text{Na}_2\text{SO}_4$ , and the data are plotted in Figure 6. Again, the LCST versus salt concentration data for each step of the two-step phase transition were fit to eq 1. For the lower temperature step, the slope of the salt concentration dependence became progressively steeper at higher polymer concentration, while the upper step was found to be insensitive to the polymer concentration. The abstracted  $c$  values for each step are provided in Table 2.

For the chaotropic anions ( $\text{Cl}^-$ ,  $\text{Br}^-$ ,  $\text{NO}_3^-$ ,  $\text{I}^-$ ,  $\text{SCN}^-$ , and  $\text{ClO}_4^-$ ), only a single LCST was observed, independent of the polymer's molecular weight or the salt concentration employed.



**Figure 5.** LCSTs of 10 mg/mL PNIPAM with different molecular weights in (a)  $\text{Na}_2\text{CO}_3$ , (b)  $\text{NaH}_2\text{PO}_4$ , and (c)  $\text{NaF}$ . The molecular weight is labeled on each figure. The symbols have the same meaning as in Figure 4.

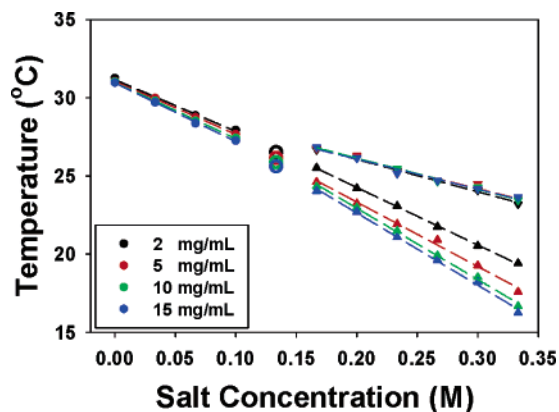
**TABLE 1: Fitted  $c$  Values from the LCST versus Salt Concentration Curves of PNIPAM with Different Molecular Weights for the Kosmotropic Anions**

Anion	$M_n$ (g/mol)	$c$ ( $^{\circ}\text{C}/\text{mol}$ )		
		initial	first step	second step
$\text{CO}_3^{2-}$	6980	-44.3	—	—
	30 700	-46.3	—	—
	121 000	-49.7	-57.4	-19.2
	360 000	-50.8	-61.3	-19.7
$\text{SO}_4^{2-}$	6980	-33.7	—	—
	30 700	-35.5	—	—
	121 000	-36.2	-45.6	-20.0
	360 000	-38.0	-50.0	-20.3
$\text{H}_2\text{PO}_4^-$	6980	-22.4	—	—
	30 700	-22.7	—	—
	121 000	-23.1	-30.1	-16.6
	360 000	-23.6	-30.9	-16.8
$\text{F}^-$	6980	-17.0	—	—
	30 700	-16.9	—	—
	121 000	-17.9	—	—
	360 000	-18.8	—	—

Figure 7 shows the LCST values of PNIPAM for all four molecular weights as a function of salt concentration. The data in Figure 7 can be modeled by eq 2, which includes a constant, a linear term, and a Langmuir isotherm<sup>5</sup>

$$T = T_0 + c[M] + \frac{B_{\max} K_A [M]}{1 + K_A [M]} \quad (2)$$

$T_0$  is again the LCST of PNIPAM in the absence of salt, and  $[M]$  is the molar concentration of the salt.  $K_A$  is the apparent



**Figure 6.** LCSTs of  $M_n$  121 000 Da PNIPAM with polymer concentrations of 2, 5, 10, and 15 mg/mL in  $\text{Na}_2\text{SO}_4$ . The symbols have the same meaning as in Figure 4.

equilibrium association constant of the chaotropic anion with the polymer.  $B_{\max}$  is the increase in the phase transition temperature due to direct ion binding at saturation. The abstracted values for  $c$ ,  $K_A$ , and  $B_{\max}$  for the chaotropic anions are listed in Table 3.

