Network Formation of Catanionic Vesicles and Oppositely Charged Polyelectrolytes. Effect of Polymer Charge Density and Hydrophobic Modification

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Introduction

Rheological control and, in particular, gel-like appearance are important in many applications and have stimulated physicochemical investigations of systems with such properties.1-5 In particular, mixtures of surfactants and polymers are frequently employed for this purpose6 and, since a variety of combinations can be used, examples of possible applications are diverse and can be exemplified by detergents, shampoos, paints, and cosmetics.6,7,8 If a surfactant is added to a polymer solution, dramatic changes in viscosity are found if the polymer is of the associating type, in particular if it is a hydrophobically modified water-soluble polymer (HMP). However, the viscosity varies in a nonmonotonic way with the surfactant concentration. There is at low surfactant concentrations a major increase in viscosity. While the introduction of ca. 1% of hydrophobic grafts on a water-soluble polymer typically leads to an order of magnitude increase in viscosity for a 1 wt % solution in the absence of surfactant, the difference can amount to several orders of magnitude in the presence of low amounts of surfactant. However, viscosity only increases up to a certain surfactant concentration; thereafter it starts to decrease strongly and at high surfactant it reaches values much below the surfactant-free situation. This behavior is well understood in terms of mixed aggregate formation between the surfactant and the HMP and of the significance of stochiometry, i.e., the ratio between the number of

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surfactant micelles and the number of hydrophobic groups on the polymer molecules.\(^\text{10}\) Particularly strong association, and thus rheological effects, are found for the case of oppositely charged polymer—surfactant pairs. In this case there may, however, also be an associative phase separation over a certain concentration range.\(^\text{11}\)

A large number of investigations has been performed on the rheological and microstructural properties of mixed solutions of HMPs and surfactants, covering combinations of different types of cationic, anionic, and nonionic polymers and surfactants.\(^\text{2,12−17}\) With few exceptions these studies have been concerned with conditions where the surfactant self-assembles into small globular micelles. However, it can be expected that larger viscosities and/or more suitable rheological characteristics can be obtained with larger surfactant aggregates, like threadlike micelles, disklike aggregates, or vesicles. This was also confirmed in an investigation where micellar growth was important changes in the surfactant aggregate shape are also reported and the mechanisms lying behind it discussed.

Experimental Section

Materials. Surfactants, sodium dodecyl sulfate (SDS) and diodecyl dimethylammonium bromide (DDAB), were obtained from Aldrich and were used without further purification. The vesicles formed had an average molecular weight of 1.04 × 10^9 and an aggregation number, N_{agg}, of approximately 15 000.\(^\text{30,31}\) Thetwodifferent cationic polyelectrolytes, P^+ (J R400) and HMP^+, (LM200), are hydroxyethyl cellulose derivatives that were manufactured by Union Carbide Chemicals and Plastics Company, Inc. Approximate structural formulas are shown in Figure 1. The J R400 N,N-trimethylammonium derivative has an average molecular weight of 1.04 × 10^9 and an aggregation number, N_{agg}, of approximately 15 000.\(^\text{30,31}\) The two cationic cellulose derivatives that were used in the present investigation: JR400 = P^+; LM200 = HMP^+.

When an aqueous solution of such vesicles is mixed with an oppositely charged polymer, three different regions can be found\(^\text{28,32}\)

(I) A phase separation region, where a precipitate forms in equilibrium with a bluish solution. The phase separation is of the associative type, the precipitate being a coacervation complex of surfactant and polymer with a (near) zero net charge.

(II) A monophasic low viscous solution where there is enough excess of surfactant in solution to dissolve the precipitate.

(III) A gel region, at excess polymer, where the precipitate is redissolved.

