

Structure, Stability, and Hydration of a Polypeptide in AOT Reverse Micelles

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It has been known for a time, both from experiments¹ and theory,² that confining a peptide chain within a geometrically restricted environment increases the relative stability of the folded state against unfolded states. This finding is of biological relevance as (in vitro) intrinsically disordered proteins have shown to be structured or folded in their native cellular environment where excluded volume effects are important. A convenient means to confine a protein within a certain region of space is through encapsulation in reverse micelles³ (RM). In this communication, we provide theoretical evidence that the folded structure of a simple peptide, alanine zwitterionic octapeptide, or A₈, unstable in solution, becomes stable in an RM of appropriate size. Our molecular dynamics simulations were carried out for realistic models of sodium 2-ethylhexylsulfosuccinate (AOT) RM in isooctane, simulated for an extended period of time. For the RM of the smaller size, we find that a helical structure is stable for the whole length of the simulation. On the contrary, the peptide very quickly takes an extended structure in larger micelles.

In a previous study, we developed a realistic molecular model for AOT reverse micelles in isooctane and reported on the structural properties of the micelles and their water core⁴ as a function of the water-to-surfactant molar ratio ($W_o = [H_2O]/[AOT]$). Here, we focus on the quaternary system A₈/AOT/water/isooctane. Studies using both experiments⁵ and simulations⁶ have shown that in water at room temperature an octaalanine is unstable in its canonical α -helix conformation. According to MD investigations,⁶ the peptide prepared as an α -helix evolves to extended structures within a few hundred or thousand picoseconds of simulation, depending on the force field. This behavior makes octaalanine an ideally simple but physically relevant system to study confinement effects for protein secondary structure.

In this work,⁷ all simulations were performed in an isobaric–isothermal (NPT) ensemble at $T = 300$ K and $P = 0.1$ MPa, as described in ref 8. In a first series of MD runs (see ref 4), two reverse micelles with 64 and 82 molecules of AOT and 320 and 574 molecules of water (named RM64 and RM82 and corresponding to $W_o = 5$ and 7, respectively) were simulated. For the current study, an additional series of runs, for a total of 26 ns (for both systems we ran two simulations of 3 and 10 ns),⁷ was performed by inserting a canonical α -helical octaalanine in the center of the RM64 and RM82 water core. In doing so, water molecules in contact with the peptide were removed, thus leaving 306 and 557 waters in the hydrophilic cores of RM64 and RM82, respectively. The new systems, named RM64+A₈ and RM82+A₈, had water-to-surfactant molar ratios of $W_o = 4.78$ and 6.79, respectively. The small reduction of the number of water molecules (less than 5 and 3% between identical micelles) allows a direct comparison with

the peptide-free RM. For all AOT RMs studied here, the solvent concentration was chosen to be compatible with the L₂ phase of AOT/water/isooctane.⁹

We first notice¹⁰ that RMs with and without the peptide are of ellipsoidal shape. As discussed in ref 4, there is experimental evidence that AOT RMs are nonspherical (see, for instance, ref 11). The ratio between the major and the minor semi-axis, $\langle a/c \rangle$, for each micelle is within 1.66 (RM64+A₈) and 1.77 (RM82+A₈) units. Inserting A₈ modifies this semi-axis ratio for both micelles and their water cores. For RM64, $\langle a/c \rangle$ increases, whereas it decreases for RM82. The radius of gyration R_g , an estimate of the RM dimensions, is less affected by the peptide as it changes only a few percent. The largest variation is seen for RM82 \rightarrow RM82+A₈, where R_g of the water core goes from 14.6 to 16.6 Å. This is probably related to the structural modifications of the confined peptide (see below).

The radial density profiles computed for all components of our systems also show that for the peptide-containing RMs the AOT aliphatic chains protect the water core from the oil and that a large interfacial zone exists occupied by the AOT headgroups and Na⁺ ions. We also observe that octaalanine is well-confined within RM64+A₈ and RM82+A₈ micelles, since no surface contacts¹² between the peptide and the solvent are detected during the simulation time. Calculations of the pair density correlation functions, $\rho(r)$, show that insertion of the peptide does not modify the hydration of the AOT headgroups. Indeed, we find that the number of water molecules bound to the AOT sulfonate groups remains very close to that found in peptide-free micelles with changes of only 0.4 units (see Table 1 in the Supporting Information). We also notice that A₈ produces small changes on the residence times, τ_w , of the waters bound to the AOT headgroups of RM82, whereas its effect is larger for RM64. If τ_w and τ_w^{bulk} are the water residence times of the system under consideration and of the bulk, the retardation ratio is defined as $\tau_w/\tau_w^{\text{bulk}}$. We find that the retardation of 2.7 in RM82 changes very little, to 3.3, in RM82+A₈. On the contrary, the retardation of 3.1 for RM64 is almost doubled, to 5.5, by the peptide. This means that water is slowed considerably in RM64+A₈. This likely prevents the peptide to break its intramolecular H-bonds and might have crucial implications for the kinetic stability of the peptide folded structure.