Subtracting the linear contribution ( $T_0 + c[M]$ ) from the concentration versus LCST curves in Figure 7 and replotting the residual data revealed Langmuir-like binding isotherms for the chaotropes (Figure 8). From this figure, the molecular weight dependence of the binding of the anions can be readily observed. It should be noted that  $\text{Cl}^-$  bound weakly enough that deviation

**TABLE 2: Fitted  $c$  Values from the LCST versus  $\text{Na}_2\text{SO}_4$  Concentration Curves for a  $M_n$  121 000 Sample of PNIPAM at Different Polymer Concentrations**

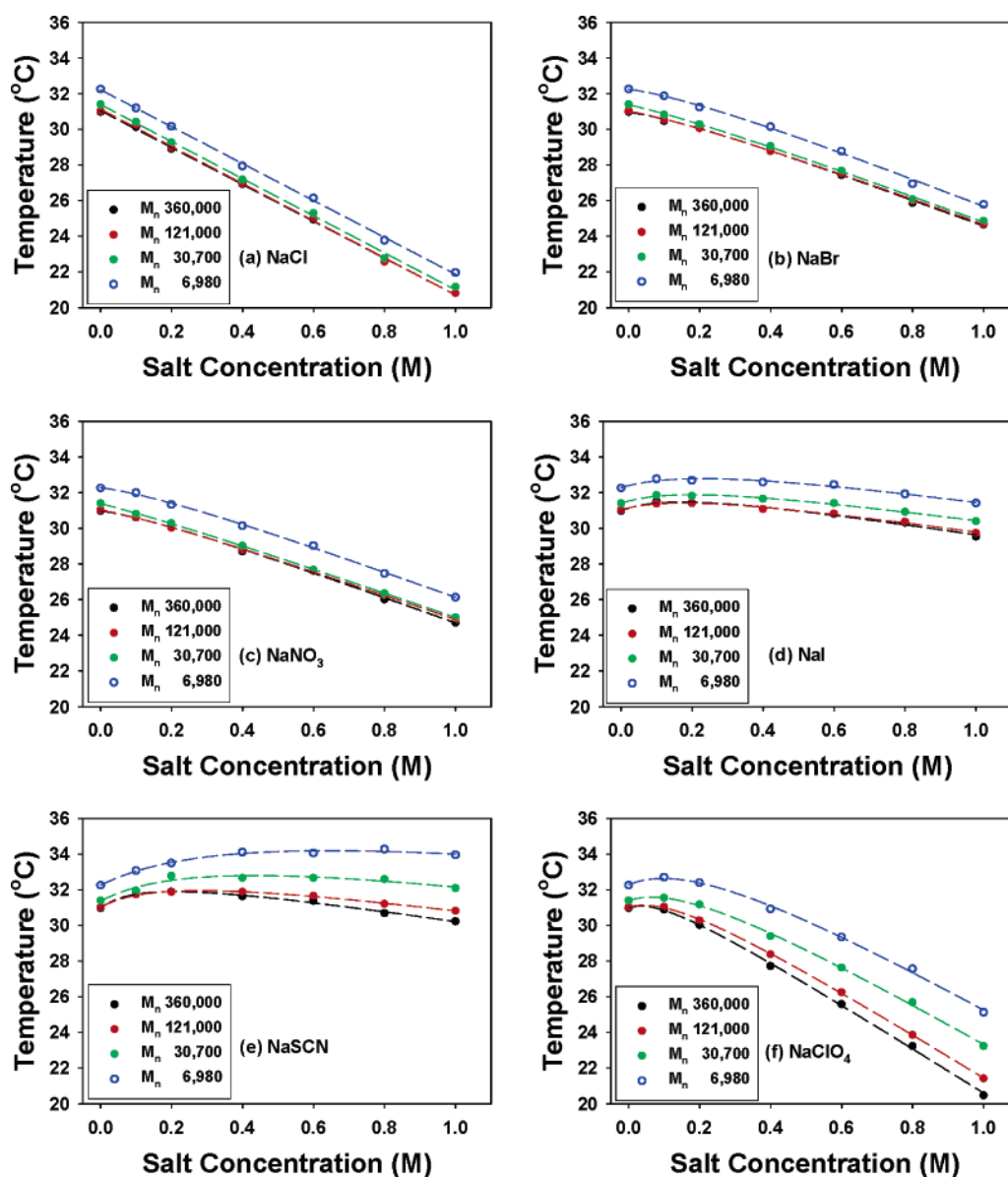
Anion	$M_n$ (g/mol)	Concn (mg/mL)	$c$ ( $^{\circ}\text{C}/\text{mol}$ )		
			initial	first step	second step
$\text{SO}_4^{2-}$	121 000	2	-33.2	-36.8	-20.7
		5	-34.2	-41.2	-19.2
		10	-36.2	-45.6	-20.0
		15	-37.5	-46.2	-19.2

