Scaled Effective Solvent Method for Predicting the Equilibrium Ensemble of Structures with Analysis of Thermodynamic Properties of Amorphous Polyethylene Glycol–Water Mixtures

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Supporting Information

ABSTRACT: Water-soluble polymers such as polyethylene glycol (PEG) are critical components of industrial processes ranging from drug delivery to water purification. However, the understanding of the microscopic structure of these polymers in water and of the thermodynamics of the mixtures is limited because available experimental techniques (such as SLS and SANS) give little information about conformations and provide even the radius of gyration only in the dilute limit ($< \sim 5$ wt % PEG). Computer simulations employing Monte Carlo (MC) and molecular dynamics (MD) techniques can provide an atomistic molecular structure; however, such approaches have difficulties in predicting the equilibrium polymer configurations of high-molecular-weight polymers at normal densities and in obtaining entropies and free energies directly from the MD. Here, we develop the scaled effective solvent (SES) method to predict the equilibrium ensemble of polymer configurations, which we illustrate for the case of a 20 kDa PEG (455 monomers) at a 25 wt % PEG aqueous solution (3339 waters per PEG chain). We evaluate the free energy and entropy of the members of this ensemble including explicit water, validating that it leads to average sizes (R_g) observed experimentally and that all members of the ensemble have favorable free energies. With the SES method validated to provide well-equilibrated polymer chains in water, it should be useful for predicting ensembles of polymer chains in polymer melts and in solvents.

1. INTRODUCTION

Understanding the nature of polymer–solvent interactions is quite important in designing optimal systems for industrial applications ranging from drug delivery to water purification and fuel cell electrolytes. Such design is hampered by limitations in experimental characterization, making it most difficult to develop an atomistic understanding beyond the limits of very dilute solutions. In order to provide this atomistic understanding, we are developing new methodologies, which we illustrate here for the case of polyethylene glycol (PEG) water mixtures, a water-soluble polymer with wide-ranging applications. Indeed, interactions of PEG with water resemble that of a protein¹ so that such studies provide insights into the more complicated interactions and function of proteins and enzymes. In particular, the substantial ability of PEG mixtures to absorb and transport water molecules² is of great importance in commercial and technological applications, such as pharmaceutical drugs delivery,^{3–7} cosmetics,^{8,9} and functioning fuel cells.¹⁰

To determine the atomic-level structure of PEG solution and the interaction of PEG with water molecules, a number of experimental methods have been applied, including dynamic light scattering (DLS),^{11–13} static light scattering (SLS),^{12,14,15} Raman spectroscopy,^{16,17} nuclear magnetic resonance

(NMR),^{18–20} infrared spectroscopy (IR),¹⁹ and small-angle neutron scattering (SANS).^{21,22} These methods provide various levels of macroscopic structural information, such as the overall size (radius of gyration, R_g) of the PEG chains and local clustering. However, such measurements are indirect and ensemble-averaged, providing little information about atomistic structures. In addition, structural characterizations based on SLS or SANS assume the Guinier condition that is valid only in the dilute regime ($< \sim 5$ wt %), where polymer–polymer interactions are negligible. Consequently, experimental characterizations of polymer structures in the semidilute regime (20–35 wt %)²³ are rare. Other important properties such as osmotic behavior of these mixtures are difficult to predict because the contribution of the second virial term to osmotic pressure is significant in this semidilute regime.

In contrast, molecular dynamics (MD) simulations can provide atomistic structures and thermodynamics for interpretation of experimental studies. However, the complexity of sampling the polymer configuration space for 20 kDa PEG

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(PEG-20 kDa) with its 455 monomer units (1365 independent torsions) is huge and becomes even more difficult when considering the water or other solvent explicitly. Thus, previous MD studies of PEG structures in water have generally been limited to low-molecular-weight polymers ($< \sim 1$ kDa^{24–27} or 22 monomer units). For practical applications, the most interesting sizes for PEGs are much larger, with molecular weights of 40–100 kDa commonly used in biomedical and biotechnical applications. In this regime, experimental limitations have not resolved such questions as whether the PEGs in water are aggregated under various conditions.^{11,28}

We report here the scaled effective solvent (SES) computational method designed to sample a sufficiently wide range of polymer conformations in solution in which we can be confident that the properties of fully equilibrated polymer chains are captured. In the SES method, we start with isolated chains but include the effect of the solvent and other polymer chains being ignored by scaling the intrachain nonbonding interactions. This mimics the effect of solvent and other molecular chains in denser polymer systems, which counterbalance the attractiveness of intrachain interactions. We then combine this SES procedure with a new first-principles-based force field developed here to describe the intermolecular interaction between water and PEG to predict the atomistic structures of PEG-20 kDa for the most interesting PEG concentration in an aqueous solution of 25 wt % (semidilute), a regime where it is difficult to extract structural data from experiment.

We then analyze the entropy and free energy of these mixtures using the two-phase thermodynamic (2PT) method.²⁹ We find an equilibrium ensemble of polymer chains with a wide range of sizes, $R_g = 34.86$ – 78.77 Å, each with similar free energies, indicating that this distribution of chains represents the equilibrium ensemble. The final thermodynamically averaged size is $R_g = 53.98$ Å for 25% PEG in water at 300 K. This is in agreement with experiment in the dilute limit, $R_g = 65$ Å, and with the $R_g = 56.68$ Å, assuming a model of segments freely jointed at the O atoms with a fixed C–O–C angle.