In Figure 1, we show typical conformations of A₈ in the RM64 and RM82 micelles. We first notice that the peptide α -helical secondary structure is conserved for RM64+A₈, whereas for RM82+A₈ the intramolecular hydrogen bonds (IHB) are disrupted. To quantify this behavior, we counted the IHB between the carbonyl and the amide groups separated by three residues, using geometrical and energy criterions¹³ (see Supporting Information). In bulk water, no IHB persisted after only a few tens of picoseconds, confirming previous results in ref 6. Conversely, in the RM64+A₈ simulation

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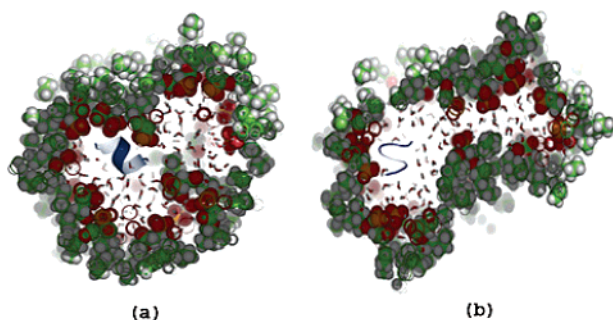


Figure 1. Views of the reverse micelles RM64+A₈ (a) and RM82+A₈ (b) at the end of the NPT simulations. These are views of the simulation box clipped by an orthogonal plane to the *X*-axis. The peptide is drawn as a cartoon in blue color. The atoms of AOT are represented by spheres, whereas the water atoms are represented by sticks. For visual clarity, the Na⁺ ions and the isooctane molecules are not shown. The pictures were drawn with PyMOL.¹⁴

three out of four IHBs were maintained during the full length of the simulation, with some fast, less than 300-ps long, hydrogen bond exchanges with water. In RM82+A₈, during the 3-ns run two of the bonds were lost after 10 ps, whereas the remaining two, CO(1) → NH(5) and CO(2) → NH(6), lasted for 0.5 and 1.5 ns, respectively. In the 10-ns run, the IHB kinetic was slower and only the CO(2) → NH(6) IHB persisted continuously for the entire trajectory. At the same time, deviations from the initial α -helical structure remained below 1.8 Å for both RM64+A₈ simulations but reached as high as 4.2 Å for the shorter and 3 Å for the longer RM82+A₈ simulation.

We find that a possible explanation for the stability of the A₈ α -helix in RM64 resides in the competition for hydration water between the AOT headgroup and the peptide. In the RM64 system, the quantity of water not bound to the sulfonate group is very limited, preventing the full hydration of A₈. This was confirmed by two extra simulations we carried out on A₈ in the modified environments: (1) a RM64 system where the sulfonate group in AOT and the Na⁺ ions were removed, for a system electrically neutral, designated as RM64+A₈* and (2) A droplet of 306 water molecules, the same number as in RM64, in isooctane, named DROP. The two new systems were run in the same conditions of temperature and pressure as the other systems for 3 and 2.5 ns, respectively. In both cases, we observe the unfolding of the peptide and then its migration to the interface between water and the modified detergent/oil phases. In the case of RM64+A₈*, the waters represent about 9% of the total surface of the solute in direct contact with the oil, and 20 modified detergent molecules are found in the solvent region at the end of the simulation run. The peptide needs 0.5 ns to unfold to an extended structure and 1.8 ns to get in stable contact with solvent and detergent molecules. For the DROP system, the unfolding to an extended structure is faster, 0.2 ns, and its migration to the oil/water interface occurs in 0.25 ns.

This communication has shown that the major force affecting the stability of octalanine in small size RM of AOT is hydration. In our simulations, the detergent and the peptide polar groups have competing affinities with water: Hydration is stronger for the AOT sulfonate groups and the Na⁺ ions than for A₈. The properties of micellar water are known experimentally to differ significantly from those of the bulk¹⁵ and affect the confined peptides/proteins structures.³ In our simulations, we find that in the smallest RM,

the dynamic of water is considerably slowed with respect to the bulk, 5.5 times, whereas not enough water is available to solvate the carbonyl and amide groups of the peptide. To conclude, our results provide a strong evidence of the crucial role played by hydration on the structure of peptides in confined hydrophilic environments. Hence, both hydration and excluded volume effects² need to be taken into account to study the secondary structure stability of proteins in restricted spaces such as RM, nanoporous silica, and the drug nanocarriers as well as the interior of ribosomes, or regions of steric occlusion inside the cell.

