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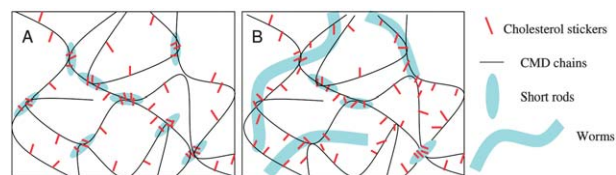
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Associative networks of cholesterol-modified dextran with short and long micelles

Hala Afifi, Marcelo A. da Silva, Cécile Nouvel, Jean-Luc Six, Christian Ligoure and Cécile A. Dreiss*

Short (rod-like) and long (wormlike) micelles based on the surfactant polyoxyethylene cholesteryl ether associate with cholesterol-modified dextran to form solid-like networks.



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■ Associative networks of cholesterol-modified dextran with short and long micelles

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The strong associative behaviours between cholesterol-modified dextran (CMD) and short and long polyoxyethylene cholesteryl ether (ChEO₁₀) micelles were investigated using rheology and small-angle neutron scattering (SANS). In solutions of short rod-like micelles (ChEO₁₀ alone), the addition of 5.0 wt% CMD induced a remarkable transition from a Newtonian system to a highly solid-like viscoelastic network, with an increase in zero-shear viscosity of over 5 orders of magnitude. The frequency sweeps at ChEO₁₀ concentrations above 2.5 wt% were fitted to a Maxwell model with 3 elements and, quite remarkably, fell onto a single master curve, while no network was formed at 2.5 wt% micelles. Viscoelastic solutions of wormlike micelles (WLMs) were obtained by adding the co-surfactant triethylene glycol monododecyl ether (C₁₂EO₃) to ChEO₁₀ solutions at a constant ChEO₁₀/C₁₂EO₃ ratio of 5/1. The introduction of CMD into the WLM solutions induced a transition to a more liquid-like behaviour (G''/G' increased), however both moduli increased by one order of magnitude. At the lowest ChEO₁₀ concentration (2.5%), the solid-like behaviour was lost. Overall, a comparable rheological response was obtained for the WLM and the short rods with CMD, however the WLM/CMD behaviour suggested a wider spectrum of relaxation processes, longer relaxation times and higher plateau moduli. SANS data from the polymer/micelles mixtures displayed a strong structural peak and were remarkably identical for both ChEO₁₀/CMD and ChEO₁₀/C₁₂EO₃/CMD systems, suggesting a very similar type of network structure, independently of the initial size of the micelles. Overall, all the results taken together show a very high affinity between the polymer and the micelles and suggest a breakup of the WLM induced by CMD. The resulting network is constituted by polymeric chains connected by micellar aggregates through hydrophobic interactions between the micellar cholesterol cores and the pendent cholesterol moieties of the polymer.

Introduction

Wormlike micelles (WLMs) are very long, semi-flexible surfactant aggregates, which, above a system-dependent concentration, known as the overlap concentration, entangle into a transient network, imparting remarkable viscoelastic properties to their solutions.^{1–3} Their behaviour is analogous to the one observed in polymer solutions, however unlike polymers wormlike micelles constantly break and reform, providing additional relaxation mechanisms. One obvious advantage over polymers therefore is that they are stable under high shear rates and can recover their supramolecular structure. They have therefore been referred to as ‘living polymers’.^{4,5} The rich rheological response of WLM has

led to their exploitation in various industrial processes such as fracturing fluids in oil fields, cooling fluids, liquid dish-washing detergents and personal care products such as shampoos and body soaps.^{6–12}

Similarly, a rich rheological behaviour has been observed in solutions of associative polymers or hydrophobically modified polymers (HMPs), which are water-soluble polymers with hydrophobic pendant groups^{13–21} (often referred to as ‘stickers’). Interest in HMP primarily started for applications in oil recovery and rheological control of paints and coatings^{22–25} but their interest now extends to the biomedical field.^{26–28}

In view of the above, the combination of HMP and wormlike micelles seems to be an attractive option. In fact, it has been shown that combining WLM and hydrophobically modified polymers can lead to significantly enhanced viscoelastic properties compared to the individual components at the same concentration.^{29,30} This has obvious environmental and cost consequences as rheological control can be achieved with less material. Surfactant/polymer interactions have been studied intensively in the past decades and are used in numerous technological applications,^{31–36} but most of these studies have dealt