from linear LCST versus concentration behavior was not observed. The  $\text{Br}^-$ ,  $\text{NO}_3^-$ , and  $\text{I}^-$  binding to PNIPAM showed relatively weak molecular weight dependence. On the other hand,  $\text{SCN}^-$  and  $\text{ClO}_4^-$ , which bound most tightly to the polymer, showed the most significant dependence on molecular weight. Specifically,  $K_A$  increased with increasing molecular weight, while  $B_{\text{max}}$  was attenuated with increasing molecular weight (Table 3).

The polymer concentration's effect on the LCST behavior of PNIPAM in the presence of  $\text{NaClO}_4$  was investigated with the  $M_n$  121 000 sample at 2, 5, 10, and 15 mg/mL. The results are shown in Figure 9a. The data were fit to eq 2. After

subtraction of the linear contribution to the curves, the data were replotted to reveal the shapes of the binding isotherms (Figure 9b). The abstracted  $c$ ,  $K_A$ , and  $B_{\text{max}}$  values as a function of polymer concentration are listed in Table 4. Significantly, the binding constant,  $K_A$ , increased with increasing polymer concentration, while  $B_{\text{max}}$  decreased with increasing polymer concentration.

Finally, the abstracted linear slopes ( $c$  values) for all of the salts were correlated to the known hydration entropy ( $\Delta S_{\text{hydr}}$ ) and surface tension increment ( $\sigma$ ) values of the anions (Figure 10). As can be seen, the correlation between  $c$  values and surface tension increments is remarkably good for all of the chaotropes. Moreover, the correlation between  $c$  and  $\sigma$  extended quite well to the upper step,  $\nabla$ , of the two-step phase transition for the kosmotropes. On the other hand, the  $c$  values for initial step,  $\bullet$ , and first step,  $\blacktriangle$ , in the two-step transition of the kosmotropes were well correlated to the entropy of hydration,  $\Delta S_{\text{hydr}}$ , of the anions. In fact, the correlations were found to be equally good at all molecular weights. This confirms the notion that the specific anion effects involve surface tension and direct anion binding for the chaotropes (Figure 1a and b).<sup>5</sup>



**Figure 7.** LCSTs of 10 mg/mL PNIPAM with different molecular weights in a salt solution with chaotropic anions at concentrations from 0 to 1.0 M: (a) NaCl, (b) NaBr, (c)  $\text{NaNO}_3$ , (d) NaI, (e) NaSCN, and (f)  $\text{NaClO}_4$ . The curves were fit to eq 2.

**TABLE 3: Fitted Values of  $K_A$ ,  $B_{\max}$ , and  $c$  Abstracted from the LCST versus Salt Concentration Curves of PNIPAM at Different Molecular Weights in the Presence of Chaotropic Anions**