2. COMPUTATIONAL DETAILS

2.1. First-Principles-Based Force Field. To obtain an accurate description of water molecules interacting with the PEG

Figure 1. Chemical structure of a PEG-20 kDa chain end-capped with hydroxyl (–OH) groups, which consists of 455 repeating monomers.

backbone, we developed a new force field (FF) by modifying the DREIDING³⁰ generic FF to reproduce the quantum mechanics (QM) energies and structures for the hydrogen bond interaction of water molecules with the ether oxygen atoms of the PEG backbone. Using dimethyl ether (DME) to represent a repeating unit of PEG interacting with H₂O, we determined these interactions using the M06-2X density functional³¹ with the Dunning correlation-corrected triple- ζ basis sets (cc-PVTZ++³² in Jaguar 7.6 program³³). Figure S1a (Supporting Information (SI)) shows the optimized structure from QM calculations (global minimum configuration) along with two other structures optimized while constraining the point group symmetry to remain C_{2v} (Figure S1b and S1c, SI). We fitted these QM results to a Morse-type off-diagonal van der Waals term and a

DREIDING-like explicit three-body hydrogen bonding term for the interactions of the ether oxygen atoms in the PEG backbone with the hydrogens in water molecule. Here, we used the 12–10 Lennard-Jones functional term for the radial term.

Figures S1 and S2 (SI) show that the FF describes well the intermolecular interactions from QM, including the global minimum configuration (Figure S1a, SI). The optimum FF parameters are listed in Table S1 (SI). The partial charge distributions (q) (Table S2, SI) of the PEG molecule were determined to fit the electrostatic potential (ESP) charges derived from QM calculation for the 1,2-ethanediol molecule (which represents the PEG model with a repeating unit; $n = 1$).

For the water–water interaction, we used the flexible three-centered (F3C) water model³⁴ with a $-0.82e^-$ partial charge on oxygen ($q_{O(wat)}$) and a $0.41e^-$ partial charge on hydrogen ($q_{H(wat)}$). The spring constant for the OH bond is 500 kcal/mol/Å², with an equilibrium bond length of 1.0 Å, and the spring constant for HOH angle is 120 kcal/mol/rad², with an equilibrium angle of 109.47°.

2.2. MD Simulations. As a prototypical PEG, we considered the 20 kDa molecular weight linear polymer, consisting of 455 repeating monomers (terminating both end groups with hydroxyl, –OH), as shown in Figure 1.

As a representative case for the semidilute regime, we focused on the detailed structures and energetics of PEG in aqueous solution with a polymer concentration of 25 wt %. The simulation cell for a 25 wt % PEG aqueous solution consisted of a single PEG-20 kDa chain (3188 atoms) solvated by 3339 water molecules, leading to 13 205 atoms in the simulation cell. We carried out MD simulations using the large-scale atomistic modeling massively parallelized simulation (LAMMPS) code.³⁵ Here, the isothermal damping constant for Nosé–Hoover NVT calculations was 100 fs, and the isobaric damping constant for Nosé–Hoover NPT calculations was 1000 fs.

3. RESULTS USING THE SOLVENT-INCLUDED COHESIVE ENERGY DENSITY (SI-CED) METHOD

Our system with the 1365 independent torsions leads to $\sim 3^{1365} \sim 10^{651}$ possible conformers, not counting the solvent. For

Figure 2. Atomistic structure of a PEG-20 kDa with $R_g = 36.57$ Å in a PEG aqueous solution with 25 wt % of PEG built using SI-CED at the final full density, $\rho = 1.05$ g/cm³ (comparable to the experimentally measured density, $\rho^{\text{exp}} = 1.04$ g/cm³). White, red, and green atoms are hydrogen, oxygen, and carbon atoms, respectively. The periodic box is shown as a blue solid line. The equilibrated structure by SI-CED is partially aggregated due to the self-solvation effect resulting from substantial intrachain interaction during vacuum dynamics (no water) and has much higher energy than the sampled structures with our new procedure (see Figure 9). Therefore, we conclude that the SI-CED is not a proper procedure for sampling high-molecular-weight polymer structures dissolved in a solvent.

Last 5 ns equilibrated structures ($f=0.3$ and $\rho=0.01$ g/cm³)

Figure 3. The $f = 0.3$ SES case. Structures of a PEG-20 kDa chain with $\rho = 0.01$ g/cm³ sampled over the last 5 ns NVT dynamics at $T =$ (a) 100, (b) 300, (c) 600, (d) 1000, and (e) 1200 K. Over the temperature range from 300 to 1200 K, the last 5 ns of equilibrated PEG-20 kDa conformations all show random coil-like behavior in which the backbone of the polymer moves randomly in three-dimensional space, while at 200 K and below, the PEG structures maintain a lamellar shape. White, red, and green atoms are hydrogen, oxygen, and carbon atoms, respectively. The periodic box is shown as a blue solid line.

such a dense polymer melt or solution, MD is far too slow to allow the system to equilibrate fully in practical time scales. Consequently, Monte Carlo (MC) or simulated annealing (SA) methods are generally used in such systems.^{36,37} However, for long polymer chains at normal density, the MC growth process can lead to nonuniform conformations due to exclusion effects as the full length of the growing polymer is reached. To overcome this limitation, Goddard and co-workers³⁸ developed the cohesive energy density (CED) method, which involves

- (1) growing the polymer at half of the expected density, 0.5ρ ,
- (2) compression of the simulation cell from 0.5ρ to 1.1ρ with five quench–anneal cycles [300–706 K ($\sim$ twice the Flory temperature; 353 K³⁹)] using NVT, and

- (3) relaxation of the final structure from step 2 using NPT MD for 0.25 ns to obtain the final structure.

Indeed, CED has been applied to predict Hildebrand solubility parameters,^{40–42} the structures of complex polymers such as Nafion⁴³ and Dendrimer-modified electrolytes,⁴⁴ and thermodynamics⁴⁵ using MD simulation.

A complication with the CED method is how to incorporate the solvent. In previous studies^{43,46} the amount of water was small, and it was sufficient to add them in at the 50% density level, right after the “polymer growth step in vacuum”. This is denoted as the solvent-included CED; SI-CED method.

An example of the structures built from SI-CED is shown in Figure 2 (25 wt % PEG at the final full density, $\rho = 1.05$ g/cm³).

