Interactions between surfactants and
\{1,4-phenylene-[9,9-bis(4-phenoxy-butylsulfonate)]fluorene-2,7-diyl\}

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Abstract

The interaction between the water-soluble anionic conjugated copolymer poly\{1,4-phenylene-[9,9-bis(4-phenoxy-butylsulfonate)]fluorene-
2,7-diyl\} (PBS-PFP) and various surfactants has been studied in aqueous solution by UV–vis absorption spectra, fluorescence and electrical
conductivity. It is suggested from the linear dependence of absorbance, fluorescence and electrical conductivity on concentration that in
the absence of surfactant, moderately stable dispersions are formed. These are affected in different ways on adding cationic, anionic or
neutral surfactants. With the cationic cetyltrimethylammonium bromide, quenching of fluorescence intensity and lifetime, and formation of
a new emission occurs at concentrations well below the critical micelle concentration (cmc). Electrical conductivity measurements indicate
a discontinuity at surfactant/polymer ratio corresponding to electroneutrality, due to complexation. With the anionic sodium dodecyl sulfate,
fluorescence quenching is also observed, but is attributed to formation of some mixed polymer/surfactant aggregate. The most striking changes
are observed with the non-ionic pentaethyleneglycol monododecyl ether (C12 E 5 ), where a blue shift in fluorescence emission, dramatic increases
in lifetime and quantum yield, and changes in electrical around the cmc are interpreted in terms of incorporation of single polymer chains in
elongated cylindrical micelles. This is supported by 1H NMR spectroscopic measurements.

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1. Introduction

Conjugated organic polymers are technologically impor-
tant materials with applications in areas such as light emitting
diodes, organic lasers, photovoltaic systems and thin film
transistors [1,2]. Particular emphasis has focused on poly-
mers and copolymers of 9,9-substituted fluorenes, since these
emit light in the blue with high efficiencies and also show
fairly high stability [3,4]. In addition, they behave as fairly
rigid “hairy-rods” [5] and also order into liquid crystalline
structures [6].

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surfactant dodecyltrimethylammonium bromide, although there is no significant change in the emission spectrum [9]. In contrast, with a neutral, water-soluble sugar substituted poly(β-phenylethylene), marked changes in emission spectra were observed on addition of various surfactants, which the authors termed surfactochromicity [13]. We have observed that with aqueous solutions of the water-soluble copolymer poly{1,4-phenylene-[9,9-bis(4-phenoxybutylsulfonate)]fluorene-2,7-diyl} (PBS-PFP, Fig. 1), a blue shift in fluorescence spectrum together with an increase in both fluorescence quantum yield and lifetime is observed on adding the nonionic surfactant pentaethylene glycol monododecyl ether (C12E5) [20]. Although the transfer of the anionic sodium dodecyl sulfate (SDS, CH3(CH2)11OSO3Na) and the neutral sodium dodecyl sulfate (SDS, CH3(CH2)10(NaCl)1Br), the anionic sodium dodecyl sulfate is not long enough to form cylindrical micelles. With these rigid rod polyfluorene derivatives, it is likely that the polymer–surfactant interactions will be somewhat different from those in more flexible polymers, where a variety of conformational changes are possible, and where association is commonly treated with a “pearl-necklace model” [21]. We present a more general study on the interaction of the anionic PBS-PFP with surfactants in aqueous solutions, considering its behaviour with the cationic cetyltrimethylammonium bromide (CTAB, CH3(CH2)15N(CH3)3Br), the anionic sodium dodecyl sulfate (SDS, CH3(CH2)11OSO3Na) and the neutral C12E5.

2. Experimental

2.1. Materials

The polymer PBS-PFP (Mw = 6500 g mol⁻¹) was synthesised by condensation of 2,7-dibromo-9,9-bis(4-sulfo-xybutoxyphenyl)fluorene (A) and 1,4-phenylenediboronic acid using Pd(PPh3)₄ as catalyst. Full details of synthesis and characterisation have been given elsewhere [20,22]. Cetyltrimethylammonium bromide (CTAB, Aldrich), sodium dodecyl sulfate (SDS, Aldrich), pentaethylene glycol monododecyl ether (C12E5, Fluka) and other reagents were of the purest grade available, and were used without further treatment. All solutions were prepared using Millipore-Q water.