anion	$M_n$ (g/mol)	$B_{\max}$ (°C)	$K_A$ (M <sup>-1</sup> )	$c$ (°C/mol)
Cl <sup>-</sup>	6980	—	—	-10.3
	30 700	—	—	-10.4
	121 000	—	—	-10.4
	360 000	—	—	-10.3
Br <sup>-</sup>	6980	2.1	2.5	-8.1
	30 700	1.3	2.3	-7.5
	121 000	1.4	2.6	-7.4
	360 000	1.5	3.4	-7.5
NO <sub>3</sub> <sup>-</sup>	6980	1.7	2.8	-7.4
	30 700	1.0	2.0	-7.0
	121 000	1.1	2.9	-6.9
	360 000	1.4	3.1	-7.4
I <sup>-</sup>	6980	2.7	3.3	-2.9
	30 700	2.3	4.3	-2.8
	121 000	2.1	4.3	-2.9
	360 000	2.2	5.1	-3.2
ClO <sub>4</sub> <sup>-</sup>	6980	6.2	3.3	-11.8
	30 700	4.8	3.9	-11.9
	121 000	3.6	5.1	-12.6
	360 000	2.6	7.7	-12.7
SCN <sup>-</sup>	6980	6.0	2.0	-2.3
	30 700	4.8	2.8	-2.7
	121 000	3.1	4.3	-2.7
	360 000	2.6	6.9	-3.1

On the other hand, the kosmotropes modulate the LCST of PNIPAM through surface tension changes and the polarization of water molecules at all of the molecular weights tested (Figure 1a and c). It should be noted that the ability to polarize the hydration waters of the pendent amide groups is related to the entropy of hydration of the anions.<sup>5</sup>

## Discussion

**Kosmotropic Anions.** Table 1 provides the molecular weight dependence of the  $c$  values for the initial slope, as well as the upper and lower steps in the two-step phase transition at higher ion concentrations. As can be seen, the  $c$  values for the initial step and the lower step decrease with increasing molecular weight of PNIPAM. The effect is most pronounced for Na<sub>2</sub>CO<sub>3</sub> and Na<sub>2</sub>SO<sub>4</sub>. It is believed that the decoupling of the amide dehydration from the hydrophobic dehydration manifests itself in the formation of partially collapsed structures which scatter less light than fully collapsed PNIPAM.<sup>5</sup> Moreover, it has been reported that molten globular and crumpled coil structures may occur in PNIPAM of very high molecular weight ( $1 \times 10^7$  Da) even in the absence of salt for very dilute polymer solutions.<sup>35,36</sup> In the present experiments, such partially collapsed states are apparently stabilized at a lower molecular weight by sufficiently high concentrations of strongly hydrated ions. It should also be noted that the initial step and the lower step in the two-step phase transition were also sensitive to the concentration of PNIPAM. Namely, the phase transition temperature was more rapidly depressed in Na<sub>2</sub>SO<sub>4</sub> with increasing PNIPAM concentration (Table 2). In sharp contrast with these results, the slope for the upper step of the two-step phase transition was found to be independent of both the molecular weight and the concentration of PNIPAM.

From a molecular point of view, it appears that hydrogen bonding between amide moieties becomes more facile with either increasing molecular weight or concentration. The former presumably occurs intramolecularly, while the latter should be the result of increased intermolecular interactions. By contrast, intramolecular and intermolecular influences are not as important for hydrophobic hydration, which is associated with the upper

phase transition step. In other words, the hydrophobic moieties' propensity to remain hydrated appeared to be independent of whether another polymer chain was nearby or not.

The phase transition of PNIPAM as a function of molecular weight appears to be at least somewhat reminiscent of the denaturation of globular proteins. It has been shown that the denaturation of sufficiently small proteins can occur as an all-or-none transition. In other words, the protein molecule denatures completely or not at all at equilibrium.<sup>37</sup> However, this is not the case for larger proteins, whose thermally induced denaturation can typically be more complex. The absence of an all-or-none transition for larger proteins, of course, is normally interpreted in terms of their domain structures.<sup>38</sup> It has been reported for PNIPAM that the coil-globule transition apparently also occurs according to a "domain" mechanism.<sup>39</sup> Domains of PNIPAM appear to contain ~90 monomer units. In the present case, only the fraction with the lowest molecular weight ( $M_n$  6980 Da, ~60 monomer units) underwent an all-or-none transition. The other fractions behaved more like large proteins in the sense that the phase transition could be resolved into two independent steps corresponding to the dehydration of different structural domains.