In our work the polymer has a positive charge and at overall charge neutralization the fraction of positive charges is 0.5; we will in this paper denote such conditions by f_p = 0.50, where f_p is defined by eq 1 below. In this work the redissolution occurs at f_p = 0.54 for the system which contains the higher charged polymer (P^+), while for the system with the lower charged and hydrophobically modified polymer (HMP^+), redissolution appears at f_p = 0.57.\(^\text{28}\) The region is referred to as the gel region, because samples are highly viscous.

Previously, the association in this system has been studied by establishing a phase map.\(^\text{28}\) For a more precise microstructural basis, cryogenic transmission electron microscopy (cryo-TEM) investigations were performed and compared with novel rheological studies. It is found that the P^+ system is characterized by relatively long relaxation times compared to the lower charged HMP^+ system. On the other hand, the latter showed higher viscosity. Rather important changes in the surfactant aggregate shape are also reported and the mechanisms lying behind it discussed.

![Schematic representation of the chemical structure of the two cationic cellulose derivatives that were used in the present investigation: JR400 = P^+; LM200 = HMP^+.](Image)

\(f_p = \frac{\text{number of positive charges}}{\text{number of positive charges} + \text{number of negative charges}}\)

Figure 1. Schematic representation of the chemical structure of the two cationic cellulose derivatives that were used in the present investigation: JR400 = P^+; LM200 = HMP^+.
to a concentration of charges of about 10 mm. Since P has a molecular weight of about 500 kD, each polymer chain has an approximate contour length of 1000 nm. LM200 is a N,N-dimethyl-N-dodecylammonium derivative that on average has approximately one positive charge every 10 nm, or about 2 mm of charge in a 1 wt % aqueous solution. Since this polymer has a lower molecular weight (approximately 100 kD), the average contour length is much shorter (on average about 200 nm). Both polymers render their charge from quaternary ammonium groups; however the ammonium group on HMP also contains a dodecyl chain. The degree of hydrophobic modification of HMP is about 5.4 hydrophobic chains per 100 sugar residues.

An estimation of the surface density of the vesicle yields approximately 1 net negative charge per nm², as calculated from the equation

\[
d = \frac{(x_{SDS} - x_{DDAB})N_{agg}}{8\pi R_{ves}^{2}}
\]

where \( d \) is the density of the negative charges in the vesicles, \( x_{SDS} \) and \( x_{DDAB} \) are the SDS and DDAB molar fractions in the surfactant mixture, respectively, \( N_{agg} \) is the aggregation number of the vesicles, and \( R_{ves} \) is the vesicle average radius. The molar fraction of polymer charge in the mixture, \( f_p \), is defined as

\[
f_p = \frac{C_p}{C_p + C_s}
\]

where \( C_p \) is the polymer charge molal concentration and \( C_s \) is the molar concentration of the net surfactant charge (\( C_s = C_{SDS} - C_{DDAB} \)).

**Methods.** Samples were prepared in the following way. A 55 mm surfactant stock solution (\( C_{surf} = C_{SDS} + C_{DDAB} \)) was prepared by mixing a SDS micellar solution and a DDAB homogenized dispersion. This resulted in a bluish solution containing vesicles bearing anionic net charge. Vesicle solutions were sonicated for 5 min. Surfactant–polymer samples were prepared by mixing solid polymer and vesicle stock solution in appropriate amounts to obtain samples along lines defined by constant surfactant content (varying polymer content) or constant polymer content (varying surfactant content).

For an easier way to correlate with the phase map, Figure 2, mixed samples are denoted in the figures and legends as JxS, where J and L represent P (JR400) and HMP (LM200), respectively. S is the polymer weight content (wt %), and y is the surfactant concentration (mm).

Most of these samples have a high viscosity and were therefore prepared in the following way: The samples were mixed with the aid of a vortex mixer before heating to a temperature of about 50 °C for approximately 30 min. After heating, the samples were again subject to vortex mixing. With increase of temperature, the viscosity of the samples decreases, which facilitates mixing. In samples containing HMP (LM200), it has actually been found that without heating, a mixture of cloudy domains and clear gel areas appears. Gentle temperature cycles do not involve important input of energy other than to disperse colloidal entities. After three to five repeated cycles (mixing–heating–mixing), depending of sample consistency, the samples were centrifuged at 25 °C for approximately 6 h to remove air bubbles. All samples submitted to rheological tests were monophasic.