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1 with low concentrations of surfactant or spherical micelles, while
a much more limited number of reports can be found on polymer/wormlike micelle interactions.^{14,29,30,34,37–43}

5 Polymers can either destroy or strengthen the transient
networks formed by wormlike micelles. Polyethylene glycol and
poly(propylene glycol) for instance were reported to induce the
disruption of the micellar network in aqueous solutions of
potassium oleate³⁴ and cetyltrimethylammonium salicylate,⁴⁴ due
to the transition from cylindrical to spherical micelles. This
behaviour was attributed to the wrapping-up of the hydrophobic
polymer chains around the surface of the surfactant micelles. As
the surface/volume ratio is larger for spherical micelles than for
cylindrical, rod-to-sphere transition takes place, optimising the
shielding of the polymer from water. In other scenarios, oppositely
charged polymer added to the anionic surfactant potassium oleate
favoured micellar growth instead,³⁷ while the hydrophobic
polymer poly(propyleneglycol) led to a drastic reduction in the
viscoelastic properties of cetyltrimethylammonium bromide
wormlike micelles without notable structural change.⁴⁰

20 With hydrophobically modified polymers (HMPs), where the
hydrophobic moieties are distributed along the polymeric chains,
different effects on the wormlike micelle structures have been
reported, which are reflected by changes in the rheological
properties.⁴¹ For instance, HMP can cross-link WLM by intercalation
of the stickers into the WLM cores, forming a network where
some relaxation modes are linked to sticker pullout time.^{41,45} The
opposite behaviour—surfactants cross-linking the polymer—has
also been observed.¹⁴ In this case, the WLMs break down and
reform as shorter (spherical or cylindrical micelles) in order to
better shield the stickers from the solvent.

30 In this contribution, we report a considerable enhancement of
the rheological response in a system based on biocompatible and
biodegradable components, namely, the association between
cholesterol-modified dextran (CMD) and the surfactant polyoxyethylene
cholesteryl ether (ChEO₁₀). We investigate and compare two types
of networks formed in the presence of the modified dextran: one
formed with short (rod-like) micelles and one formed with long
entangled wormlike micelles.

35 Polyoxyethylene cholesterol surfactants (ChEO_n) are unique
nonionic surfactants which are characterised by a strong segregation
between their hydrophilic headgroups and hydrophobic tails compared
to conventional alkyl ethoxylated surfactants.⁴⁶ Long EO-chain
cholesterol surfactants form ellipsoids or short cylindrical micelles.^{47,48}
However, when they are mixed with short-chain polyoxyethylene
alkyl ether-type nonionic surfactants, such as triethylene glycol
monododecyl ether (C₁₂EO₃) or monoglycerides, they form
viscoelastic solutions of long wormlike micelles.^{49–51}

50 Dextran is a biocompatible polymer which was modified in
order to optimise the associative interactions with the surfactant's
tails, through hydrophobic substitution by cholesterol stickers.
By blending CMD and ChEO₁₀ micelles, cholesterol-cholesterol
interactions between the pendant units of the modified polymer
and the hydrophobic cores of the micelles were expected to take
place, leading to synergistic effects in the rheological response
of the polymer/micellar mixtures.

55 In the first part of this paper, the rheological behaviour of both
the short (ChEO₁₀) and long (ChEO₁₀/C₁₂EO₃) micelles with
CMD is studied and compared using both oscillatory and steady

shear experiments. Remarkable changes in the viscoelastic
properties of both systems are observed upon addition of CMD,
demonstrating a strong association between the polymer and the
micelles. In the second part, small-angle neutron scattering
measurements are used to bring structural insight into the
networks. The combination of the results from both techniques
enables us to propose a picture of the organisation of the polymer
chains and micelles.

10 Materials and methods

Materials

Polyoxyethylene cholesteryl ether (ChEO₁₀) was purchased from
Ikeda Corporation, Yokohama, Japan. Triethylene glycol
monododecyl ether (C₁₂EO₃) was obtained from Aldrich
Chemical Co. Ltd., UK. Dextran T40 ($\bar{M}_n = 33\,800\text{ g mol}^{-1}$,
polydispersity index (I) = 1.27, as characterised by size exclusion
chromatography coupled to multi-angle laser light scattering
(SEC-MALLS) in water (0.1 M NaNO₃, 6.15×10^{-3} M NaN₃)
was purchased from Pharmacia Biotech and dried under reduced
pressure at 100 °C overnight. Cholesterol and 1,6-hexyl diisocyanate
were purchased from Aldrich and used without any further
purification. Toluene (VWR), pyridine (Aldrich) and dimethyl
sulfoxide (DMSO, Fischer scientific 99%) were vacuum-distilled
before use.

