Effect of a Clathrate-Forming Counterion on Micellar Solubilization

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Abstract - Clathrate hydrates are supramolecular solids in which small, hydrophobic species are trapped inside a network of hydrogen-bonded water cages. While the best known examples are the natural gas hydrates, similar crystalline structures can also be formed from salts of tetra-n-butylammonium hydroxide (TBAOH). When TBAOH is reacted with medium to long chain carboxylic acids it produces surfactant compounds capable of forming either clathrate hydrates or micelle solutions in aqueous solution. However, little is known about how this dual nature affects the functional properties of the surfactants. Here we show a direct comparison of the solubilization of the solvatochromic dye, Nile Red (NR) by sodium 10-undecenoate (NaU) and tetrabutylammonium 10-undecenoate (TBAU). We found from the fluorescence emission spectra that the environment around NR was more polar for TBAU micellar solutions than for NaU. Molecular dynamics simulations indicated that for both systems, the NR had partial exposure to water at the surface of the micelle. The primary difference was a significant structuring of the counterions and water molecules as part of the micelle/NR complex for TBAU. Our results demonstrate an active role of role of the clathrate-forming counterion in the formation of the micelle and the solubilization of the substrate. We anticipate that the more polar environment near the surface of the micelle induced by the TBA/H$_2$O structuring will enhance the solubilization power of the micelle for more polar substrates. Furthermore, we expect the more structured shell around the micelle core to enhance the isolation of the substrate from the environment, a valuable feature for applications such as targeted drug delivery.

Keywords: micelle, solubilization, clathrate, fluorescence spectroscopy, molecular dynamics, model system, targeted drug delivery

1. Introduction

Micelles form from amphiphilic surfactant compounds in aqueous solution by aggregating with their hydrophobic tails in a roughly spherical core surrounded by a shell composed of their hydrophilic head groups. For ionic compounds, the hydrophilic shell is surrounded by an inner, ionic Stern layer, and a more diffuse, ionic Gouy-Chapman layer. Lipophilic substrates are solubilized in the hydrophobic “pseudo phase” of the micelle cores. [1]

Clathrate hydrate formation is governed by the same hydrophobic interactions that govern micelle assembly, but since the solute particles tend to be small, the water cavities surrounding them are also small. [2] The water cavities also exhibit structured, regular polyhedra composed of fully hydrogen-bonded water pentagrams and hexagrams connected in a supramolecular network. The best known examples are the natural gas hydrates that have important energy and environmental applications. [3] Closely related clathrate-hydrate crystal structures can also be formed from salts of tetra-n-butylammonium (TBA) hydroxide, where each butyl group supports the structure of one water cage. [4], [5], [6]

At this time, the number of compounds known to promote both micelle and clathrate hydrate formation is small and the number of studies of their dual nature is limited. [7] Micelles of TBA carboxylates exhibit unusually high solubility and small aggregation numbers [8], and relatively high solubilization power [7]. Micelles of TBA perfluorooctanoate have exceptional thermodynamic properties. [9]

The overall goal of this research is to develop a better understanding of the role of a clathrate-forming counterion on the structure and function of aqueous, ionic micelles. The specific objective of the work is to investigate the molecular structure of the micelle/substrate complex formed from tetra-n-butylammonium 10-undecenoate and the lipophilic dye, Nile Red, NR (The chemical structure of Nile Red is shown in figure 1.
Fig. 1: Nile Red (NR).

The 10-undecenoate anion was selected because its micelles are well characterized [10], polymerizable [11], and its oligomers have shown potential as a model for targeted drug delivery systems [12]. The solubilization power, and pH-mediated release of lipophilic substrates from TBAU micelles are also relevant features for a drug delivery system model [7], [13]. Finally, Nile Red is a lipophilic substrate with solvatochromic properties suitable for describing the polar environment of the micelle. [14], [15]

2. Materials and Methods

2.1. Materials

All solutions were made up with reagent grade, ASTM Type II water (Ricca Chemical Company). Stock solutions of sodium and tetrabutylammonium 10-undecenoate were prepared by neutralizing 10-undecenoic acid (Acros, 99%) with sodium hydroxide (Fisher NF/FCC) or tetrabutylammonium hydroxide (Acros, 40% in water) after standardizing the base with potassium hydrogen phthalate (Fisher, ACS). Nile Red (Sigma Aldrich) was technical grade. Methanol (Pharmco-Aaper), ethanol (Acros), propional (Fisher), butanol (Fisher), pentanol (Matheson, Coleman & Bell), and octanol (Acros) were all reagent grade or better. All substances were used as received, without further purification.

2.2. Fluorescence Spectroscopy

Fluorescence spectra were taken in a temperature-controlled lab (295.0 ± 0.8 K) with a Vernier Spectrovis Plus spectrophotometer. All samples were filtered (GE Whatman PVDF 0.45 μm) into glass cuvettes and measured at room temperature using an excitation frequency centered at 500 nm. The manufacturer’s specifications for accuracy and precision are 3.0 nm and 1.0 nm respectively.

2.3. Molecular Dynamics

All simulations employed Gromos 54a7 force fields [16], calculated with the Automated Topology Builder at the University of Queensland, Brisbane, Australia [17], [18], [19], and the simple point charge (SPC) water model [20]. Micelle/substrate complexes were preformed with Packmol [21], [22]. The aggregation numbers for the preformed micelles were 46 for NaU [10] and 26 for TBAU, extrapolated from [8]. Both systems were placed in dodecahedral boxes with 2.0 nm from the micelle to the faces of the box, solvated, provided with sodium ions (sodium 10-undecenoate, NaU, micelles), and equilibrated before the final production runs of 50 ns at 300 K (Gromacs 5.1.4 [23], [24], [25], [26]). Radial distribution functions were determined for the final 20 ns of each simulation with the Gromacs program, gmx rdf [27] and the snapshots were obtained with the molecular visualization program, VMD [28].