A Reologica StressTech rheometer equipped with an automatic gap setting was used for the rheological measurements. The instrument can be run either in the controlled stress or in the controlled strain mode. A cone and plate geometry was employed, and to minimize evaporation, a solvent trap system was used. Prior to any oscillatory deformation test, the linear viscoelastic region for each sample was determined by stress sweep tests.

Measurements within the linear region ensure that the measured rheological properties are independent of the applied stress and (near) equilibrium properties are probed.

In addition to oscillatory shear measurements, dynamic viscosity and shear-thinning were also studied by rotational shear tests. All experiments were run at a constant temperature of 25 °C.

A Philips CM120 Bio-twin microscope was used for cryogenic transmission electron microscopy (cryo-TEM) studies. Samples were placed in a vitrification chamber in which temperature and humidity can be controlled. The sample chamber was set at controlled 25 °C (ca. 5 mL) was put on a copper grid supported carbon coated holey film. Immediately after blotting with filter paper to obtain a thin liquid film over the grid, the sample is plunged into liquid ethane (at its melting temperature). The vitrified film is then transferred under liquid nitrogen to the electron microscope.

**Results**

**Polyacation–Vesicle Complexes and Phase Behavior.** The pseudoternary surfactant system SDS/DDAB/...
water is used as a basis for the present investigation.\textsuperscript{31,38} (Any system of two electrolytes without common ion must from a thermodynamic point of view be treated as a quaternary system. Thus separating phases do not have compositions falling in the triangular diagram representing the isothermal mixing of water and the two surfactants. Here we are concerned with the heterogeneous solution phases, where this complication does not enter.) By themselves the two surfactants have critical packing parameters (CPPs) that suggest formation of aggregates curving toward oil (CPP\textsubscript{SDS} = 0.33 and CPP\textsubscript{DDAB} = 0.8). This is largely due to the ionic headgroups of these molecules. By mixture of anionic SDS with cationic DDAB, the effective headgroup area in the mixed aggregates decreases since opposite charges favor closer packing. At certain mixing ratios, lamellar structures form and a limited vesicular region can be found.\textsuperscript{31} In the present study \(X_{\text{SDS}} = 0.71\) was chosen; here small unilamellar vesicles (SUVs) are the dominant aggregates while multilamellar vesicles (MLVs) are sparsely detected. At a higher concentration (\(C_{\text{SDS}} > 50\) \text{m}) a heterogeneous region with a bluish solution and a cloudy phase separation of lamellar liquid crystal appears. For low concentrations (\(C_{\text{SDS}} < 5\) \text{m}), a catanionic solid (precipitate or flocs) is present in equilibrium with solution.\textsuperscript{31}

In a previous study, the effect of adding two polycations to different surfactant compositions in this system has been reported.\textsuperscript{28} The same (pseudoquaternary) polymer–surfactant systems were used here, and interestingly it was found in the present investigation that the presence of poly electrolyte prevents phase separation at surfactant concentrations lower than 5 \text{m} (this has been checked down to 2 \text{m}). Composition maps with macroscopic appearance of samples indicated are shown in Figure 2. The line indicates \(f_{p^+} = 0.50\).

Rheological Characterization. Samples within the gel region have been investigated with respect to the rheological behavior.

