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# Probing weak intermolecular interactions in self-assembled nanotubes

*Laurent Bouteiller<sup>1\*</sup> and Paul van der Schoot<sup>2\*</sup>*

<sup>1</sup> UPMC Univ Paris 06, UMR 7610, Chimie des Polymères, F-75005 Paris, France, and CNRS, UMR 7610, Chimie des Polymères, F-75005 Paris, France

<sup>2</sup> Group Theory of Polymers and Soft Matter, Technische Universiteit Eindhoven, P.O. Box 513, 5600 MB Eindhoven, The Netherlands, and Instituut voor Theoretische Fysica, Universiteit Utrecht, Leuvenlaan 4, 3584 CE Utrecht, The Netherlands

laurent.bouteiller@upmc.fr; p.p.a.m.v.d.schoot@tue.nl

**ABSTRACT.** Extreme confinement affects the physical properties of fluids but little quantitative data is available. We report on studies of a bisurea compound that self-assembles into nanotubes to probe solvent confinement on the Å scale. By applying a statistical model to calorimetric data obtained on solvent mixtures, we show that the thermodynamic stability of the nanotubes is an extremely sensitive function of the solvent composition because solvent interactions inside and outside the nanotubes are different. We are able to measure energetic effects as small as 0.01kT and relate them to the differences in molecular structure of the solvents.

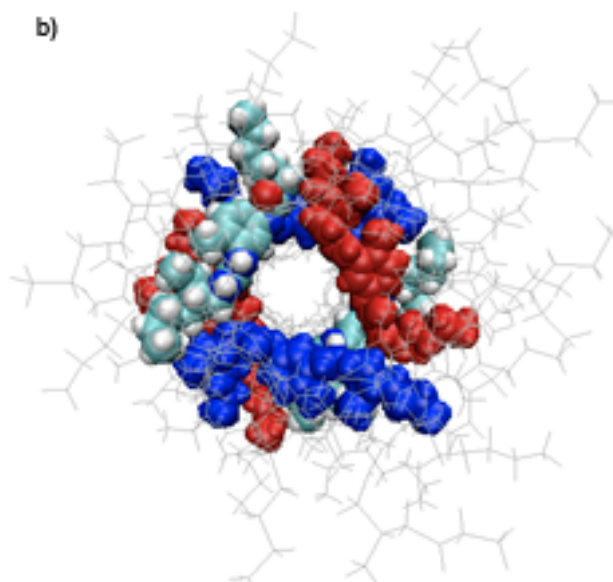
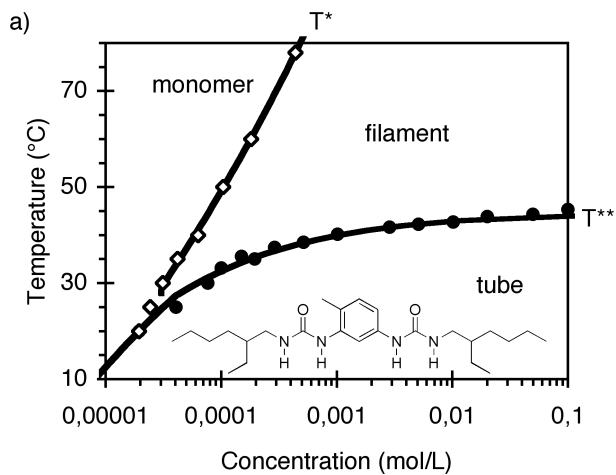
**KEYWORDS.** self-assembly; nanotube; confinement, supramolecular.

## **Introduction**

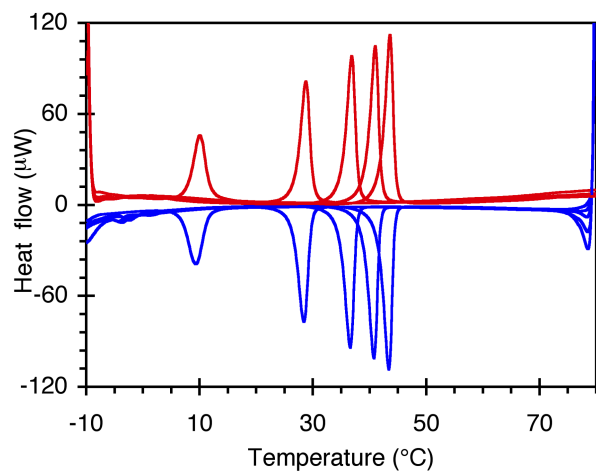
Physical properties of fluids are known to change under the influence of confinement and more so the stronger the confinement: fluids in carbon nanotubes a few nanometers across exhibit lower friction