Last 5 ns equilibrated structures ($f=1.0$ and $\rho=0.01$ g/cm³)

Figure 4. The $f = 1.0$ SES case. Structures of a PEG-20 kDa chain with $\rho = 0.01$ g/cm³ sampled over the last 5 ns NVT dynamics at $T =$ (a) 100, (b) 300, (c) 600, (d) 1000, and (e) 1200 K. Over the temperature range from 900 to 1200 K, the last 5 ns of equilibrated PEG-20 kDa conformations all show random coil-like behavior in which the backbone of the polymer moves randomly in three-dimensional space, while at 800 K and below, the PEG structures maintain a lamellar shape. White, red, and green atoms are hydrogen, oxygen, and carbon atoms, respectively. The periodic box is shown as a blue solid line.

We see that for 25 wt % PEG in water, SI-CED leads to PEG chains containing nonuniform polymer globules. This occurs because of the strong intrachain interaction (i.e., self-solvation effect) during the polymer growth procedure in vacuum prior to water insertion. These partially globular conformations from SI-CED bias the sampling toward the initial self-solvated conformation, which survives even after compression/expansion cycles using the temperature annealing procedures in CED.

On the basis of these results, we concluded that the SI-CED strategy is not appropriate for a broad sampling of high-molecular-weight polymer structures immersed in a solvation box. In fact, we show below that the PEG–water mixture

sampled and equilibrated with SI-CED leads to a much higher energy than the structures built with our new procedure and that the size of the SI-CED chain is at the small size limit of our final distributions (vide infra).

4. SES-CED METHOD

4.1. SES Concept. In order to develop an efficient but unbiased sampling of polymer configurations for high degrees of solvation, we developed the SES-CED method. SES-CED samples the polymer chain conformations of an isolated chain but with reduced nonbonding interactions (van der Waals, hydrogen bonds, and electrostatic) scaled by a factor f to account

Figure 5. The scaled standard deviation of R_g , $\delta R_{g-sc} = (\text{standard deviation of } R_g) / (\text{average } R_g)$, for $f = 0.3$ over the last 5 ns NVT dynamics for each temperature ranging from 100 to 1200 K using a large simulation cell with $\rho = 0.01 \text{ g/cm}^3$. We find that low temperature leads to a globular or lamella structures with small δR_{g-sc} ($< \sim 0.01$), and high temperature leads to a random coil (R-coil) structure with $\delta R_{g-sc} > \sim 0.13$. At the highest temperature, the value of δR_{g-sc} is similar to the value expected for an infinite Gaussian coil, 0.26. To determine a precise θ temperature (T_t) to serve as the boundary between these two regimes, we used the hyperbolic tangent function, $\delta R_{g-sc}(T) = a[\tanh\{(T - T_t)/b\} + 1]$, where T_t is indicated by the blue point.

for the solvent being ignored (the effective solvent). The idea is that in the real polymer system, there are attractive interactions of the polymer chain with solvent and other polymer chains ignored in the single-chain vacuum representation, while the intrachain interactions are included explicitly. This case ($f = 1.0$) favors globule or lamella formation. Using a reduced f for the isolated single-chain calculations reduces the bias of the intrachain interactions, allowing chain structures to more closely resemble the actual chain conformations in a polymer melt or a water–polymer mixture while still allowing rapid and efficient sampling of the polymer chain over all possible conformations and sampling the multiple basins in the potential energy surface.

4.2. Phase Diagram of Polymer Chain Conformations (As a Function of f and T). In order to choose a proper scaling factor, f , we performed MD simulations on the isolated polymer chain to sample a wide range of PEG chain sizes in a large simulation box ($140 \times 140 \times 140 \text{ Å}^3$) with $\rho = 0.01 \text{ g/cm}^3$. Here, we varied f from 0.1 to 1.0 and carried out NVT MD for 10 ns at each temperature of 100–1200 K. The R_g values for the last 5 ns of each case are shown in Figure S3 (SI). Note that we use the term R_g to be $\langle R_g^2 \rangle^{1/2}$. Snapshots of the conformations for $f = 0.3$ and 1.0 are shown in Figures 3 and 4, respectively, and snapshots for the other cases are shown in Figures S4–S6 (SI). These results show that there is a clear transition temperature (T_t) at $\sim 300 \text{ K}$ for $f = 0.3$ and $\sim 1000 \text{ K}$ for $f = 1.0$. We denote this transition temperature as the coil–globule or θ temperature. Defining the scaled fluctuations in the radius of gyration as

$$\delta R_{g-sc} = \frac{(\text{standard deviation of } R_g)}{(\text{average } R_g)}$$

we find the following:

- Below T_t , intrachain attraction dominates, leading to polymer chain conformations that adopt lamella-like chain structures that change little with time ($\delta R_{g-sc} < \sim 0.04$). This is the poor solvent condition, in which the polymer chains favor globular

a) Relative Standard Deviation R_g (Å)

b) Average R_g (Å)

c) Maximum R_g (Å)

Figure 6. Phase diagram for conformations of an isolated PEG chain showing the dramatic change in properties above the θ temperature characterized by rapidly fluctuating random coil conformations and below the θ temperature characterized by frozen lamella conformations. (a) Relative standard deviation of R_g , (b) average value of R_g , and (c) maximum value of R_g . The relative standard deviation of R_g , δR_{g-sc} , is the normal standard deviation of R_g divided by the average R_g . Color bars indicate the magnitudes of R_g . The black squares with a least-squares fit indicate the θ temperature (T_t) separating the two regimes, which are quite distinctive.

conformations to minimize polymer–solvent contacts and maximize contacts between polymer segments.

Figure 7. (a) Red line: the overall R_g distribution obtained over the temperature range from 100 to 1200 K for scaling factors (f) (from $f = 0.1$ to 1.0) using a large simulation box ($140 \times 140 \times 140 \text{ \AA}^3$) with $\rho = 0.01 \text{ g/cm}^3$. Blue and green lines: R_g distributions obtained by using $f = 0.3$ (blue) or 1.0 (green). (b) Comparison of the R_g distribution obtained by using $f = 0.3$ and varying the temperature in a $63.52 \times 63.52 \times 63.52 \text{ \AA}^3$ simulation box, leading a density of $\rho = 0.13 \text{ g/cm}^3$ (if we insert 75 wt % water, then the total density should be 0.52 g/cm^3 , which is half of the density expected from experiment $\rho^{\text{exp}} = 1.04 \text{ g/cm}^3$) (magenta line) with the R_g distribution obtained by using $f = 0.3$ and varying the temperature in a large simulation box with $\rho = 0.01 \text{ g/cm}^3$ (blue line).