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Supporting Information Available: The details of our simulation protocols and an extended analysis of our results are reported in the Supporting Information. This material is available free of charge via the Internet at <http://pubs.acs.org>.

References

- (1) (a) Thompson, K. F.; Gierash, L. M. *J. Am. Chem. Soc.* **1984**, *106*, 3648–3652. (b) Gallay, J.; Vincent, M.; Nicot, C.; Waks, M. *Biochemistry* **1987**, *26*, 5738–5747. (c) Eggers, D. K.; Valentine, J. S. *Protein Sci.* **2001**, *10*, 250–261. (d) Bolis, D.; Politou, A. S.; Kelly, G.; Pastore, A.; Temussi, P. A. *J. Mol. Biol.* **2004**, *336*, 203–212. (e) Eggers, D. K.; Valentine, J. S. *Protein Sci.* **2001**, *10*, 250–261.
- (2) (a) Minton, A. P. *Biophys. J.* **1992**, *63*, 1090–1100. (b) Zhou, H. X.; Dill, K. A. *Biochemistry* **2001**, *40*, 11289–11293.
- (3) (a) Luisi, P. L.; Magid, L. J. *Crit. Rev. Biochim.* **1986**, *20*, 409–74. (b) Nicot, C.; Waks, M. *Biotechnol. Genet. Eng. Rev.* **1996**, *13*, 267–314. (c) Shastry, M. C. R.; Eftink, M. R. *Biochemistry* **1996**, *35*, 4094–4101. (d) Thena de Gomez Puyou, M.; Gomez-Puyou, A. *Crit. Rev. Biochem. Mol. Biol.* **1998**, *33*, 53–89. (e) Fiori, S.; Renner, C.; Cramer, J.; Pegoraro, S.; Moroder, L. *J. Mol. Biol.* **1999**, *291*, 163–175. (f) Lundgren, J. S.; Heitz, M. P.; Bright, F. V. *Anal. Chem.* **2000**, *72*, 3775. (g) Peterson, R. W.; Anbalagan, K.; Tommos, C.; Wand, A. J. *J. Am. Chem. Soc.* **2004**, *126*, 9498–9499.
- (4) Abel, S.; Sterpone, F.; Bandyopadhyay, S.; Marchi, M. *J. Phys. Chem. B* **2004**, *108*, 19458–19466.
- (5) Chakrabarty, A.; Baldwin, R. L. *Adv. Protein Chem.* **1995**, *46*, 141–150.
- (6) (a) Weber, W.; Hunenberger, P. H.; McCammon, J. C. *J. Phys. Chem. B* **2000**, *104*, 3668–3675. (b) Margulis, C. J.; Stern, H. A.; Berne, B. J. *J. Phys. Chem. B* **2002**, *106*, 10748–10752.
- (7) See the Supporting Information for a full description of the simulation parameters and procedures. For each RM containing peptide, we ran two simulations with different initial conditions of 3 and 10 ns, respectively. If not specified, results reported in this communication are from the 10-ns trajectory.
- (8) Marchi, M.; Procacci, P. *J. Phys. Chem. B* **1998**, *102*, 5194–5197.
- (9) *Structure and reactivity in reverse micelles*; Pileni, M. P., Ed.; Elsevier: Amsterdam, 1989.
- (10) The structural results discussed below are summarized in Table 1 of the Supporting Information.
- (11) Yano, J.; Milhofer, H. F.; Wachtel, E.; Garti, N. *Langmuir* **2000**, *16*, 9996–10004.
- (12) We compute the contact surface between species A and B as the sum of the surface of the Voronoi^{12a} facets shared between all the atoms of A with those of B. For a reference to the Voronoi polyhedra the reader should refer to: (a) Voronoi, G. F. *J. Reine Angew. Math.* **1908**, *134*, 198–287. Our calculations were carried out according to ref 12b: (b) Procacci, P.; Scateni, R. *Int. J. Quantum Chem.* **1992**, *42*, 1515–1528.
- (13) An IHB was assumed to be formed if the distance between the donor (NH) and acceptor (CO) group was smaller than 0.25 nm. For the energy criterion, we used the Kabsch and Sander function: An HB between (NH) and (CO) exists if the energy is ≤ -0.5 kcal/mol (for further details see Kabsch, W.; Sander, C. *Biopolymers* **1983**, *22*, 2577–2637).
- (14) DeLano, W. L. *The PyMOL Molecular Graphics System*; DeLano Scientific: San Carlos, CA, 2002; <http://pymol.sourceforge.net/>.
- (15) Levinger, N. *Science* **2002**, *298*, 1722–1723.

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