2.2. Apparatus and methods

2.2.1. Spectroscopic measurements

Absorption spectral measurements were made in 1 cm quartz cuvettes on a Shimadzu UV-2100 spectrophotometer. The steady-state fluorescence spectra were measured with a Fluorolog 3-22 instrument with 0.5 nm excitation and emission bandwidths. Fluorescence decays were measured using a home-built Time-Correlated Single Photon Counting apparatus with an N2 filled IBH 5000 coaxial flashlamp or an IBH NanoLED (λexc = 373 nm) as excitation source. This was coupled to a Jobin-Ivon monochromator, Philips XP2020Q photomultiplier, and Canberra instruments time-to-amplitude converter and Multichannel Analyser. Alternate measurements (1000 counts per cycle), controlled by Decay® software (Biodinâmica-Portugal), of the pulse profile at 337, 356 or 373 nm and the sample emission were performed until 1 to 2 x 10⁴ counts at the maximum were reached [23]. The fluorescence decays were analysed using the modulating functions method of Striker with automatic correction for the photomultiplier “wavelength shift” [24]. All experiments were carried out at room temperature.

Solution electrical resistances were measured with a Wayne-Kerr model 4265 Automatic LCR meter at 1 kHz. A Shedlovsky-type conductance cell was used [25]. The cell constant (approximately 0.8465 cm⁻¹) was determined to ±0.02% from measurements with KCl (reagent grade, recrystallised and dried using the procedure and data from [26]). Measurements were made at 25.00 ± 0.01 °C in a Grant thermostatic bath.

1H NMR spectra of PBS-PFP were run on solutions in D2O on a Varian Unity 500 MHz spectrometer.

2.3. Sample preparation

As indicated in Section 3.1, dispersions of PBS-PFP in water are not stable over extended periods of time. However, at a PBS-PFP concentration around 6 x 10⁻⁴ g dm⁻³ it was possible to prepare systems which were stable enough for fluorescence and electrical conductivity measurements by keeping under continuous stirring overnight and making the relevant measurements immediately afterwards.

3. Results and discussion

3.1. Behaviour of PBS-PFP in aqueous solution

The polymer PBS-PFP did not dissolve readily in water. However, by stirring overnight it was possible to obtain a relatively stable dispersion, although on leaving a solid precipitated out of this. With freshly prepared solutions, a relatively broad absorption was observed, with a maximum at 381 nm (Fig. 2). The absorbance of this increased linearly with polymer concentration, in agreement with the Beer–Lambert law. Similarly, a linear increase in fluorescence with concentration

![Fig. 1. Structure of PBS-PFP.](image-url)
was observed. The precipitation probably results from association of PBS-PFP aggregates in solution, which have previously been reported for \( p \)-phenylene and fluorene based polymers \([4,20,27,28]\) and other conjugated polyelectrolytes \([29]\). The absorbance behaviour will be further discussed with electrical conductivity results in Section 3.3.

3.2. Fluorescence spectral and lifetime measurements

Solutions of PBS-PFP in water showed a structured fluorescence between 400 and 550 nm, which is similar to that observed with other phenyl-fluorene copolymers \([29]\). Upon addition of various concentrations of CTAB, a decrease in intensity of the fluorescence was observed and was accompanied by formation of a new emission around 525 nm. There were no changes in the emission spectra of the dominant species (Fig. 3(a)). This contrasts with the behaviour of MPS-PPV with cationic surfactants, where a dramatic increase in fluorescence intensity was observed. These differences possibly arise from the relatively rigid-rod structure of PBS-PFP, compared with the random coil MPS-PPV.

As with other polymer–surfactant systems \([21]\), all these changes occur at surfactant concentrations (ca. \( 4 \times 10^{-6} \) mol dm\(^{-3} \)) very much lower than the cmc of CTAB (\( 8 \times 10^{-4} \) mol dm\(^{-3} \) \([30]\)). The results strongly suggest complexation between the oppositely charged surfactant and polymer favoured by electrostatic interactions. However, because of the rigid structure of PBS-PFP, we feel that the polymer–surfactant aggregates have a somewhat different structure from the pearl-necklace structure seen in more flexible polymer–surfactant systems. The origin of the new band at 525 nm is discussed in detail elsewhere \([22]\). These changes were accompanied by a decrease in the fluorescence lifetime observed at the main band and appearance of a second component with rather longer lifetime at 525 nm (Table 1).

In contrast, on addition of the anionic SDS to aqueous solutions of PBS-PFP, although quenching of fluorescence occurred (Fig. 3(b)), no new bands were observed, and there was no observable change in the fluorescence lifetime of the polymer within the time resolution of our system. Again, emission maxima did not shift on addition of the surfactant.