Figure 8. Starting with the PEG chain having $R_g = 55.67 \text{ \AA}$, the DoS for the PEG from the Fourier transform of the velocity autocorrelation function by the 2PT analysis at 300 K for the 25 wt % PEG aqueous solution. The red line is the DoS of the PEG, $S_{\text{PEG}}(v)$. Blue and green lines are gas-like $S_{\text{PEG}}^s(v)$ and solid-like $S_{\text{PEG}}^l(v)$ components of $S_{\text{PEG}}(v)$, respectively. Most of the diffusional contribution is from the water dynamics. The SI shows the total DoS and that of the water for several cases.

• Above T_v , thermal (entropic) effects dominate, leading to fluctuating random coil structures. This is the good solvent condition, in which the polymer chains favor a random coil structure that maximizes configurational entropy, leading to extended conformations.

To determine the T_t for each scaling factor, we consider how $\delta R_{g-\text{sc}}$ changes with temperature. These are shown in Figure 5 for the case of $f = 0.3$ and in Figure S7 (SI) for the other cases. Here, we see that at the lowest temperatures, the lamella or globule region leads to very small fluctuations, $\delta R_{g-\text{sc}} < \sim 0.04$, while the highest temperatures lead to $\delta R_{g-\text{sc}} = 0.19\text{--}0.26$, close to the infinite chain limit for a Gaussian coil of $\delta R_{g-\text{sc}} = 0.26$. (for details, see Table S3 and S4, SI).

To define the θ temperature (T_t) forming the boundary between these two regimes, we used the hyperbolic tangent function, $\delta R_{g-\text{sc}}(T) = a[\tanh\{(T - T_t)/b\} + 1]$. For $f = 0.3$, Figure 5 shows that this leads to $T_t = 252 \text{ K}$.

Figure 6 shows the phase diagram for the PEG conformations as a function of f and temperature, where T_t is indicated by the black squares. The lamella structure regime shows characteristics such as small $\delta R_{g-\text{sc}}$, small maximum R_g , and small average R_g due to very limited movements dominated by the strong intrachain interactions. In contrast, the random coil (R-coil) structure regime exhibits opposite characteristics such as large $\delta R_{g-\text{sc}}$ approaching the limit of 0.26 for infinite Gaussian chains, large maximum R_g , and moderate average R_g .

4.3. Choosing the Initial Ensemble of Polymer Chain Conformations. For a good solvent such as water, we expect that moderately dilute solutions of PEG at 300 K should lead to R-coil structures; therefore, we considered only these cases in choosing the initial ensemble of polymer chain conformations.

Figure 7a (red solid line) shows the overall R_g distribution obtained in a large simulation box ($140 \times 140 \times 140 \text{ \AA}^3$) with $\rho = 0.01 \text{ g/cm}^3$ by averaging over all temperatures (from 100 to 1200 K) and scaling factors ($f = 0.1\text{--}1.0$) but including only cases in the R-coil regime (we note that temperature-dependent R_g distributions are shown in Figure S8, SI). This leads to a broad range of PEG structures (red solid line) with $\Delta R_g \approx 120 \text{ \AA}$, and the most probable value of R_g is $64.44 \text{ \AA}$.

Also shown in Figure 7a are the R_g distributions obtained in a large simulation box ($140 \times 140 \times 140 \text{ \AA}^3$) with $\rho = 0.01 \text{ g/cm}^3$ by using $f = 0.3$ (blue solid line) and 1.0 (green solid line) for all temperatures from T_t to 1200 K. The single case of $f = 0.3$ (blue) leads to an R_g distribution similar to the overall R_g distribution (red). Indeed, the peak of the distribution for the case of $f = 0.3$ is at $R_g = 65.58 \text{ \AA}$, very close to the peak of the overall distribution, $R_g = 64.44 \text{ \AA}$. Therefore, we conclude that MD simulations using only $f = 0.3$ but varying the temperature up to $\sim 1200 \text{ K}$ lead to a distribution of conformations appropriate for spanning the conformations of a PEG chain in solution. This can be compared to studies⁴⁷ by Jang and Goddard on linear and cyclic polyethylene, where $f = 0.28\text{--}0.35$ led to θ temperatures in good agreement with experimental measurements. [In addition, $f = 0.1$ can be another option because it also leads to similar range of R_g as in the case of $f = 0.3$ (see Figure S9, SI).]

As discussed above, sampling of PEG structures using SES dynamics with $f = 0.3$ within the large simulation box with $\rho =$

Figure 9. Thermodynamics calculated by the 2PT method over the last 1 ns of NVT dynamics. (a) Helmholtz free energies, (b) entropies, (c) internal energies, and (d) total MD energies. Red circles show the results from the solvent-included CED (SI-CED) calculations. The blue triangles show the results from using the SES-CED model (with $f = 0.3$) starting the simulation from the half of the experimental value ($0.5 \rho^{\text{exp}}$) as the initial density, followed by density and temperature annealing to near the experimental density (ρ^{exp}). The magenta squares show the SES results starting the simulation from the experimental density (ρ^{exp}) as the initial density. The error bars are the standard deviation of each energetic property over 100 points. The standard deviation of R_g (x-axis error bar) is smaller than the marker size for all samples. The MD energy here is the total energy (sum of the kinetic energy and potential energy) from the last 1 ns of NVT dynamics, -5520.58 ± 135.39 kcal/mol (SI-CED), -5722.74 ± 137.72 kcal/mol (SES-CED), and -5734.67 ± 137.23 kcal/mol (SES), without zero point energy and thermal corrections. These results show that the SES values are equivalent to the SES-CED values, indicating no extra advantage to the CED process for this system. In contrast, the simple SI-CED leads to significantly higher energies. Note however that the entropy is similar to that from SES, indicating that the main error in SI-CED is strain in the conformation that has not been fully relaxed by the CED process, leading to ~ 200 kcal/mol of extra strain energy (0.45 kcal/mol per monomer).