Figure 3 shows typical mechanical spectra for the two polymer systems, with storage, \(G'\), and loss, \(G''\), moduli, as functions of the oscillation frequency. Following other associating polymer systems, the storage and loss moduli increase with increasing frequency, i.e., with decreasing observation time. The HMP\textsuperscript{+} system shows a commonly observed behavior of \(G''\) being larger at low frequencies and \(G'\) at high frequencies, i.e., of a progressive changeover from dominating viscous effects at low frequencies to dominating elastic properties at high frequencies. From this behavior, a formation of a three-dimensional network can be deduced and the lifetime of the cross-links can be estimated from the inverse of the angular frequency at which \(G'\) and \(G''\) cross. For the P\textsuperscript{+} samples, in general \(G''\) exceeds \(G'\) over the entire frequency range accessible; i.e., the cross-link lifetimes are very long. This behavior was found for almost all P\textsuperscript{+} samples, except for low surfactant and polymer concentrations, where relaxation times could be easily measured. Typically thus the P\textsuperscript{+} system displays a gel-like behavior for all studied frequencies; HMP\textsuperscript{+} samples are viscoelastic with a gel-like behavior at high enough frequencies and a liquidlike behavior for low frequencies. On the other hand, HMP\textsuperscript{+} samples show higher storage moduli than the system composed of the polycation without hydrophobic modification. Analogous observations were found from the analysis for other conditions, i.e., 5, 7.5, 10, 12.5, 15, and 20 \text{m} surfactant concentration (in the P\textsuperscript{+} system) and 2, 2.5, 5, and 6.5 \text{m} surfactant concentration (in the HMP\textsuperscript{+} system). Table 1 illustrates the variation of the relaxation times and storage moduli at high frequencies with composition. In the case of the P\textsuperscript{+} system, some of the relaxation times (indicated in Table 1 as a lower limit) are either below the accessible frequency or display a noisy regime. Both relaxation time and storage modulus generally increase with polymer and surfactant concentration, as seen in Table 1. However, a maximum in the storage modulus and relaxation time is observed in the surfactant concentration sweep and will be discussed below.

Figure 4 shows the dependence of \(G'\) on the polymer concentration (at a constant surfactant concentration of 5 \text{m}). An increase of the polyelectrolyte concentration, and thus of the number of possible bridges, is accompanied by an increase of the elastic and loss moduli. This was seen particularly at intermediate and high frequencies.

Concerning the P\textsuperscript{+} system, samples are generally characterized by a larger \(G''\) than \(G'\) in the investigated frequency range. Thus, relaxation times are long and all samples either are self-supporting (test tubes can be tilted upside down without sample flow) or flow very slowly, as is illustrated in Figure 5. For low amounts of polymer and surfactant, for instance 0.455, elastic properties are only displayed at high frequencies.

With HMP\textsuperscript{+}, the situation is quite different. Despite \(G''\) moduli having larger numerical values, the relaxation times are shorter, as seen by a crossover of \(G'\) and \(G''\) within the studied frequency interval. The HMP\textsuperscript{+} polymer thus can be seen to give mixtures with higher viscosity as compared to the P\textsuperscript{+} polymer that gives more elastic mixtures (longer lived cross-links). A rheological comparison of the two polymers in samples with the same composition (and far away from the phase separation line) was performed and underlines the described tendencies. The presence of equal weight percentage of polycation signifies that the number of positive charges is five times larger in the P\textsuperscript{+} sample and that there are five times as
many HMP$^+$ chains as there are P$^+$ chains; the total length of P$^+$ chains is obviously similar to that of HMP$^+$ chains.

The storage modulus was also compared for high frequencies (again HMP$^+$ has higher storage modulus than P$^+$), which gives information about the number of active links (bridges).\(^{(39)}\) Note that $G_\infty$ (plateau of $G'$ at high frequencies) is proportional to the bridge density; some stiff gels (as in our case) cannot achieve the plateau and $G'$ at high frequencies is used as a factor to give qualitative information about the active chain density.

The complex viscosity, $\eta^*$, and the storage modulus, $G'$, are larger in the HMP$^+$ system, which gives evidence for the importance of the hydrophobic modification. P$^+$ samples are more elastic, as seen by a longer relaxation time which is even outside the frequency interval studied, and, thus, significantly longer than the relaxation time for HMP$^+$.