All solutions of surfactants, polymers and their mixtures were
made in D₂O (99% purity, Sigma Chemical Co. UK), except those
for the cryo-TEM measurements which were prepared in deionised
water. All chemicals were used as received unless stated. A glass
pycnometer was used to measure the density of ChEO₁₀.

35 Samples preparation

All surfactant solutions were prepared by mixing specific
amounts of stock solutions of ChEO₁₀ and C₁₂EO₃ (20 wt%) in
D₂O at room temperature and were vortex-mixed. The polymer-
containing samples were prepared using a 10 wt% stock solution
of the CMD. All solutions were left to rest at least 48 hours at
room temperature and were equilibrated at 25.0 °C (in a water-
bath) for 24 hours before conducting the rheological
measurements.

45 Polymer synthesis

The synthetic pathway of the cholesterol-modified dextran
(CMD) is a two-step procedure. First, cholesteryl *N*-(6-isocyanatohexyl)carbamate
was synthesised using a procedure previously established⁵² followed
by its condensation with dextran T40. Cholesterol (10 g, 25.8 mmol)
was reacted with 1,6-hexyl diisocyanate (62.5 mL, 0.37 mol) in 256 mL
of dry toluene in the presence of 5 mL pyridine at 80 °C for 48 hours.
The reaction was monitored by thin-layer chromatography (TLC)
and after the reaction was completed the solvent was removed
in vacuo. After adding 600 mL of petroleum ether, the residue
was stored overnight at −10 °C. The precipitate obtained was
separated and dried *in vacuo* at 30 °C overnight to give a white
powder of cholesteryl *N*-(6-isocyanatohexyl)carbamate.

The second step in the synthetic procedure was carried out by adding 0.169 g (0.3 mmol) of cholesteryl *N*-(6-isocyanatohexyl) carbamate to a solution of 4 g dextran (24.8 mmol equiv.) in 100 mL dry dimethyl sulfoxide (DMSO) containing 8 mL pyridine, the reaction mixture was left to stir for 8 hours at 80 °C. The disappearance of cholesteryl *N*-(6-isocyanatohexyl)carbamate was monitored by TLC. Ethanol (500 mL) was added to the reaction mixture and the resulted suspension was left at 4 °C overnight. After filtration, the precipitate was purified by dialysis against ethanol/water (1 : 1) for 48 hours, followed by pure water for another 48 hours. The obtained suspension was then freeze-dried to give a white powder of cholesterol-modified dextran (CMD) with a degree of substitution of 1 cholesterol group per 100 glucose units as determined by ¹H NMR in CDCl₃.

Rheological measurements

Steady shear and small-amplitude dynamic shear measurements were performed on a dynamic strain-controlled rheometer ARES (TA instruments) using cone-and-plate geometry (25 mm, 0.02 rad), with a Peltier unit for temperature control and a solvent trap to minimise evaporation. Samples were allowed to rest before the start of the experiment to ensure dissipation of any pre-shearing due to loading.

The frequency sweep tests were performed within the linear viscoelastic regime (determined previously by dynamic strain sweep measurements). Measurements were carried out in duplicate or triplicate, with very good reproducibility. The results reported in this work are examples of typical data obtained, not averages. The zero-shear viscosity (η_0) was obtained by extrapolation from the flow curves. Values of the relaxation time τ_r and plateau modulus G_0 were obtained by fitting the frequency sweeps with a multi-element Maxwell model:

$$G'(\omega) = \sum_k G_k \frac{\omega^2 \tau_k^2}{1 + \omega^2 \tau_k^2} \quad (1)$$

$$G''(\omega) = \sum_k G_k \frac{\omega \tau_k}{1 + \omega^2 \tau_k^2} \quad (2)$$

where G' and G'' are the storage and loss moduli, respectively; G_k and τ_k are the G_0 and τ_r for the k^{th} element, respectively; ω is the oscillation frequency and G_0 is defined as:

$$G_0 = \sum_k G_k \quad (3)$$

The fitting was done using the non-linear fitting tool from Microcal Origin™ 6.0 software. Initially, a set of parameters were obtained for the G' curves; this set was then used as initial parameters for fitting G'' and the results were used to generate a new fit for G' data. This process was repeated until a single set of parameters was obtained that provided the best fit for both moduli (which was not necessarily the best fit for each modulus individually). The purpose of the fitting with multiple elements is to provide a more quantitative interpretation of the rheological data. The number of elements chosen was based solely on the minimal number needed to obtain a good fit and they are not attributed a physical meaning. However, they give a useful measure of the departure from the pure Maxwell behaviour.

Samples for rheological measurements were vortex-mixed and kept in a water bath at 25 °C for 24 hours to ensure equilibration before performing the measurements. Samples were made in D₂O to ensure that they were equivalent to the samples used for SANS measurements.

SANS measurements

Small-angle neutron scattering experiments were performed on a LOQ instrument at ISIS pulsed neutron source (ISIS, Rutherford-Appleton Laboratory, STFC, Didcot, Oxford) and on a V4 instrument at the Hahn-Meitner Institute, Berlin, Germany. The LOQ uses incident wavelengths from 2.2 to 10.0 Å, sorted by time-of-flight, with a fixed sample–detector distance of 4.1 m, which provided a range of scattering vectors (q) from 0.009 to 0.29 Å⁻¹. V4 covers a scattering vector (q) range from 0.004 to 0.37 Å⁻¹; the neutrons are collected on a two-dimensional position sensitive detector which was placed at 1, 4 and 12 m horizontally from the sample.

The samples used in the SANS experiments were prepared in D₂O to optimise the contrast with the protonated surfactants and polymer. The samples were placed in clean disc-shaped quartz cells (Hellma) of 1 and 2 mm path length and the measurements were carried out at 25 °C. All scattering data were first normalised for sample transmission and then background-corrected using a quartz cell filled with D₂O (this process also removes the inherent instrumental background arising from vacuum, windows, *etc.*) and finally corrected for the linearity and efficiency of the detector response using instrument-specific software package. The data were then converted to the differential scattering cross-sections (in absolute units of cm⁻¹) using the standard procedures at ISIS⁵³ and HMI.

SANS data fitting

Data presenting a typical Kratky behaviour in a [$q^2 I(q)$ vs. q] representation ('bell-shaped' scattering curve with a linear slope at low q) were tested against the Kratky–Porod wormlike chain model, which describes the scattering from elongated objects characterised by their local rigidity.^{54,55} In the q range $q > b^{-1}$, where b is the statistical unit length (or persistence length), the scattering cross-section of an infinitely thin wormlike chain is expressed by:

$$\frac{\partial \Sigma}{\partial Q}(q) \approx \frac{\pi c (\Delta \rho)^2}{N_A \delta^2} \times \frac{M_L}{q} \quad (4)$$

where $\Delta \rho$ is the contrast, M_L the mass per unit length, equal to (M/n), where n is the number of segments and M the total mass. This expression can be modified by incorporating a Guinier-like factor to take into account a finite cross-section.⁵⁴

$$\frac{\partial \Sigma}{\partial Q}(q) \approx \frac{\pi c (\Delta \rho)^2}{N_A \delta^2} \times \frac{M_L}{q} \times \exp\left(-\frac{q^2 R_g^2}{2}\right) \quad (5)$$

where R_g is the cross-sectional radius of gyration, which is related to the cross-sectional radius by $R = \sqrt{2} R_g$. By plotting the data as $\ln\left(q \times \frac{\partial \Sigma}{\partial Q}(q)\right)$ vs. q^2 , a linear plot is obtained with a gradient of $\left(-\frac{R_g^2}{2}\right)$ and an intercept of $\ln\left(\frac{\pi c (\Delta \rho)^2 M_L}{N_A \delta^2}\right)$.