than in micron-sized capillaries,<sup>1</sup> solvent mixtures in contact with sufficiently thin capillaries fractionate,<sup>2</sup> and phase transitions change order if trapped in Vycor glass if the pore size is small enough.<sup>3</sup> This has ramifications for a wide variety of fields, including geology,<sup>4</sup> biology,<sup>5,7</sup> chemical engineering<sup>8</sup> and supramolecular chemistry.<sup>9</sup> Still, little is known let alone understood about the effects of confinement on fluids in tubes that are so thin that only a single file of fluid molecules fit in.<sup>10</sup> In this paper, we report on studies of a bisurea compound that self-assembles into nanotubes in a large number of solvents in order to probe solvent confinement on the Å scale. Self-assembled nanotubes are ideally suited for this, because their thermodynamic stability is an extremely sensitive function of the solvent composition. By applying a statistical model to experimental data obtained on twelve binary solvent mixtures, we find that solvent interactions inside the nanotubes are different from those in the bulk. For almost all mixtures tested, interactions between unlike solvent molecules are less favourable inside the nanotubes than outside. The coupling between nanotube self-assembly and intermolecular interactions with (and between) encapsulated solvent molecules also provides an unprecedented insight into how small differences in intermolecular interactions have a large impact on supramolecular assemblies and highlights the important role of solvent therein, a role that so far has not received the attention that it merits in the experimental and theoretical literature.<sup>11</sup>

The compound we focus attention on belongs to a class of bisureas known to self-assemble, through inter-molecular hydrogen bonding, into filaments below a polymerization temperature  $T^*$ , and into narrow tubes for temperatures  $T \leq T^{**} \leq T^*$ .<sup>12,13</sup> While the polymerisation of monomers into filaments is cooperative, the filament-tube transition at  $T^{**}$  is even sharper and resembles a phase transition. It is characterized by a peak in the heat capacity and can be pinpointed accurately as a function of concentration, solvent type and so on.<sup>14,15</sup> An example of a state diagram is given in Figure 1. It turns out that the filament-tube transition is extremely sensitive to the solvent in which the bisureas are dissolved. The polarity of the solvent plays an important role, but more significant is the size and shape of the solvent molecules.<sup>16</sup> Molecular modeling and host-guest studies have shown that the inner diameter of the self-assembled tubes measures at most 7 Å, suggesting there is barely enough space to allow for a single file of solvent molecules.<sup>16,17</sup> Molecules wider than the optimal radius distort the self-assembled nanotube and hence destabilize it. Also, the longer the solvent molecules, the more orientational entropy is lost upon their encapsulation. Clearly, interactions between pairs of solvent molecules in- and outside a tube must also be different because confinement frustrates their optimal positioning.



**Figure 1.** (a) Chemical structure of bisurea EHUT and its phase diagram in toluene.<sup>15</sup> (b) Optimized nanotube structure determined by molecular mechanics and dynamics (6 individual molecules are represented with a different colour).<sup>13</sup>