Figure 6 shows the storage modulus as a function of $f_p^+$ and vesicle volume fraction, $\phi_v$, calculated from an expression relating vesicle volume, surfactant concentration, $C_{surf}$, and aggregation number, $N_{agg}$

$$\phi_v = 4\pi R_{ves}^3 C_{surf}N_A/3N_{agg}$$  \hspace{1cm} (3)
where \( N_{\text{agg}} \) is defined as

\[
N_{\text{agg}} = \frac{8\pi R_{\text{ves}}^2}{a_s} \tag{4}
\]

where \( a_s \) is the average surfactant headgroup area, \( R_{\text{ves}} \) is the vesicle radius, \( N_A \) is Avogadro's number, and \( C_{\text{surf}} \) is the total surfactant concentration. Equations 3 and 4 can be related to obtain the expression 26,31

\[
\phi_v = a_s R_{\text{ves}} N_A C_{\text{surf}} / 6 \tag{5}
\]

The total polymer concentration is 2 wt \% for the \( P^+ \) system and 2.5 wt \% for the HMP\(^+\) system. A value of \( a_s \) of 35 Å\(^2\) has been used, as justified previously.\(^{31}\) Note that the range of \( \phi_v \) in Figure 6 is very different for the two polymers. When compared under similar conditions, the storage modulus is considerably higher for HMP\(^+\) than for \( P^+ \). Higher \( f_p^+ \) values correspond to lower vesicle volume fractions; this is more evident for HMP\(^+\) system, since HMP\(^+\) has lower charge density than \( P^+ \). Additionally, Figure 7 represents storage modulus as a function of \( f_p^+ \) for 1 wt \% of \( P^+ \).

From both figures it can be inferred that first the storage modulus increases with surfactant concentration. However, at higher surfactant contents it passes via a maximum; after that it decreases, indicating a weaker network.

An analogous behavior is found in the shear rate dependence of viscosity, exemplified in Figure 8. This figure also demonstrates that all the samples are shear-thinning with a Newtonian plateau at low shear rates. The progressive addition of surfactant leads to stronger gels, as seen as an increase of Newtonian plateau. The sample J2S27.5 is an exception since it is close to the neutralization line.

It was also found that in the \( P^+ \) system, the viscoelastic properties increase monotonically on increasing the total polymer/surfactant concentration, as illustrated in Figure 8.
Network Formation of Catanionic Vesicles

since aggregate shapes are strongly altered by an in-
tvesicles and SUVs (Figure 10c), while in the samples of
surfactant sample, J1S2, there is coexistence of faceted
aggregates coexist with faceted vesicles and holey vesicles
(Figure 10e, f).

Figure 9. Complex viscosity and storage modulus as a function of polymer concentration for the P•+ system. It can be seen that both properties increase monotonically as the concentration is increased; f0•+ is kept constant (f0•+ = 0.6557).

9, for the case of keeping f0•+ constant (at f0•+ = 0.6557, equidistant to the neutralization line).

cryo-TEM investigation of microstructure. To complement the rheological measurements, the micro-
structural effects connected with the polymer–vesicle association have been investigated by cryogenic trans-
mision electron microscopy. For this technique, during sample vitrification the temperature is so rapidly lowered
that amorphous rather than crystalline ice is obtained; in this way, self-assembly structures generally remain intact.

For the case of vesicles and other large structures, artifacts due to rapid freezing should then be quite small. A more
serious difficulty consists of obtaining shear-free thin films
in the cryo-grid with using highly viscous samples, such as those found for HMP•+. Therefore, studies for this
polymer had to be limited to a range of polymer concentrations lower than those of the gel-like samples. The
polymer effect on the surfactant self-assembly should, nevertheless, be a local effect rather than associated with
the long-range formation of a network. Therefore, we believe that the main structural effects of polyelectrolyte addition
can be correctly assessed and, in particular, the differences between the two polymers used.