SANS data were also fitted using a model for cylinders. For N randomly oriented rods of length L and radius R the form factor $P(q)$ is given by:

$$P(q) = N \int_0^{\pi/2} F^2(q) \sin(\gamma) d\gamma \quad (6)$$

$$\text{with } F(Q) = (\Delta\rho)V \frac{\sin\left(\frac{1}{2}qL \cos \gamma\right)}{\frac{1}{2}qL \cos \gamma} \frac{2J_1(qR \sin \gamma)}{qR \sin \gamma} \quad (7)$$

where $J_1(x)$ is the first order Bessel function and γ is the angle between the q vector and the axis of the rod.

The data modelling program FISH⁵⁶ was used to fit the SANS data. FISH uses standard iterative least-squares fitting in which selected parameters of the chosen model can be refined to optimise the fit. Parameters were refined from several starting points to ensure that a global (rather than a local) minimum was found. A pre-factor, referred to as the ‘scale’ factor, containing parameters such as volume fraction and contrast was left to float and the value returned by the fit checked against its calculated value to confirm consistency of the fit. A margin of $\pm 20\%$ difference was deemed acceptable. Samples were measured in D₂O to maximise the contrast. Scattering length densities of $4.19 \times 10^{-7} \text{ \AA}^{-2}$ and $1.33 \times 10^{-8} \text{ \AA}^{-2}$ were used for ChEO₁₀ and C₁₂EO₃, respectively. A volume-averaged scattering length density was calculated for the wormlike micelles, using the experimental volume fractions of each surfactant.

Cryogenic Transmission Electron Microscopy

Cryo-TEM measurements were performed with a Zeiss TEM 902A instrument, operating at 80 kV in zero-loss bright-field mode. Digital images were recorded under low dose conditions with a BioVision Pro-SM Slow Scan CCD camera system. An underfocus of 1–2 μm was used in order to enhance the contrast. The preparation procedure has been described in detail elsewhere.⁵⁷ Specimens for examination were prepared in a climate chamber with temperature and humidity control (temperature 25 °C and a relative humidity of approximately 98–99%). Thin films of sample solution were prepared by placing a small drop of the sample on a copper grid supported perforated polymer film, covered with thin carbon layers on both sides. After the drop was blotted with filter paper, thin sample films (10–500 nm) spanned the holes in the polymer film. Immediately after blotting, the samples were vitrified by plunging them into liquid ethane, held just above its freezing point. Samples were kept below $-165 \text{ }^\circ\text{C}$ and protected against atmospheric conditions during both transfer to the TEM and examination. Several images were taken of each sample studied, which were all consistent (a representative example is presented here).

Results and discussion

In this work, we compare the effect of cholesterol-modified dextran (CMD) on the associative structures and viscoelastic properties of two types of micelles: short micelles (ChEO₁₀) and wormlike micelles (formed by mixtures of ChEO₁₀ with C₁₂EO₃).

The polymer concentration is kept constant at 5.0 wt%. The micellar concentration (based on the amount of ChEO₁₀) is varied from 2.5 to 10 wt%, while keeping ChEO₁₀/C₁₂EO₃ weight ratio constant at 5 : 1. Upon mixing the different stock solutions, highly viscoelastic solutions form almost instantaneously. In the first part of this study, the linear and non-linear rheological properties of these solutions are discussed. In the second part, the results from SANS measurements on the same systems are analysed, from which a structural arrangement of the micelles/polymer networks is proposed.

Rheological properties

Viscoelastic behaviour. In this study, we compare both the linear and non-linear rheological behaviour of the ChEO₁₀/CMD and ChEO₁₀/C₁₂EO₃/CMD systems. ChEO₁₀ has been found to aggregate in solution as ellipsoids or short rods of *ca.* 170 \AA length.⁴⁸ Solutions of ChEO₁₀ micelles show a Newtonian behaviour with a viscosity of *ca.* 2 mPa s at 25 °C, hence very close to pure water. The polymer solutions display a very weak shear-thinning response with a viscosity only slightly higher than water (data not shown), even at 5.0%. Unlike other HMPs which have been found to self-assemble and lead to a strong viscoelastic response,^{58,59} CMD does not form a network on its own. Instead, its solutions are slightly turbid, with the presence of particles visible to the eye, showing limited solubility of the polymer, probably due to aggregation driven by hydrophobic interactions between the cholesterol pendant groups. Note that dextran T40, which was modified to give CMD, is well below the entanglement molecular weight at the concentration studied,¹⁷ which therefore should also be the case for CMD.^{17,60} We therefore do not expect to see relaxation processes due to polymer reptation in our systems. The addition of 5.0% CMD to ChEO₁₀ or ChEO₁₀/C₁₂EO₃ solutions at concentrations above 2.5% induces a remarkable increase in the viscoelastic properties of the micellar solutions and leads to the disappearance of undissolved polymeric particles, as already observed for other HMP/surfactant micelles mixtures.⁶¹ At 2.5% micellar concentration instead, the short ChEO₁₀ solutions remain liquid-like, while the ChEO₁₀/C₁₂EO₃ wormlike solutions, originally solid-like, become more liquid-like upon addition of the polymer.