0.01 g/cm³ leads to R-coil PEG structures exhibiting a wide size range, $\Delta R_g \approx 110$ Å, with the most probable value of $R_g = 65.58$ Å. Therefore, we performed NVT MD using $f = 0.3$ on the single isolated polymer chain in a $63.52 \times 63.52 \times 63.52$ Å³ simulation box, leading to a density of $\rho = 0.13$ g/cm³ (if we insert 75 wt % water, then the total density should be 0.52 g/cm³, which is half of the density expected from experiment $\rho^{\text{exp}} = 1.04$ g/cm³). This procedure is to obtain a wide range of the initial ensemble of polymer chain conformations at half of the target density. Figure 7b shows that in the case of $f = 0.3$, compared with the case of a very low density system ($140 \times 140 \times 140$ Å³) (blue solid line), the range of R_g of the chains from the $63.52 \times 63.52 \times 63.52$ Å³ simulation box decreased due to interactions with the image chains at higher density.

4.4. SES-CED Method. The steps of the SES-CED method are as follows:

- (1) Grow the polymer chain using standard MC.
- (2) Carry out MD calculations on the isolated chain using an appropriate range of the nonbonding scaling factor f and a box of sufficiently low density to avoid chain–chain interactions with its periodic image. For our PEG-20 kDa system aimed at 25 wt % aqueous solution, we started with a $63.52 \times 63.52 \times 63.52$ Å³

simulation box, which would lead to a density of $\rho = 0.52$ g/cm³ after adding water (assuming that 75 wt % water is included), which is 50% of the density measured from experiment,⁴⁸ $\rho^{\text{exp}} = 1.04$ g/cm³, for 25 wt % PEG. The actual density before adding water is 0.13 g/cm³. This procedure was described above in section 4.3. From the distribution in Figure 7b for $f = 0.3$ and $\rho = 0.13$ g/cm³, we selected 21 different PEG chains spaced evenly over the range of $R_g = 33.80$ – 95.68 Å for use in the SES-CED procedure.

(3) Using the equilibrated polymer chains from step 2, introduce water molecules into each simulation cell but change f back to 1 because we now account for explicit solvation. To find favorable locations for the addition of water molecules without perturbing the polymer conformation, we carried out grand canonical Monte Carlo (GCMC) simulations while fixing the final number of water loading to the target value (3339 for the 25 wt % case).

(4) Then, starting with the chain–water distributions from step 3, compress the simulation cell gradually (uniformly over ~ 3 ns) from half of the expected density ($0.5\rho^{\text{exp}} = 0.52$ g/cm³) up to 1.1 times the expected density ($1.1\rho^{\text{exp}} = 1.14$ g/cm³) while carrying out temperature annealing between 300 and 706 K

Figure 10. Gaussian distribution for each energetic property, (a) Helmholtz free energies (A), (b) entropies (S), (c) internal energies (E), and (d) total MD energies for the cases of SI-CED (green line), SES-CED (blue line), and SES (magenta line). These figures show that the SES/SES-CED method leads to well-equilibrated PEG conformations in a 25 wt % PEG aqueous solution, compared to the SI-CED method. Considering all 21 of the SES-CED configurations at the ensemble, we calculated $A = 21\,572.37 \pm 84.51$ kcal/mol, $S = 52\,595.63 \pm 188.42$ cal/mol/K, and $U = 37\,485.77 \pm 55.74$ kcal/mol. Considering all 13 of the SES configurations at the ensemble, we calculated $A = 21\,561.05 \pm 84.37$ kcal/mol, $S = 52\,593.32 \pm 185.63$ cal/mol/K, and $U = 37\,475.39 \pm 55.68$ kcal/mol.

(approximately twice the Flory temperature; 353 K^{39}). This was done for 5 cycles.

(5) Starting with the chain–water distributions with the density of $1.1\rho^{\text{exp}}$ from step 4, carry out NPT MD at 1 atm and 300 K for 4–15 ns to determine the equilibrium density (ρ^{eq}). The full SES-CED procedure leads to final densities of the 21 samples (with explicit waters) ranging from $\rho^{\text{eq}} = 1.0555$ to 1.0571 g/cm^3 , which agrees with the experimental density⁴⁸ (ρ^{exp}) of 1.0417 g/cm^3 within 1% for 25% PEG. This provides confidence that the water–PEG interaction is well described by our new FF and that the SES-CED procedure leads to well-equilibrated structures.

(6) At the predicted equilibrium density from step 5, perform NVT MD for 10–20 ns. The 1 ns trajectories are used for the thermodynamic and structural analyses by using 2PT method.²⁹ We performed thermodynamic analyses on the sampled polymer conformers immersed in solvent to calculate the thermodynamic preference of the polymer configuration in solvent.

4.5. Thermodynamic Analysis Using the 2PT Density of States (DoS) Method. To determine the thermodynamic properties of the PEG–water systems, we used 2PT theory, which partitions the vibrational density of states (DoS) from the Fourier transform of the velocity autocorrelation function into diffusional contributions (leading to $\text{DoS}(\nu) \neq 0$ at $\nu = 0$) and a solid-like vibrational contribution (leading to $\text{DoS}(\nu) = 0$ at $\nu = 0$).

The trajectories from the last 1 ns NVT dynamics were used for the thermodynamic and structural analyses. We used the 2PT method²⁹ to predict the thermodynamic properties of the polymer conformations immersed in solvent. This is used to calculate the thermodynamically preferred polymer configuration in solvent. Because 2PT requires analysis only over a period of 20 ps to get an accurate free energy and entropy, we restarted the trajectories every 10 ps over the last 1 ns of NVT and did the 20 ps 2PT analysis 100 times to provide both thermodynamic quantities and their fluctuations.

Figure 8 shows the DoS of PEG for a representative case, $R_g = 55.67\text{ Å}$. [Figures S10–S12 (SI) show the full DoS and the partition into PEG and water parts for three representative cases, $R_g = 34.86$, 55.67 , and 78.77 Å .] This DoS leads to a partition function from which we calculate the Helmholtz free energy (A), entropy (S), and internal energy (E) for the 21 independent samples.