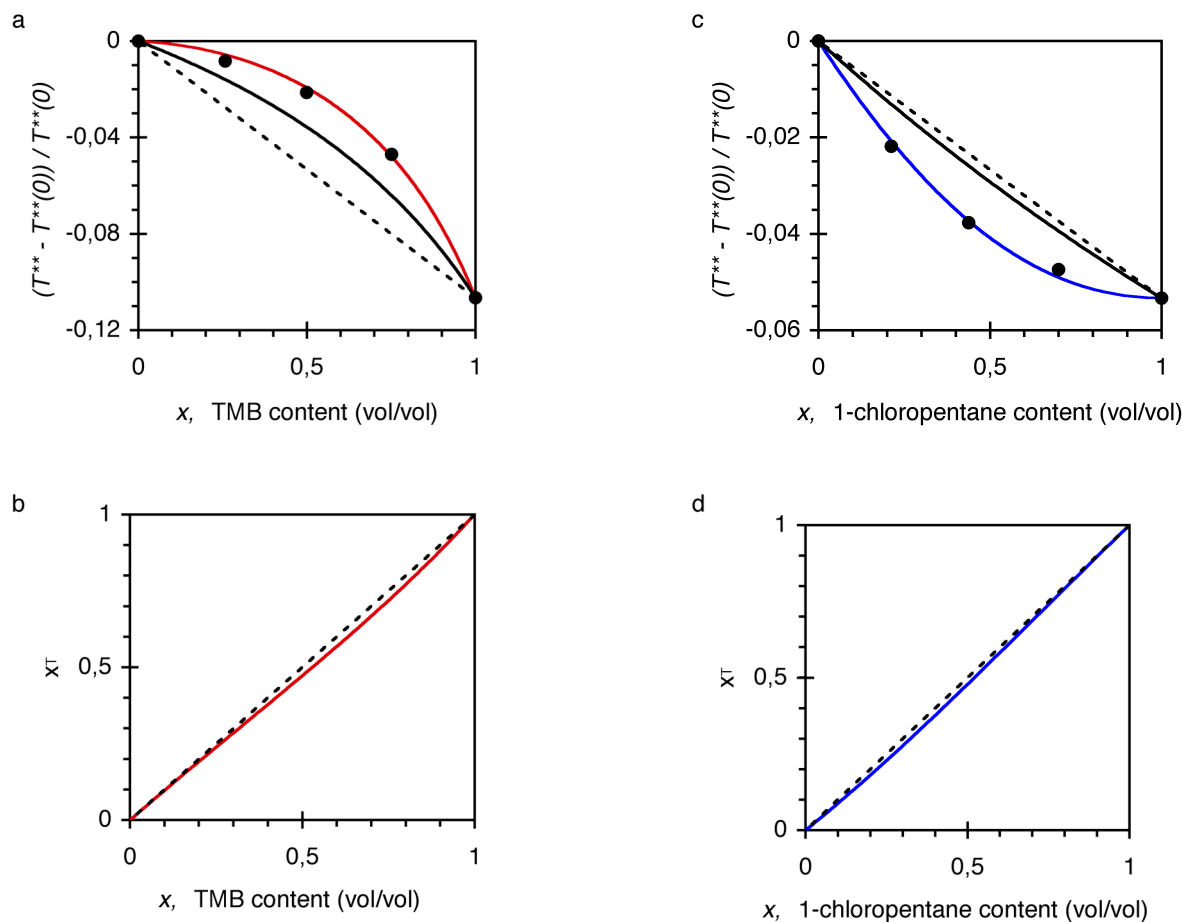


**Figure 2.** Calorimetric characterization of the tube to filament transition. DSC thermograms (heating in red; cooling in blue) for solutions of EHUT in toluene/1,3,5-trimethylbenzene (TMB) mixtures. The molar composition of the solvent is: 0/100, 21/79, 43/57, 70/30, 100/0, from left to right. [EHUT] = 0.01 mol/L.

By investigating how the filament-tube transition temperature varies with composition in a variety of solvent mixtures, and by applying a theoretical model that describes the impact of solvent on the stability of self-assembled nanotubes, we are able to probe extremely small changes in interaction free energies between the various components caused by the confinement. The competition between tube-solvent and solvent-solvent interactions explains the strongly non-linear dependence of the filament-tube transition temperature  $T^{**}$  on the bulk solvent composition. Competition between self-assemblies amplifies small differences in interactions that otherwise cannot straightforwardly be measured nor obtained from computer simulation.<sup>14</sup>

## Results and Discussion

Figure 2 shows Differential Scanning Calorimetry (DSC) traces measured for bisurea EHUT solutions in toluene/1,3,5-trimethylbenzene (TMB) mixtures. The endothermic transition is characteristic for the transition from the tube to the filament structure. For the twelve binary solvent mixtures studied, the heating and the cooling runs are symmetrical confirming the reversibility of this transition. The transition temperature,  $T^{**}$ , was taken at the maximum of the endothermic peak and is plotted versus solvent composition in Figures 3a and S1. The variation is clearly non-linear with a positive departure of as much as 10°C from the expected linearity if the solvent inside and outside the tubes would behave identically. Of course, a change of solvent may affect the nanotube precise structure, which is expected to have a direct influence on the transition temperature. However, a conformational change induced by a direct solvent-nanotube interaction affects only the portion of the tube in direct contact with the solvent molecule. Therefore, direct solvent-nanotube interactions cannot explain the non-linearity observed.



**Figure 3.** Influence of bulk solvent composition on tube stability and on solvent composition in the tubes. Normalized transition temperature ( $T^{**}$ ) versus bulk solvent composition ( $x$ ) for EHUT in toluene/TMB mixtures (a) or heptane/1-chloropentane mixtures (c) ( $T^{**}(0) = T_1^{**}$  and  $T^{**}(1) = T_2^{**}$ ). Drawn lines: curve fit with  $J = h_J = -0.025$  (a: red),  $J = h_J = 0.016$  (c: blue) or  $J = h_J = 0$  (black). Calculated volume fraction of TMB (b) or 1-chloropentane (d) in the tubes versus bulk fraction.