Fig. 1 shows the frequency sweep curves for the ChEO₁₀/CMD systems studied in this work: 5.0% CMD with 5.0, 7.5 and 10% ChEO₁₀. All samples show a characteristic viscoelastic behaviour, reminiscent of the Maxwell model. However, the data do not fit to a single-element Maxwell model and a multi-element model was used instead. The straight lines show the fits obtained for a 3-element Maxwell model (Fig. 1). The values of the parameters obtained are reported in Table 1. There is a very clear, nearly linear increase of the moduli G_{01} , G_{02} , G_{03} and the relaxation times τ_{r1} , τ_{r2} , τ_{r3} with the concentration of ChEO₁₀. For simplicity, the rest of the discussion focuses on G_0 , the sum of all three moduli obtained from the fits (eqn (3)), and on τ_r , the longest relaxation time, which is substantially higher than the other two, and correlates with the hydrophobic junctions lifetime.⁶² This clear trend with concentration is reflected by the construction of a master curve from all three mechanical spectra, obtained by dividing all frequencies by the frequency value at the cross-over point, where $G' = G''$, and equally dividing all the

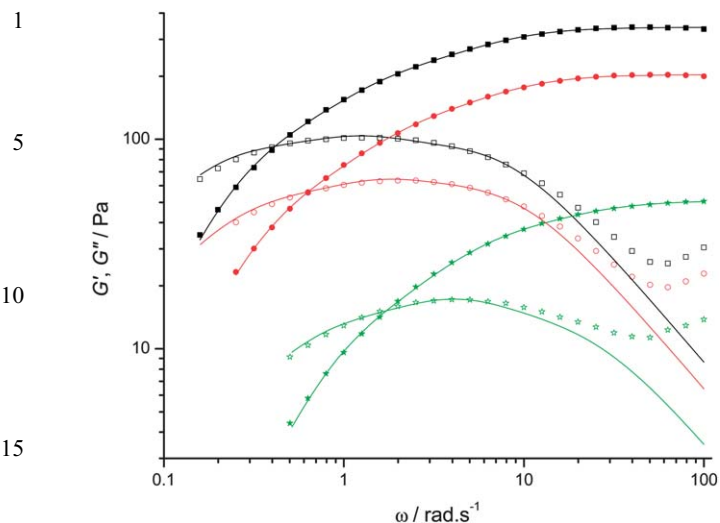


Fig. 1 Variation of the storage modulus G' (filled symbols) and loss modulus G'' (open symbols) as a function of the oscillatory shear frequency in solutions of 10% (■), 7.5% (●) and 5.0% (★) ChEO₁₀ with 5.0 wt% CMD. Solid lines are fits to the Maxwell model with 3 elements.

Table 1 Summary of the rheological parameters extracted from the frequency sweep measurements and viscosity curves of ChEO₁₀/CMD and ChEO₁₀/C₁₂EO₃/CMD solutions at 10%, 7.5% and 5.0% of ChEO₁₀ and fixed ChEO₁₀/C₁₂EO₃ ratio (5 : 1)

	ChEO ₁₀ /CMD			ChEO ₁₀ /C ₁₂ EO ₃ /CMD		
	5.0%	7.5%	10%	5.0%	7.5%	10%
G_1/Pa	14.7	59.3	109	20.2	29.4	54.3
G_2/Pa	23.7	75.6	123	31.4	34.0	60.9
G_3/Pa	12.5	68.6	110	28.8	56.9	62.8
G_4/Pa	—	—	—	23.7	55.3	66.1
G_5/Pa	—	—	—	14.6	24.5	138
G_0/Pa	50.9	204	342	119	200	382
$G''_{\text{dip}}/\text{Pa}$	11.3	19.6	25.5	19.4	43.9	—
τ_{r1}/s	1.2	2.9	4.0	52.2	72.0	34.0
τ_{r2}/s	0.24	0.61	0.81	6.70	5.55	4.70
τ_{r3}/s	0.05	0.14	0.16	0.92	0.730	0.69
τ_{r4}/s	—	—	—	0.15	0.120	0.097
τ_{r5}/s	—	—	—	0.02	0.013	0.008
τ_r/s	1.2	2.9	4.0	52.0	72.2	34.0
$\eta_0/\text{Pa s}$	29.9	367	688	—	—	—