Previous cryo-TEM investigations of these systems have shown that the polyelectrolytes induce a modification in
the vesicle shape and integrity, in particular for the P•+ system.28 In the current study, a larger number of samples
have been extensively monitored by the imaging technique. Figure 10 shows representative micrographs, where
typical features are indicated by arrows. The polymer-
free vesicle solution shows the presence of mainly unilamellar, roughly spherical vesicles (Figure 10a). It can be
seen that HMP•+ associates with the vesicles and creates
only small effects on their shape (Figure 10b); dispersed
spherical vesicles are shown, where they undergo minor
changes in their size and shape.28 The TEM images give
evidence for a strong interaction of P•+ with the vesicles,
since aggregate shapes are strongly altered by an in-
creasing amount of polymer. In 1 wt % P•+ and 2 mm
surfactant sample, J 152, there is coexistence of faceted
vesicles and SUVs (Figure 10c), while in the samples of
2 wt % P•+ with 10 mm surfactant (J 2510), dislike
aggregates coexist with faceted vesicles (Figure 10d). At
2 wt % P•+ with 20 mm surfactant (J 2520), dislike
aggregates coexist with faceted vesicles and holey vesicles
(Figure 10e, f). At 0.4 wt % P•+ and 7.5 mm surfactant
(J 0.457.5), the mixture is close to phase separation
boundary and large irregularly shaped bilayer patches
can be observed (Figure 10f).

There is no obvious increase in size of vesicles with
either polymer. The polymer-free vesicular solutions
contain unilamellar vesicles with diameters in the range
20–70 nm and an average size of 30 nm; with polymer
addition the vesicle size distribution observed in the
pictures remains apparently unchanged. The cryo-TEM
images also indicate the possibility to stabilize vesicle
solutions at 2 mm of surfactant when the polymer HMP•+
is present, whereas the polymer-free samples at concentra-
tion below 5 mm are phase-separated (solution + precipitate).

Discussion

The general and most significant findings in the rheological characterization of the mixed polymer–vesicle
systems investigated in this work can be summarized as
follows: (i) the hydrophobically grafted polycation, HMP•+,
affects the viscosity and the storage and loss moduli to a
larger extent than the hydrophilic polycation, P•+; (ii)
the number of cross-links is larger with HMP•+ than with P•+;
(iii) the cross-links are considerably more long-lived with
P•+ than with HMP•+. From the variations of the rheological
parameters with composition, we suggest that these
findings are quite general, i.e., they do not depend on the
particular polymer samples used here, for example, they
are not due to the differences in molecular weight.

Otherwise, the current systems show several character-
istic features of polymer solutions in general and as-
sociating polymer solutions in particular, such as shear-
thinning behavior and viscoelastic properties with
mitating viscous effects for long observation times and
dominating elastic effects for short times.

The observed rheological properties can be related to
two main factors: the polyelectrolyte–vesicle interactions and the effects of the polymer on surfactant self-assembly.

Let us start by addressing the first factor. The polymer–
vesicle interactions at play are of electrostatic and
hydrophobic origin. For P•+, only electrostatic interactions
account for association, whereas for HMP•+ the hydrophobic
grafts can be directly inserted in the vesicle bilayer. The
surface charge density of the vesicles has been estimated
to be of the order of 1 charge per nm2 (cf. eq 1). The P•+
polymer with a high charge density and a mean charge-
to-charge contour length of about 2 nm can efficiently
match the vesicle surface charge. For the lower charge
density HMP•+, where the charge separation is of roughly
10 nm, the situation is less optimal for the interaction
with the anionic vesicles. Moreover, P•+ has a longer overall
particle size than HMP•+ and thus can more effectively
wrap around vesicles, with long contact regions along the
polymer chain, and even connect two or more vesicles.