This leads to the results shown in Figure 9a–c. Because the dynamics of PEG in the solvated system leads to only marginal changes in its chain conformation during the 1 ns MD simulation, we regard each calculated thermodynamic quantity as the local value for the given PEG conformation, while we consider that the Boltzmann average over all 21 conformations provides the thermodynamic quantities for the equilibrium polymer–water system. This leads to an average over the entire

Figure 11. The final chain conformations of a PEG in aqueous solutions with 25 wt % PEG for the 21 cases of SES-CED. The final values of R_g are shown in the figure. White, red, and green atoms are hydrogen, oxygen, and carbon atoms, respectively. The periodic box is shown as a blue solid line. For all cases, the final densities range from 1.0555 to 1.0571 g/cm³, comparable with the experimental density⁴⁸ of 1.0417 g/cm³. The initial values of R_g were (a) 41.95, (b) 33.80, (c) 38.81, (d) 36.03, (e) 48.23, (f) 45.03, (g) 51.28, (h) 63.77, (i) 54.40, (j) 60.66, (k) 57.56, (l) 73.14, (m) 79.34, (n) 66.82, (o) 76.20, (p) 70.00, (q) 82.48, (r) 85.01, (s) 90.59, (t) 88.90, and (u) 95.68 Å.

phase space from separate integrations over the subvolume of each nearly fixed PEG conformation.

Figure 9a–c shows that all 21 samples obtained from SES-CED lead to very similar free energies (with standard deviations of $\sigma_A = 22.92$ kcal/mol/cell, $\sigma_E = 17.05$ kcal/mol/cell, and $\sigma_S = 32.94$ cal/mol/K/cell) with respect to their statistical fluctuations due to the solvent molecules.

Figure 10 shows the result of Boltzmann weighting the distributions of energetic properties for each of the 21 cases in Figure 9 to obtain the thermodynamic average over the whole ensemble. This shows that the Helmholtz free energy from SES-CED is substantially more stable than that from SI-CED.

The similarity of the energetics ($\Delta A \approx 104$ kcal/mol/cell compared with a total average $A = 21\,572.37$ kcal/mol/cell) for the large range of polymer conformations ($\Delta R_g \approx 44$ Å) suggests that the conformational dynamics of the PEG in this semidilute regime (25 wt %) is similar to the polymer dynamics in a θ solvent.⁴⁹ That is, the characteristics of the PEG conformation are determined by the short-range properties of the polymer backbone.

4.6. Analysis of Results. Figure 9d shows that all 21 PEG structures lead to nearly equal total MD energies with a range from -5753.39 to -5679.58 kcal/mol, where the fluctuations within each system are ± 137.72 kcal/mol. This gives us

Final PEG conformations after SES

Figure 12. The final chain conformations of a PEG in aqueous solutions with 25 wt % PEG for the 13 cases of SES. The final values of R_g are shown in the figure. White, red, and green atoms are hydrogen, oxygen, and carbon atoms, respectively. The periodic box is shown as a blue solid line. For all cases, the final densities range from 1.0557 to 1.0573 g/cm³, which compare with the experimental density of 1.0417 g/cm³. The initial values of R_g were (a) 30.35, (b) 35.77, (c) 42.87, (d) 39.38, (e) 46.38, (f) 49.98, (g) 53.58, (h) 57.06, (i) 60.68, (j) 64.24, (k) 67.74, (l) 71.11, and (m) 72.30 Å.

confidence that the procedure leads to well-equilibrated chain–solvent systems so that this range of sizes is expected to occur over the time scales of the experiment.

In contrast, the SI-CED method (red dot in Figure 9d) leads to a MD energy higher by ~ 200 kcal/mol and to an R_g that is small compared to that of the range of structures from SES-CED.

We emphasize that the introduction of SES dynamics without explicit water molecules in steps 1 and 2 is crucial to achieve the wide range of polymer chain conformations within a short (~ 40 ns) MD simulation time scale. This is because the conformational dynamics of the PEG chain in explicit water solvent changes only very slowly, with a standard deviation in R_g of < 0.60 Å within 36 ns of MD.

The final 21 PEG chain configurations of the equilibrated PEG–water system with 25 wt % PEG chains sample a large

range of PEG conformations with R_g varying from 34.86 to 78.77 Å. Figure 11 shows the final chain conformations for the 21 cases of SES-CED.

On the basis of the Boltzmann distribution of free energies predicted for the 21 cases, we re-evaluated the R_g , leading to $R_g = 53.98$ Å, which is the final thermodynamically averaged size from the SES-CED method for 25 wt % PEG in water at 300 K. These results are quite different than the result from SI-CED, which leads to $R_g = 36.57$ Å, but are comparable to $R_g = 65$ Å from viscosity experiments⁵⁰ in the dilute limit.

To interpret this R_g using the freely jointed polymer (Gaussian random coil) model would require a segment length of $l = 13.16$ with $N = 147$ segments to preserve the total length, leading to $(R_g^2)^{1/2} = l(N/6)^{1/2} = 65.14$ Å.

This suggests that the ether oxygen atoms serve as the joints (with the rigid O—CH₂—CH₂—O units). This leads to $l = 4.251$ Å, which, with $N = 455$, leads to $\langle R_g^2 \rangle^{1/2} = 37.02$ Å. This suggests that we use the freely rotating chain in which the C—O—C angle is fixed at the minimum energy value of 113.7° . This leads to $\langle R_g^2 \rangle^{1/2} = 37.02[(1 + \cos \theta)/(1 - \cos \theta)]^{1/2} = 56.68$ Å, where $\theta = 180 - 113.7 = 66.3^\circ$, which is in excellent agreement with the simulations.