### Table 1

<table>
<thead>
<tr>
<th>CTAB (mol dm(^{-3} ))</th>
<th>Fluorescence lifetime (ns)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Observed at 420 nm</td>
</tr>
<tr>
<td>0</td>
<td>0.36</td>
</tr>
<tr>
<td>( 3.3 \times 10^{-7} )</td>
<td>0.25</td>
</tr>
<tr>
<td>( 6.7 \times 10^{-7} )</td>
<td>0.24</td>
</tr>
<tr>
<td>( 7.9 \times 10^{-8} )</td>
<td>0.22</td>
</tr>
<tr>
<td>( 3.1 \times 10^{-8} )</td>
<td>0.27</td>
</tr>
<tr>
<td>( 2.0 \times 10^{-8} )</td>
<td>0.29</td>
</tr>
</tbody>
</table>

* Biexponential decay.

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Fig. 2. Absorption spectra of PBS-PFP in water at various concentrations. Inset: Beer–Lambert plot of absorption at 381 nm as function of polymer concentration.

Fig. 3. Fluorescence spectra of aqueous solutions of PBS-PFP upon addition of: (a) CTAB; (b) SDS; (c) C\(_{12}\)E\(_{5}\). Note blue shift in fluorescence spectra in (c) on adding surfactant. In (a) and (b) the Rayleigh scattering peak is also shown.
It would seem that although there is some aggregation of polymer and surfactant, there is no specific interaction driving this.

Very different behaviour from that with either CTAB or SDS was observed with aqueous solutions of PBS-PFP in the presence of the neutral C12E5. A blue shift in the emission maxima (Fig. 3(c)) was seen, and was accompanied by dramatic increases in both fluorescence quantum yield and lifetime (Fig. 4). We have suggested [20] that the spectral changes, which are very similar to those observed with a phenyl-fluorene copolymer on going from thin films to iso-

3.3. Electrical conductivity of solutions

Since PBS-PFP is a polyelectrolyte, electrical conductivity measurements are likely to be informative. We have studied the effect of the PPS-PFP concentration on the electrical conductivity in aqueous solution. The molar conductivity (A) was calculated using

\[ A = (\kappa - \kappa_0)/c \times 1000 \]  

where \( \kappa \) and \( \kappa_0 \) are the electrolytic conductivities of solution and \( c \) is the polymer concentration (1.577 \times 10^{-6} \text{ mol dm}^{-3}) to 1.495 \times 10^{-6} \text{ mol dm}^{-3}). The experimental electrolyotic conductivity was around 3.5 \times 10^{-3} \text{ S m}^{-1} and the molar conductivty depends on the square root of concentration, in agreement with the Kohlrausch equation [25]

\[ A = A^0 - A_0 c^{1/2} \]  

where \( A^0 = 0.362 \pm 0.011 \text{ S m}^{-1} \text{ mol}^{-1}, \) \( A = 8.50 \pm 0.27 \text{ S mol}^{-1/2} \text{ m}^{2/2} \). The molar limiting conductivity \( A^0 \) is similar to that of other polyelectrolytes in aqueous solution [32].

The facts that the behaviour of aqueous solutions of PBS-PFP, both with respect to the Beer–Lambert and Kohlrausch laws, are linear functions of concentration suggest that in the absence of surfactants the polymer aggregates may be considered as separate chemical entities.

Upon addition of CTAB to an aqueous solution of PBS-PFP (8.6 \times 10^{-6} \text{ (moles monomer)} \text{ dm}^{-3}), a linear increase in electrical conductivity was observed up to surfactant concentrations 1.35 \times 10^{-3} \text{ mol dm}^{-3}, where a change of slope and second linear region was seen. As with the fluorescence measurements, this break occurs at surfactant concentrations much lower than the cmc. Further, it is observed at a monomer surfactant ratio of approximately 1:2. Since each monomer has two anionic groups, this corresponds roughly to charge neutralisation. Results are summarised in Table 2.

Electrical conductivity measurements were also made on aqueous solutions of PBS-PFP in the presence of SDS. In this case, very different behaviour was observed from the system with CTAB, and all the electrolytic conductivity data could be fitted to a single straight line over the surfactant concentration range 3 \times 10^{-4} \text{ to } 5 \times 10^{-2} \text{ mmol dm}^{-3}. At higher concentrations, phase separation occurred. From the electrical conductivity behaviour and the fluorescence measurements, some kind of aggregation of the surfactant and polymer is occurring which reduces the free SDS concentration in solution. This may be a polymer–surfactant anionic mixed micelle.