Previous studies show that in systems of strong electro-
static association with low content of screening electrolyte,
the interaction forces can yield gels with very long
relaxation times.40

For the hydrophobically modified polymer, the charged
stickers can easily anchor in the bilayer, but the smaller
number of charges by which the polymer is attached to
each vesicle implies shorter relaxation times. Association
of different parts of the polymer molecule with several
vesicles may result in the formation of a three-dimensional
polymer–vesicle network; the electrostatic interactions

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Polymer systems, vesicles with regions of zero or almost molecular packing in the surfactant bilayer. For both interaction but also the local vesicle charge and the charged polymer will affect not only the vesicle is essential for thermodynamic stability. An oppositely where equilibrium vesicles are present and a net charge polymer addition. Such changes are relevant in systems vesicle structure undergoes considerable changes upon aggregate structure and emphasizing the differences between the two polymer systems discussed so far.

Besides the polymer-vesicle interactions, modifications on vesicle structure, fluidity, and integrity due to the association could also underpin the rheological behavior. The cryo-TEM imaging unequivocally shows that the network formation in the P+/vesicle system (left) and in the HMP+/vesicle system (right). For simplicity, the polymer chains are represented in a stretched way.

Reinforce to some extent the strength of association. An increase of surfactant concentration promotes closer surfactant aggregates and hence the possibility of cluster formation. This could explain the increase of relaxation time since the cluster network is more protected against deformation. Moreover, the increase of surfactant, and therefore the number of close vesicles, increases the number of bridges between the polymers and the surfactant aggregates, which causes an increase of the storage modulus, G'. Figure 11 summarizes in a schematic way the association model proposed here, assuming no changes in aggregate structure and emphasizing the differences between the two polymer systems discussed so far.

A high charge density polymer is expected to affect an oppositely charged vesicle in two main ways. First, the polymer charge will affect the curvature of the surfactant films. A surfactant film with a positive curvature due to a net charge will have its curvature reduced on charge neutralization. Second, the conformational state of the surfactant alkyl chains can also be changed. For ionic surfactants, it is known that the Krafft point, or the chain melting temperature in the case of bilayer-forming amphiphiles, are strongly affected by electrostatic effects, e.g., addition of electrolyte or increase in counterion valency. The screening of the headgroup repulsions by an electrolyte stabilizes both the crystalline and the micellar/bilayer state, but the effect is more pronounced for the more organized state where the area per molecule is smaller. Hence, the addition of electrolyte typically increases the Krafft point or the chain melting temperature. A polyion, which can be seen as a giant counterion, is similarly expected to increase the melting temperature and thus induce surfactant crystallization in the vesicles in a certain temperature range.

A freezing of the alkyl chains, with loss of liquidlike motional freedom, affects surfactant aggregation profoundly. The formed network probably inhibits crystal packing and sedimentation, keeping the solidlike surfac-
tant in solution. If upon freezing the vesicles remain kinetically trapped, there may be changes in the surfactant films to a less fluid state, a consequence of which may be the formation of planar regions in the vesicles—hence the observed faceted vesicles. An alternative mechanism could be an in-plane surfactant segregation induced by the polymer “adsorption” onto the bilayer, owing to the preferential interaction of the polycation with the anionic surfactant SDS. The consequent heterogeneous surfactant packing in the bilayer plane could also induce morphological changes. Segregation effects have been previously seen for mixed ionic–nonionic lipid vesicles upon interaction with an oppositely charged polymer. However, such effects are probably less likely in our system since surfactant–surfactant interactions here are stronger than for an ionic–nonionic surfactant film.

Formation of hollow icosahedra in catanionic surfactant mixtures has been detected for crystallized chains in previous studies, where it was also found that the aggregates were stabilized by the presence of pores located at the vertexes of theicosahedra. Such states would be less stable at high polymer concentration, and vesicles would be easily disrupted. A first consequence could then be the formation of holes in the vesicles; a second more dramatic consequence could be the fragmentation of vesicles into more or less planar bilayer fragments or disks. Disks seem to be the preferred arrangement of the surfactant aggregates when the polymer positive charge concentration is high enough.