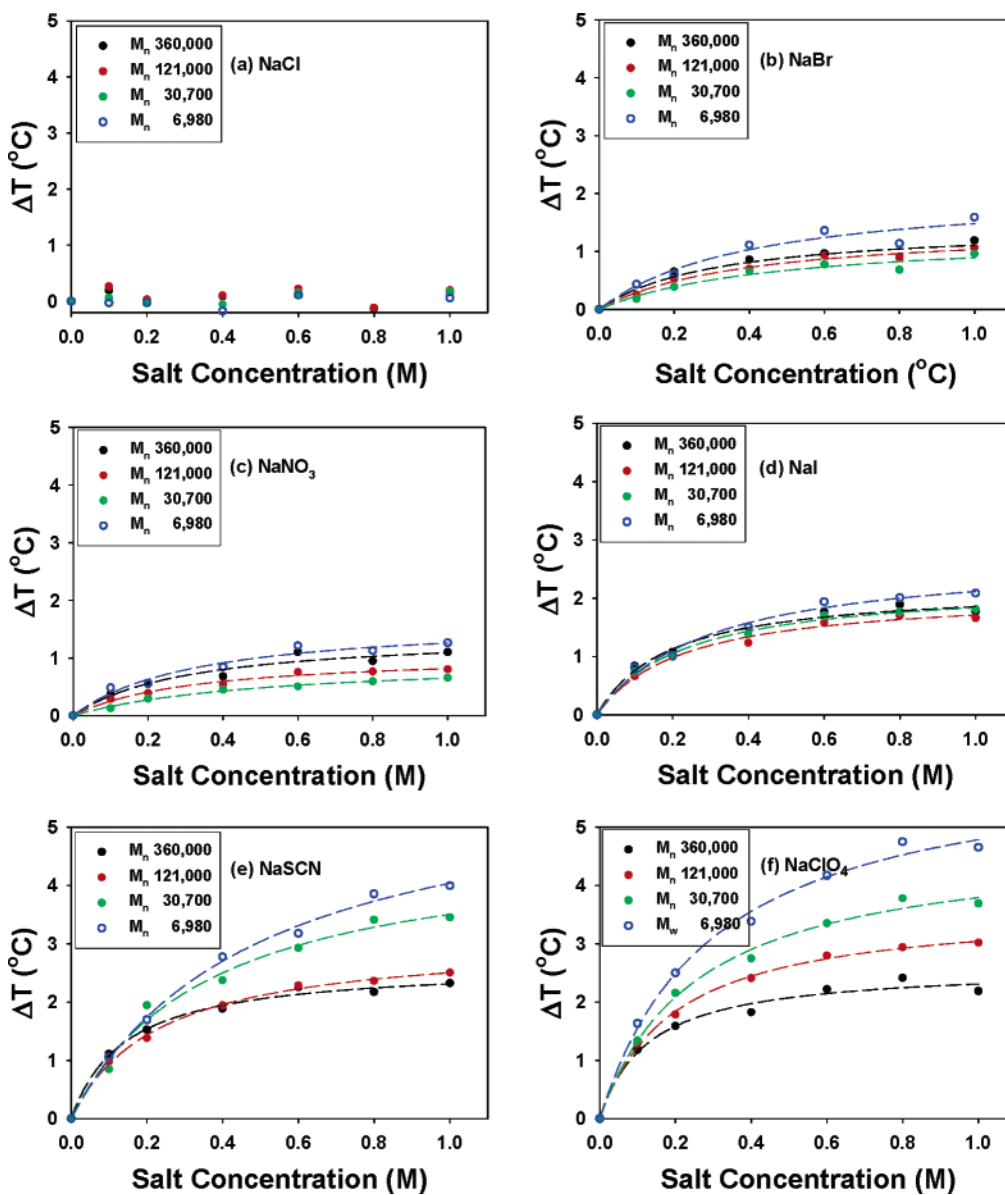
**Chaotropic Anions.** Chaotropic anions have two influences on the LCST of PNIPAM.<sup>5</sup> First, the surface tension of the hydrophobically hydrated moieties increases with added salt (Figure 1a). Second, the anions can shed their hydration shells and directly bind to the polymer (Figure 1b). As shown in Tables 3 and 4, the  $c$  values do not change with the molecular weight or concentration of PNIPAM, within experimental error, for any of the anions. This is consistent with the results obtained for the upper step of the two-step transition for the kosmotropic ions. On the other hand, direct ion binding (putatively involving the amide groups) does depend upon the molecular weight and PNIPAM concentration. For the most strongly chaotropic anions, ClO<sub>4</sub><sup>-</sup> and SCN<sup>-</sup>, a clear trend can be seen for  $B_{\max}$  and  $K_A$ . Namely, the value of  $B_{\max}$  falls with increasing molecular weight, while the value of  $K_A$  rises. The same trend is observed as a function of polymer concentration. The molecular weight effect is presumably caused by increased intramolecular interactions, while the concentration dependence is caused by intermolecular interactions. Such opposite behavior of  $K_A$  and  $B_{\max}$  could perhaps appear to be counterintuitive since one might expect a larger  $K_A$  value to be associated with a stronger salting-in effect (larger  $B_{\max}$ ). These results, however, can be understood by considering a multiple binding site model as described below.