An obvious explanation for the non-linear behaviour observed is fractionation of the solvent. The bisurea tubes are more stable in toluene than in TMB as judged from the higher  $T^{**}$ , therefore one expects the tubes to encapsulate toluene preferentially implying its concentration inside the tubes to be larger than outside. This could explain the positive deviation of the curve. In order to test this hypothesis, we measured the composition dependence of the bisurea transition temperature for diverse solvent mixtures (Table 1). Remarkably, for some mixtures a *negative* deviation was measured, e.g., heptane and 1-chloropentane (Figure 3c). If the negative deviation were the consequence of solvent fractionation, then this would mean that the tubes favour the solvent in which the tubes are the least

stable. Arguably, this is unphysical and shows that solvent fractionation cannot be the sole reason for the non-linear variation of the transition temperature.

**Table 1.** Values for the parameters deduced from the composition dependence of  $T^{**}$  for EHUT in mixed solvents (compositions probed: approximately 0/100, 25/75, 50/50, 75/25, and 100/0).

entry	solvent 1	solvent 2	$T_1^{**}$ <sup>a</sup> (°C)	$T_2^{**}$ <sup>a</sup> (°C)	$h^b$ ( $k_B T$ )	$\Delta h^c$ ( $k_B T$ )	$\Delta \varepsilon^d$ ( $k_B T$ )	$T\Delta s^e$ ( $k_B T$ )	$J^f$ ( $k_B T$ )
1	toluene	TMB	43.5	9.8	-2.0	0.9	0.12	0.8	-0.025
2	p-xylene	TMB	49.0	9.8	-2.0	0.9	0.13	0.8	-0.020
3	octane	heptane	75.4	72.9	-1.5	0.2	0.01	0.2	0.001
4	terbutylbenzene	p-xylene	50.9	49.0	-2.1	0.1	0.01	0.1	0.001
5	terbutylbenzene	toluene	50.9	43.5	-2.1	0.1	0.05	0.1	0.002
6	p-xylene	toluene	49.0	43.5	-2.0	0.0	0.03	0.0	0.004
7	terbutylbenzene	TMB	50.9	9.8	-2.1	1.1	0.14	0.9	0.010
8	heptane	p-xylene	72.9	49.0	-1.4	-0.6	0.14	-0.8	0.010
9	heptane	1,9-decadiene	72.9	65.9	-1.4	-0.1	0.03	-0.2	0.010
10	heptane	toluene	72.9	43.5	-1.4	-0.6	0.17	-0.8	0.015
11	heptane	1-chloropentane	72.9	54.4	-1.3	-0.3	0.08	-0.4	0.016
12	heptane	TMB	72.9	9.8	-1.4	0.3	0.20	0.1	0.035 (0.04) <sup>g</sup>

<sup>a</sup> transition temperature for EHUT, measured by DSC; uncertainty  $\pm 0.2^\circ\text{C}$ ;

<sup>b</sup> transition enthalpy for EHUT in solvent 1, measured by DSC; uncertainty  $\pm 0.15 k_B T$ ;

<sup>c</sup> difference in transition enthalpy for EHUT in solvents 1 and 2 ( $\Delta h = h_2 - h_1$ ), measured by DSC; uncertainty  $\pm 0.3 k_B T$ ;

<sup>d</sup> excess free energy; uncertainty  $\pm 0.03 k_B T$ ;

<sup>e</sup> uncertainty  $\pm 0.33 k_B T$ ;

<sup>f</sup> interaction parameter between solvent 1 and 2, deduced by fitting the DSC data of mixtures, assuming that  $h_J = J$ .

<sup>g</sup> interaction parameter between solvent 1 and 2, deduced by fitting the DSC data of mixtures, assuming that  $h_J = 0$ .

In order to provide an explanation for our observations, we apply a statistical theory based on the law of mass action and a model for the interaction of the solvent mixture with the assemblies, described in the Supporting Information. There are seven model parameters, five of which are fixed by DSC of the solutions in the pure solvents 1 and 2, see Table 1. These include the transition temperatures  $T_1^{**}$  and  $T_2^{**}$  in the two solvents, and the difference between binding enthalpies  $h_1 \equiv h$  and  $h_2 \equiv h + \Delta h$  of single bisureas in tubes and in filaments.<sup>18</sup> The difference of the solvent-contribution  $\Delta\varepsilon$  to the excess free energy of binding of the monomers in a tube in the two types of solvent is also fixed by the pure solvent data, because the transition temperature difference  $T_2^{**} - T_1^{**}$  is proportional to  $\Delta\varepsilon$ .<sup>19</sup> Only two unknowns are left: a free energy  $J$  and its associated enthalpy  $h_J$ , describing the effect of confinement on the interaction between the two types of solvent. If  $J > 0$ , unlike contacts are penalized by the confinement in the nanotube whilst if  $J < 0$  the reverse is true.