The insertion of the charged grafts, in the case of the HMP + seems to be less disruptive of the original vesicle shape. In studied charged vesicle systems where low molar fractions of an oppositely charged surfactant have been added, only a small increase in the chain melting temperature is observed and the vesicle structure remains intact. One could expect by comparison that a polymer with a relative small number of charged hydrophobic chains inserted in the surfactant films will have a relatively minor effect on vesicle shape and fluidity.

These changes in the state of the surfactant aggregates can be expected to influence the formation and properties of the polymer–surfactant network. For instance, a liquid–solid transition for the surfactant film in the network would have the effect of considerably lengthening the lifetimes of cross-links. The high relaxation times deduced for the P + system could then be explained, not only by optimal polymer–vesicle interactions but also by the crystallization of the surfactant in the vesicles. The detachment of a polycation from a crystallized vesicle surface would have a high free energy of activation. Regarding the formation of disclike structures, one can speculate that an additional crowding effect, perhaps similar to the card-house effect found for clays, could well contribute to an enhanced network formation and strength. HMP + has a lower charge density, and the effects on the surfactant packing are weaker than with P +. Disks are detected to a negligible extent for the HMP + system, and considerably fewer vesicles acquire a pronounced faceted shape compared to those in P + mixtures. Figure 12 illustrates schematically the relation between the aggregates formed in the two polymer systems and the polymer concentration.

One final point to be discussed is the viscosity maximum as a function of composition, observed for the HMP + system.

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Figure 12. A schematic representation of the surfactant aggregate shape in the presence of (A) P + and (B) HMP +, from relative low polymer concentrations (left pictures) to relative high polymer concentrations (right pictures).

(Figures 6–8). For mixed systems of a HMP and a micelle-forming surfactant, the rheological properties show a markedly nonmonotonic behavior with a maximum as a function of composition; as mentioned above this is related to the stoichiometry, i.e., the ratio between hydrophobic groups on the polymer molecules and the number of micelles. In the vesicle case, the aggregation number is orders of magnitude higher than in the case of micelles and a simple estimate shows that the same mechanism cannot apply. Instead we see that the decrease in viscosity occurs as the two-phase region indicating associative phase separation is approached. Previous microscopy observations show the presence of large irregularly shaped “membrane-like” fragments at fP + = 0.55, suggesting that these are precrystallites. Therefore, we interpret the decrease as a result of onset of microscopic phase separation and association that reduces the extension of the polymer chains.

It should be pointed out that the picture of the possible mechanisms underlying the rheology and network formation of the polymer–vesicle systems described here remains largely speculative at this stage. Inferences were made essentially from the comparison of polymer structures and from the inspection of microscopy data. Clearly, a more in-depth picture requires the collection of more experimental data for these systems (or even variation of the cosolute chemistry). In this respect, we are currently carrying out calorimetric and rheological measurements as a function of temperature and composition, to be presented in a forthcoming report.

Conclusions

A highly viscous gel-like mixture is obtained by mixing anionic vesicles with two different cationic polyelectrolytes. Variations in polyelectrolyte and surfactant concentration strongly influence gel properties. For the more highly charged polymer, an elastic gel is found and relaxation times are long, explained by both the charge density and a proposed surfactant crystallization in the vesicles. With the shorter and less charged hydrophobically modified polyelectrolyte, the number of cross-links is higher, while the mixtures have shorter relaxation times. cryo-TEM shows that the vesicle structure is affected by the polycations in different ways. Polygon-shaped vesicles and disclike
fragments appear due to the presence of the hydrophilic polyelectrolyte since the polycation is likely to induce a crystallization of the vesicle surfactant film and thus reduce its flexibility. The higher charged polyelectrolyte induces more marked changes in the vesicle structure than the lower charged polymer.

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