A third type of behaviour was observed with solutions of PBS-PFP in the presence of C12E5. As both the surfactant and polymer concentrations were varied in these experiments, the results are shown in a three-dimensional plot. In this case, on adding C12E5, ionic conductivity, and hence molar conductivity, increased with surfactant concentration, with a sharp increase around 3 \times 10^{-3} \text{ mol dm}^{-3}, where micellisation starts (Fig. 5). Possible reasons for the increase in conductivity on adding C12E5 are discussed elsewhere [20]. What is important for this discussion is that the break point in the curve is very close to the surfactant cmc.

\[
\begin{array}{cccc}
\text{Surfactant} & \Delta \tau_c (\text{mM}) & A (\text{S cm}^{-1}) & B (\text{S cm}^{-1} \text{ M}^{-1}) & R^2 & \text{cmc (mM)} \\
\text{CTAB} & 3 \times 10^{-4} \text{ to } 7 \times 10^{-4} & 1.566 & 157185 & 0.998 & 0.365 \\
& 1 \times 10^{-2} \text{ to } 0.21 & 2.716 & 72220.1 & 0.998 & 0.0028 \\
\text{SDS} & 3 \times 10^{-4} \text{ to } 5 \times 10^{-2} & 1.039 & 7444.41 & 0.999 & - \\
& 3 \times 10^{-2} \text{ to } 0.20 & \text{Positive deviation from straight line} & & & \\
\end{array}
\]
Fig. 5. Effect of C12E5 and polymer concentrations on the molar conductivity of PBS-PFP solutions.

\( (5 \pm 2 \times 10^{-5} \text{ mol dm}^{-3}) \) [31]. Combining this observation with the fluorescence data gives strong evidence for a model where at surfactant concentrations above the cmc, PBS-PFP becomes incorporated as isolated, rigid-rod molecules in long cylindrical micelles. Support for this from NMR data will be given in the next section.

3.4. NMR studies

From \(^1\)H NMR spectra, self-diffusion and proton relaxation studies, it has been shown that with increasing surfactant concentration C12E5 forms elongated rod-like micelles [33,34]. This has been confirmed by dynamic light scattering [34] and small angle neutron scattering [35]. NMR studies in such systems are expected to show considerable line broadening due to slow tumbling [36]. The \(^1\)H NMR spectra were run of PBS-PFP in D2O solutions. Although in our previous report [20] we assigned sharp signals in the 6.8–7.9 ppm region (Fig. 6(a)) to the polymer, we now believe these are due to trace impurities, and that no signals attributable to PBS-PFP are observed in this region in the absence of surfactant. However, upon addition of C12E5 at concentrations above the surfactant cmc, broad signals due to the polymer are observed (Fig. 6(b)). These results are completely consistent with suggestions from the other techniques that PBS-PFP is incorporated as isolated polymer chains in elongated, cylindrical C12E5 micelles. Small angle neutron scattering experiments [37] suggest that when no surfactant is present, the polymer exists as solid-like aggregates, which provides a clear explanation for the inability to observe NMR signals under these conditions.

4. Conclusions

The interactions of surfactants with relatively rigid-rod conjugated polymers are of interest both from theoretical and practical viewpoints. The behaviour is likely to be rather different from systems involving more flexible polymers, where pearl-necklace structures are invoked. In these studies, we have shown that the anionic PBS-PFP interacts strongly with CTAB at concentrations well below the surfactant cmc. From both changes in the fluorescence behaviour and electrical conductivity as functions of surfactant concentration, we suggest that this is due to complexing of the cationic head group of the surfactant by the sulfonate group of the polymer through electrostatic interactions. It is probable that the alkyl chains of the surfactant become orientated close to the polymer backbone. It is hoped to test this through molecular simulations. With SDS, there is evidence of aggregation from both fluorescence and electrical conductivity measurements. Although we do not have any evidence for the type of species present, we suspect it may be some type of anionic mixed micelle. Much clearer indications of micellisation came from fluorescence, conductivity and \(^1\)H NMR studies on PBS-PFP in the presence of the neutral surfactant C12E6. All the results strongly support a model where isolated polymer chains are incorporated in elongated, cylindrical polymer micelles. Studies in progress of the behaviour of this polymer in various solvents are consistent with it being present as isolated chains in good solvents, but forming polymer aggregates in poor ones. Within this model, the interior of C12E5 micelles would seem to be a favourable environment for the polymer.

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