Because  $J \equiv h_J - T_1^{**} s_J$ , we for simplicity consider the enthalpy-dominated case where  $s_J = 0$  and the entropy-dominated one where  $h_J = 0$ . Hence, in the curve fitting to our experimental data we have a single adjustable parameter,  $J$ , and examine the two cases  $h_J = 0$  or  $h_J = J$ . Figures 3a and 3c show curve fits for two solvent mixtures. Irrespective of whether  $h_J = 0$  or  $J$ , a near perfect fit is obtained suggesting our results are robust albeit that we cannot discriminate between enthalpic and entropic confinements effects.

The curve with a positive deviation (Figure 3a) gives a good agreement with a *negative* value for  $J$ , while the curve with a negative deviation (Figure 3c) gives a *positive* value for  $J$ . We applied our model to the data obtained for all twelve solvent mixtures investigated: all of them can be fitted accurately requiring a single adjustable parameter  $J$ . The numerical values that we find (Table 1), are mere fractions of the thermal energy, yet setting  $J = 0$  does not describe the data adequately (see Figures 3a and 3c). As before, setting  $h_J$  equal to 0 or  $J$  has no discernible effect, except for mixtures of the solvents heptane and TMB. For this single case, we therefore have a larger uncertainty for  $J$ :  $0.035 \leq J/k_B T \leq 0.04$ . Still, the robustness of the model is remarkable; all the data can be explained on the basis of the strength of interactions between two unlike solvent molecules confined in one spatial dimension relative to that in three. Apparently, unlike contacts are penalised under confinement for all mixtures, except TMB/toluene and TMB/p-xylene. That the magnitude of  $J$  that we find is small, at



most  $0.04 k_B T^0$  seems reasonable, given that the solvents selected do not possess strongly interacting groups and in bulk seem to behave like ideal mixtures.<sup>21</sup>

In spite of this, it is possible to detect a clear trend in our results. The four mixtures in which the smallest interaction energy is found ( $|J| \leq 0.004 k_B T$ ) involve similar solvents: two linear alkanes (entry 3) or two aromatic hydrocarbons with a single substituent (entries 4 through 6). All the mixtures for which  $|J| \geq 0.01 k_B T$  concern less symmetrical systems: (i) a linear alkane and an aromatic hydrocarbon (entries 8, 10 and 12); (ii) a linear alkane and an unsaturated or chlorinated alkane (entries 9 and 11); (iii) an aromatic hydrocarbon with a single substituent and an aromatic hydrocarbon with substituents in the 1, 3 and 5 positions (entries 1, 2 and 7). The correlation between the value of the interaction parameter  $J$  and the similarity of solvent molecules suggests that the main physical effect is captured by our model.

Table 1 also presents our findings for the difference between the solvent contributions to the excess free energy of binding of the monomer in a tube,  $\Delta\varepsilon$ , and its breakdown in the contributions from the enthalpy  $\Delta h$  and entropy,  $T\Delta s = \Delta h - \Delta\varepsilon$ . The enthalpies are up to ten times larger than the free energies. This shows the important role that solvent entropy plays in the driving force for the supramolecular assembly of molecules.

We now return to the fractionation of the solvents in- and outside the nanotubes. Results of our calculations in Figures 3b and 3d show that it must be insignificant, at most a few %. This is to be expected because the interaction free energies we find are small, see Table 1. Also, the quasi 1D character of the solvent in the tubes makes it sensitive to fluctuations that diminish any tendency toward fractionation. This confirms that fractionation cannot explain our findings and must be due to the effects of confinement on the interactions between the solvent molecules in the tubes.

## Conclusion

In conclusion, the interaction free energy between neighbouring solvent molecules placed in a single file configuration, caused by confinement in a nanotube, differs from that in a bulk mixture. Although that for all mixtures tested the difference is very small, we are able to conclude that the larger the difference in the molecular structure of the solvent molecules, the larger the influence of confinement on the net interaction between them. In most but not all mixtures interactions between unlike species are penalised under conditions of extreme confinement. It is not clear why this is so, and hopefully our

experiments inspire (ab initio) calculations of intermolecular interactions and help increase our understanding of the consequences of extreme confinement on molecules.

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**Supporting Information Available.** Experimental procedures and description of the theoretical model. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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- (19) See details in Supporting Information.
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# Table of Content Graphic