For a PNIPAM chain having  $n$  apparently independent binding sites, the number of anions bound may range from 0 to  $n$ . Therefore, the average binding number,  $\nu$ , must lie between 0 and  $n$ . The value for  $\nu$  is related to the equilibrium association constant,  $K_A$ , by the following equation<sup>40</sup>

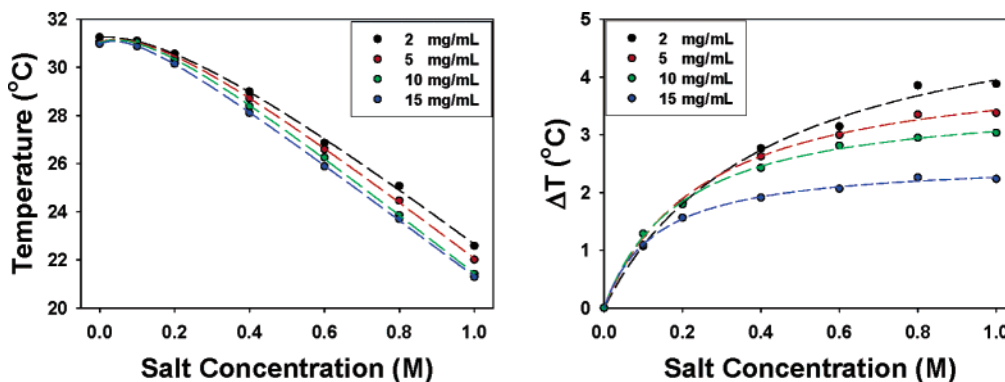
$$\nu = \frac{n[M]K_A}{1 + K_A[M]} \quad (3)$$

where  $[M]$  is the molar concentration of anion. In this model,  $n$  is related to  $B_{\max}$  in eq 2, while  $\nu$  is correlated to the actual temperature increase in the LCST observed at any given salt concentration due to direct anion binding. Accordingly, increasing  $n$  leads to a larger value for  $B_{\max}$ . The fraction of occupied sites,  $Y$ , can be found by rearranging eq 3 as follows

$$Y = \frac{\nu}{n} = \frac{[M]}{1/K_A + [M]} = \frac{[M]}{K_D + [M]} \quad (4)$$



**Figure 8.** The residual LCST data for the chaotropic anions from Figure 7 after removing the linear portion of the LCST versus concentration curves. The lines represent Langmuir isotherm fits to the data points.



**Figure 9.** (a) LCSTs of  $M_n$  121 000 Da PNIPAM with the polymer concentrations of 2, 5, 10, and 15 mg/mL in NaClO<sub>4</sub>. The curves were fit to eq 2. (b) The residual LCST data after removing the linear portion of the LCST versus concentration curves from Figure 9a. The lines represent Langmuir isotherm fits to the data points.