5. SES METHOD

Our analysis on the results from the SES-CED calculations suggests that it might not be necessary to start with the half density cell. This leads to the SES method in which we start with the simulation box of $50.42 \times 50.42 \times 50.42$ Å³, leading to an initial density of $\rho = 1.04$ g/cm³ = ρ^{exp} (after including water molecules) rather than $0.5\rho^{\text{exp}}$ as in the previous SES-CED method. From the distribution obtained in the box with ρ^{exp} , we selected 13 different PEG chains spaced evenly for use in this new SES procedure. By modifying step 2 of SES-CED in this way, we excluded the compression annealing from the SES-CED (step 4) and used only the *NPT* dynamics (step 5) to adjust the final simulation box size to have the optimal density. Because compression annealing of the simulation box is not involved here, we are able to transfer the PEG chain sampled from SES dynamics into the explicit solvation box. The equilibrated final structures are shown in Figure 12. The final densities of the 13 samples (with explicit waters) range from 1.0557 to 1.0573 g/cm³, which are within 1% of the experimental density⁴⁸ of 1.0417 g/cm³.

The trajectories from the last 1 ns *NVT* dynamics are used for the thermodynamic and structural analyses. Using the 2PT method, we perform thermodynamic analyses on the sampled polymer conformers immersed in solvent to calculate the thermodynamic preference of the polymer configuration in solvent. Figure 9 confirms that these 13 new structures have a range of energies and sizes similar to those of the 21 structures from the SES-CED procedure.

Calculating the Helmholtz free energies based on a standard Boltzmann distribution for the 13 cases leads to the Helmholtz free energy distribution in Figure 10a, which is slightly more stable than that for SES-CED. Using this Boltzmann-weighted distribution, we calculate $R_g = 55.48$ Å for 25 wt % PEG, which is similar to the value of R_g (53.98 Å) calculated from SES-CED. Thus, the SES method lead to a broad range of well-equilibrated PEG conformations in water, in contrast to SI-CED, which leads to only one size of PEG conformation and a much higher energy (see Figures 9 and 10).

The optimum R_g from SES compares well with experimental ensemble-averaged R_g of PEG conformations in the dilute concentration regime; $R_g = 57.11$ – 69.17 Å from SLS measurements,^{12,15,51} and $R_g = 65$ Å from viscosity measurements.⁵⁰

This SES calculation indicates that properly equilibrated chain–water structures can be constructed directly at the target density without the density annealing steps. We expect that this SES procedure can be useful in future simulations.

6. CONCLUSIONS

We present and validate here the new SES method for sampling a wide range of polymer chain sizes for a high-molecular-weight (20 kDa) PEG chain dissolved in water for polymer concentrations of 25 wt %.

Our detailed thermodynamic analysis shows that the polymer configurations are determined mostly by short-range properties of the polymer backbone allowing substantial size variations in the PEG chains ($\Delta R_g = 43.91$ Å from SES-CED and $\Delta R_g = 38.98$ Å from SES) for solvation in the semidilute regime of 25 wt %. This is because the stiffness of the PEG backbone opens sufficient space for water molecules to form a strong first solvation shell (via a short-range interaction), stabilizing the polymer independently.

We expect that the SES method will be useful to investigate comprehensively the solvent–polymer interaction with MD simulations, as illustrated with this study of PEG–water systems. Such MD studies should provide atomistic insights into PEG–water interactions in the semidilute regime (where experimental characterization on the structure is almost impossible), which is directly related to the nonlinear increase of osmotic behavior of PEG.

■ ASSOCIATED CONTENT

Supporting Information

The comparison between QM calculation results and the results calculated by our optimum FF is shown in Figures S1 and S2, the optimum FF parameters are listed in Table S1, the partial charge distributions of a PEG-20 kDa are shown in Table S2, the R_g variation with time for a scaling factor ranging from 0.1 to 1.0 and the temperature ranging from 100 to 1200 K is illustrated in Figure S3, the snapshots of the conformations with time and temperature for $f = 0.1, 0.5$, and 0.7 are shown in Figures S4–S6, the determination of the θ temperature for each scaling factor is explained in Figure S7, the scaled standard deviations of R_g at the highest temperature for each scaling factor are listed in Table S3, the scaled standard deviation of R_g for the ideal Gaussian chains with a different number of monomers or polymer chains is shown in Table S4, the R_g distribution of random coils sampled with $f = 0.3$ as a function of temperature is illustrated in Figure S8, the R_g distributions of random coils sampled with $f = 0.1$ or 0.3 are shown in Figure S9, and the density of states for three representative cases $R_g = 34.86, 55.67$, and 78.77 Å are shown in Figures S10–S12. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

The authors declare no competing financial interest.

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■ REFERENCES

- (1) Bekiranov, S.; Bruinsma, R.; Pincus, P. *Phys. Rev. E* **1997**, *55*, 577–585.
- (2) Harris, J. M. *Poly(Ethylene Glycol) Chemistry: Biotechnical and Biomedical Applications*; Plenum: New York, 1992; p 385.