3. Results

3.2. Fluorescence Spectroscopy

Fluorescence spectra of NR in various normal alcohols were taken to correlate the wavelength of maximum peak intensity, λmax, with relative polarity of the solvent [29]. The results, plotted in Figure 2 show a very good correlation, R2 = 0.98. In addition, the measured value of λmax for methanol, 638.7 nm is in good agreement with the value of 637.9 nm reported by Stuart, et al. [14].
Fluorescence spectra for NR in NaU and TBAU micelle solutions are shown in Figure 3. For each micelle system, the values of $\lambda_{\text{max}}$ did not depend significantly on concentration. The value of $\lambda_{\text{max}}$ for NR in NaU micelles, 630.4 nm, is very close to that of butanol, 630.5 nm, indicating a relative polarity of 0.59 compared to a value of 1.00 for water. The value of $\lambda_{\text{max}}$ for NR in TBAU micelles, 638.5 nm, is very close to that of methanol, 638.7 nm, indicating a relative polarity of 0.76.

2.3. Molecular Dynamics

The snapshots of the molecular dynamics simulations in Figure 4 illustrate the qualitative differences between the two micelle/substrate complexes. Both systems show compact, roughly spherical micelles with partial exposure of the substrate at the surface. For the NaU micelle, the sodium counterions can be observed close to the micelle surface, Stern layer, as well as distributed in the more diffuse Gouy-Chapman layer. In contrast, most of the TBA ions in the TBAU micelle appear to be an integral part of the micelle surface structure. This is attributed to the hydrophobicity of the TBA counterion.
Fig. 4: Snapshots of simulations of micellar solubilization of Nile Red (red). The image on the left shows the sodium (yellow) 10-undecenoate (grey) micelle and the image on the right shows the tetrabutylammonium (blue) 10-undecenoate (grey) micelle. For clarity, water molecules are not shown in these images.

The radial distribution functions for the two counterions relative to the NR substrate are shown in Figure 5, where $g(r)$ is the probability of finding any atom of the counterion at a distance, $r$, from any atom of the Nile Red. The diffuse counterion distribution for sodium ions observed in Figure 4 are indicated quantitatively in this figure with weak peaks at about 0.5 nm (Stern layer) and about 1 nm (Gouy-Chapman layer). In contrast, the distribution for TBA ions shows two prominent peaks: an inner peak at 0.50 nm and a secondary peak at 0.84 nm. The well-defined peaks in the radial distribution function for TBA indicate a high degree of structuring for the counterions in this system.

Fig. 5: Radial distribution functions of the counterions about the Nile Red (NR) substrate. The sodium ion distribution is represented by the yellow line and the tetrabutylammonium ion distribution is represented by the blue line.

The radial distribution functions for water relative to the NR substrate in the two micelle systems are shown in Figure 6, where $g(r)$ is the probability of finding any atom of water at a distance, $r$, from any atom of the Nile Red. Again, there is very little structuring apparent for the NaU micelles, while it is pronounced in the TBAU micelles. The peak for water at 0.45 nm is just inside the TBA peak at 5.0 nm. The valley at 0.55 nm is just outside.
We believe that the correspondence of the counterion and water radial distribution functions for the TBAU micelles and the lack of similar structuring of either the counterion or water in the NaU micelles is compelling evidence that the TBA counterions cause the structuring of water at the surface of the TBAU micelles. This is consistent with the known TBA/water structuring in clathrate hydrate formation. [4], [5], [6] We believe this is not an artefact of the simulation model since it agrees with our observation of increased polarity of the environment near NR in TBAU micelles from the fluorescence spectroscopy experiments.

4. Conclusion

The picture that emerges for the sodium 10-undecenoate, NaU, micelle conforms to the standard model of a roughly spherical micelle surrounded by diffuse, but identifiable Stern and Gouy-Chapman layers. The picture of the tetrabutylammonium, TBAU, micelle is fundamentally different: the counterions are an integral part of the micelle, forming an outer shell and entrapping water molecules as part of the shell.

The packing of the TBA ions into the shell of the micelle may explain, in part, why the aggregation numbers of TBA micelles is unusually low. The hydrocarbon component of the micelle now includes a significant contribution from the counterion. The trapping of water within the micelle’s shell may be involved in the unusual thermodynamics of TBA micelles. The restructuring of water on TBA micelle formation almost certainly will affect both the enthalpy and entropy of micellization.

1. Two types of applications of micelles incorporating structured counterion/water shell can be foreseen:

   The polar nature of the shell should change the solubilization characteristics of the micelle, possibly increasing the solubilization power of the micelle for substrates with hydrophilic moieties. The shell should isolate any substrate from interaction with the environment. This would be desirable in the design of drug delivery systems.

   2. The shell should isolate any substrate from interaction with the environment.

Finally, it should be noted that the counterion/water shell, like the micelle itself is dynamic. At this temperature, the tetrabutylammonium 10-undecenoate, TBAU, micelles are very micelle-like. There is no evidence of a crystalline, clathrate-like structure for the micelle or the shell.

Acknowledgements

Emily Ogle was supported in part by the University of Cincinnati, Women in Science and Engineering (WISE) program. The molecular force fields were calculated at the University of Queensland, Brisbane, Australia.
References