As can be seen from eq 4, a higher fraction of occupied sites is correlated with a larger value for  $K_A$ . This can also be expressed in terms of the equilibrium dissociation constant,  $K_D$ , where  $K_D = 1/K_A$ . Either way, stronger binding is related to a higher fraction of bound chaotropic anions.

As was also found for experiments with kosmotropic anions, the amide side chains should have increased opportunities to undergo hydrogen bonding with increasing chain length and polymer concentration. These effects should be intramolecular and intermolecular, respectively. Chain–chain interactions may

**TABLE 4: Fitted Values of  $K_A$ ,  $B_{\max}$ , and  $c$  Abstracted from the LCST versus  $\text{NaClO}_4$  Concentration Curves for a  $M_n$  121 000 Sample of PNIPAM at Different Polymer Concentrations**

anion	$M_n$ (g/mol)	concn (mg/mL)	$B_{\max}$ ( $^{\circ}\text{C}$ )	$K_A$ ( $\text{M}^{-1}$ )	$c$ ( $^{\circ}\text{C}/\text{mol}$ )
$\text{ClO}_4^-$	121 000	2	5.6	2.4	-12.6
		5	4.3	3.9	-12.4
		10	3.6	5.1	-12.6
		15	2.6	7.5	-11.9

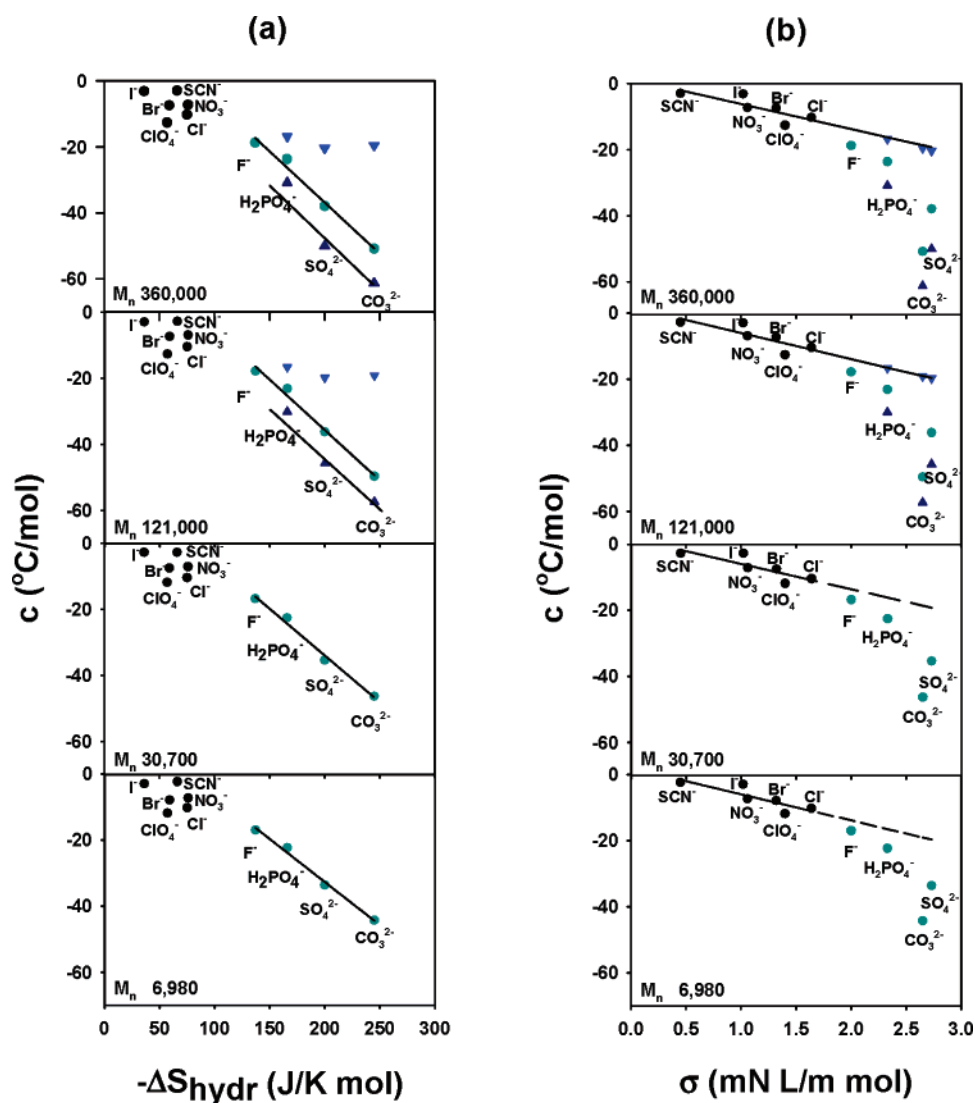
affect both the number of available binding sites as well as the apparent  $K_A$  value. In fact,  $n$  should be reduced on electrostatic grounds. This is because the proximity of the amides will be closer, making anion binding to neighboring amide moieties more costly. Such an effect will increase with increasing molecular weight or polymer concentration since these chain–chain interactions become more pronounced. Moreover, reducing  $n$  will also reduce  $B_{\max}$ . This is exactly what is found by increasing either the molecular weight or the concentration of PNIPAM for the strongest chaotropes (Tables 3 and 4). On the other hand, the apparent value of  $K_A$  should be increased for single anion binding to more closely interacting polymer chains.