- (3) Veronese, F. M.; Pasut, G. *Drug Discovery Today* **2005**, *10*, 1451–1458.
- (4) Huang, Y. Y.; Chung, T. W.; Tzeng, T. W. *Int. J. Pharm.* **1999**, *182*, 93–100.
- (5) Molineux, G. *Cancer Treat. Rev.* **2002**, *28*, 13–16.
- (6) Knop, K.; Hoogenboom, R.; Fischer, D.; Schubert, U. S. *Angew. Chem., Int. Ed.* **2010**, *49*, 6288–6308.
- (7) Liu, Z.; Robinson, J. T.; Sun, X. M.; Dai, H. J. *J. Am. Chem. Soc.* **2008**, *130*, 10876–10877.
- (8) Le Coz, C. J.; Heid, E. *Contact Dermatitis* **2001**, *44*, 308–319.
- (9) Fruijtier-Polloth, C. *Toxicology* **2005**, *214*, 1–38.
- (10) Ye, Y.; Rick, J.; Hwang, B. *Polymers* **2012**, *4*, 913–963.
- (11) Devanand, K.; Selser, J. C. *Nature* **1990**, *343*, 739–741.
- (12) Polverari, M.; van de Ven, T. G. M. *J. Phys. Chem.* **1996**, *100*, 13687–13695.
- (13) Ho, D. L.; Hammouda, B.; Kline, S. R. *J. Polym. Sci., Part B: Polym. Phys.* **2003**, *41*, 135–138.
- (14) Polik, W. F.; Burchard, W. *Macromolecules* **1983**, *16*, 978–982.
- (15) Kawaguchi, S.; Imai, G.; Suzuki, J.; Miyahara, A.; Kitano, T. *Polymer* **1997**, *38*, 2885–2891.
- (16) Maeda, Y.; Tsukida, N.; Kitano, H.; Terada, T.; Yamanaka, J. *J. Phys. Chem.* **1993**, *97*, 13903–13906.
- (17) Maxfield, J.; Shepherd, I. W. *Polymer* **1975**, *16*, 505–509.
- (18) Derkaoui, N.; Said, S.; Grohens, Y.; Olier, R.; Privat, M. *J. Colloid Interface Sci.* **2007**, *305*, 330–338.
- (19) Liu, K.; Parsons, J. L. *Macromolecules* **1969**, *2*, 529–533.
- (20) Faraone, A.; Magazu, S.; Maisano, G.; Migliardo, P.; Tettamanti, E.; Villari, V. *J. Chem. Phys.* **1999**, *110*, 1801–1806.
- (21) Hammouda, B.; Ho, D.; Kline, S. *Macromolecules* **2002**, *35*, 8578–8585.
- (22) Hammouda, B.; Ho, D. L.; Kline, S. *Macromolecules* **2004**, *37*, 6932–6937.
- (23) Kozer, N.; Kuttner, Y. Y.; Haran, G.; Schreiber, G. *Biophys. J.* **2007**, *92*, 2139–2149.
- (24) Tasaki, K. *J. Am. Chem. Soc.* **1996**, *118*, 8459–8469.
- (25) Smith, G. D.; Bedrov, D.; Borodin, O. *Phys. Rev. Lett.* **2000**, *85*, 5583–5586.
- (26) Smith, G. D.; Bedrov, D.; Borodin, O. *J. Am. Chem. Soc.* **2000**, *122*, 9548–9549.
- (27) Borodin, O.; Bedrov, D.; Smith, G. D. *Macromolecules* **2001**, *34*, 5687–5693.
- (28) Boils, D.; Hair, M. L. *J. Colloid Interface Sci.* **1993**, *157*, 19–23.
- (29) Lin, S. T.; Blanco, M.; Goddard, W. A., III. *J. Chem. Phys.* **2003**, *119*, 11792–11805.
- (30) Mayo, S. L.; Olafson, B. D.; Goddard, W. A., III. *J. Phys. Chem.* **1990**, *94*, 8897–8909.
- (31) Zhao, Y.; Truhlar, D. G. *Theor. Chem. Acc.* **2008**, *120*, 215–241.
- (32) Dunning, T. H. *J. Chem. Phys.* **1989**, *90*, 1007–1023.
- (33) *Jaguar*, version 7.6; Schrödinger, LLC: New York, 2009.
- (34) Levitt, M.; Hirshberg, M.; Sharon, R.; Laidig, K. E.; Daggett, V. *J. Phys. Chem. B* **1997**, *101*, 5051–5061.
- (35) Plimpton, S. J. *Comput. Phys.* **1995**, *117*, 1–19.
- (36) Sadanobu, J.; Goddard, W. A., III. *Fluid Phase Equilib.* **1998**, *144*, 415–425.
- (37) Sadanobu, J.; Goddard, W. A., III. *J. Chem. Phys.* **1997**, *106*, 6722–6729.
- (38) Belmares, M.; Blanco, M.; Goddard, W. A., III; Ross, R. B.; Caldwell, G.; Chou, S. H.; Pham, J.; Olofson, P. M.; Thomas, C. J. *Comput. Chem.* **2004**, *25*, 1814–1826.
- (39) Venohr, H.; Fraaije, V.; Strunk, H.; Borchard, W. *Eur. Polym. J.* **1998**, *34*, 723–732.
- (40) Jaramillo-Botero, A.; Blanco, M.; Li, Y. Y.; McGuinness, G.; Goddard, W. A., III. *J. Comput. Theor. Nanosci.* **2010**, *7*, 1238–1256.
- (41) Henz, B. J.; Chung, P. W.; Andzelm, J. W.; Chantawansri, T. L.; Lenhart, J. L.; Beyer, F. L. *Langmuir* **2011**, *27*, 7836–7842.
- (42) Rai, N.; Wagner, A. J.; Ross, R. B.; Siepmann, J. I. *J. Chem. Theory Comput.* **2008**, *4*, 136–144.
- (43) Jang, S. S.; Molinero, V.; Cagin, T.; Goddard, W. A., III. *J. Phys. Chem. B* **2004**, *108*, 3149–3157.
- (44) Jang, S. S.; Goddard, W. A., III. *J. Phys. Chem. C* **2007**, *111*, 2759–2769.
- (45) Jeon, J.; Kim, H.; Goddard, W. A., III; Pascal, T. A.; Lee, G. I.; Kang, J. K. *J. Phys. Chem. Lett.* **2012**, *3*, 556–559.
- (46) Jang, S. S.; Molinero, V.; Cagin, T.; Goddard, W. A., III. *Solid State Ionics* **2004**, *175*, 805–808.
- (47) Jang, S. S.; Cagin, T.; Goddard, W. A., III. *J. Chem. Phys.* **2003**, *119*, 1843–1854.
- (48) Cruz, R. D.; Martins, R. J.; Cardoso, M. J. E. D.; Barcia, O. E. *J. Solution Chem.* **2009**, *38*, 957–981.
- (49) Adam, M.; Delsanti, M. *Macromolecules* **1985**, *18*, 1760–1770.
- (50) Kato, T.; Nakamura, K.; Kawaguchi, M.; Takahashi, A. *Polym. J.* **1981**, *13*, 1037–1043.
- (51) Devanand, K.; Selser, J. C. *Macromolecules* **1991**, *24*, 5943–5947.