This is, at least in part, related to a multivalency effect, whereby a given anion has greater opportunities to interact with the polymer. Analogous effects for protein–ligand binding have already been hypothesized upon increased interactions between the receptor sites.<sup>41</sup>

It should be noted that the reasoning for the direction of change for  $K_A$  and  $B_{\max}$  for the strongest chaotropes should also apply to the weaker chaotropes such as  $\text{Br}^-$ ,  $\text{NO}_3^-$ , and  $\text{I}^-$ . In fact, the apparent value of  $K_A$  increases in all cases with increasing molecular weight. The magnitude of  $B_{\max}$ , however, is down by a factor of 2–4 for the weaker binding anions at identical molecular weights compared with that of  $\text{ClO}_4^-$  and  $\text{SCN}^-$ . This makes the trends more difficult to discern within the signal-to-noise limitations of our experiments.

## Conclusions

The effects of molecular weight and concentration on the mechanisms of specific anion interactions with PNIPAM were investigated. It was found that amide dehydration in the presence of kosmotropes was more facile at higher molecular weight and



**Figure 10.** (a) Plot of hydration entropy of the anions versus  $c$  for PNIPAM with different molecular weights. (b) Plot of the surface tension increment of the anions versus  $c$  for PNIPAM with different molecular weights. Initial slopes are shown with filled gray-green circles; the first slope of the split transition is ( $\blacktriangle$ ), and the second slope of the split transition is ( $\blacktriangledown$ ) for kosmotropes; ( $\bullet$ ) is for the chaotropes. The lines shown represent least-squares fits to the data for the first slope of the split transition and for the slope of the initial transition for the kosmotropes. Hydration entropy values for the anions were obtained from ref 42. The surface tension increments of the anions were obtained from ref 43, except for  $\text{H}_2\text{PO}_4^-$ , which was from ref 1, and  $\text{CO}_3^{2-}$ , which was from ref 44.



polymer concentration. This was consistent with increased chain–chain interactions of both intramolecular and intermolecular origin. Moreover, the apparent association constants with the chaotropes increased as a function of molecular weight and PNIPAM concentration on largely similar grounds. In stark contrast with this, the slope of the LCST change versus salt concentration for hydrophobic hydration of the polymer was not altered by modulation of either the polymer's molecular weight or concentration in the presence of either kosmotropes or chaotropes.

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**Supporting Information Available:** Representative light-scattering curves are provided. A note on the effects of the ions on the pH of the polymer solutions is also provided. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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