moduli by the respective moduli value at the cross-over point. The curve obtained is shown in Fig. 2. All the frequency sweep curves overlap remarkably well, except at very high frequency, thus suggesting that the rheological changes observed are linked to a concentration effect rather than structural rearrangements.⁶³⁻⁶⁵ As mentioned before, CMD aqueous solutions are turbid, suggesting the presence of aggregates of collapsed coils. CMD/ChEO₁₀ solutions instead are free of particles. This suggests that the micelles protect CMD's hydrophobic stickers from the solvent, allowing the coils to expand. Increasing the amount of surfactant correlates directly with an increase in active junction sites, by binding of the stickers from the polymer into the micelles, thus demonstrating a strong affinity of the polymer stickers for the micelles, as they associate with them rather than with themselves.^{64,66} This linear dependence of the rheological response with ChEO₁₀ concentration also supports the absence

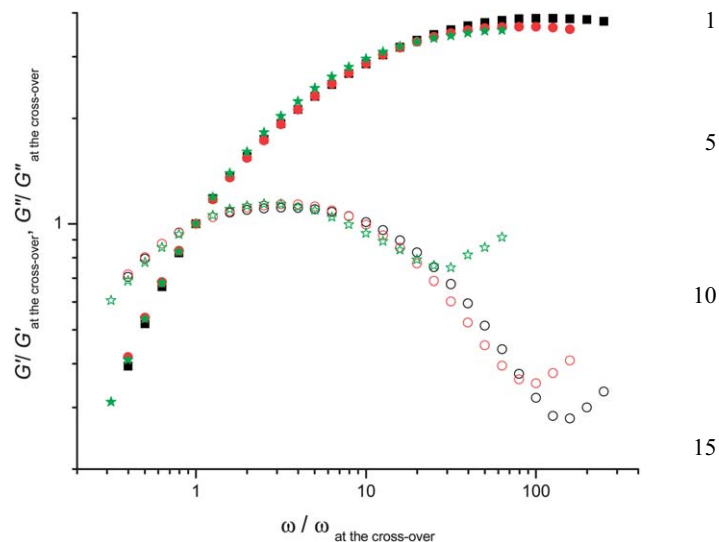


Fig. 2 A master curve of the three curves presented in Fig. 1: solutions of 10% (■), 7.5% (●) and 5.0% (★) ChEO₁₀ with 5.0 wt% CMD. The master curve was obtained by dividing all frequencies by the frequency value at the cross-over point ($G' = G''$), and equally dividing all the moduli by the respective moduli value at the cross-over point.

of entanglements; relaxation processes seem to be dominated by the association/dissociation of the hydrophobic stickers from the micellar aggregates. Note however that this concentration effect breaks up at 2.5% micelles, since the solutions remained liquid-like after adding the polymer, suggesting that this concentration of micelles is insufficient to provide connectivity. The master curve also highlights the departure from the model at high frequencies. For the systems studied here, the Maxwell behaviour holds as long as the relaxation processes are dominated by the dynamics of the junctions: as the frequency increases, this process becomes less important, while breathing or Rouse-like motion starts to dominate.⁶⁷ In the frequency sweeps, the transition is marked by a sudden increase of G'' in the high frequency range, deviating from the expected behaviour of the Maxwellian model. This departure taking place at the same frequency for all samples reflects the fact that while rheological parameters such as τ_r and G_0 depend on ChEO₁₀ concentration, the transition to breathing or Rouse-like relaxation processes depends on the nature of the supramolecular structures and not on their concentration.

The combination of ChEO₁₀ with C₁₂EO₃ leads to the growth of short ChEO₁₀ micelles into elongated wormlike micelles, as shown previously.^{48,49} A more detailed structural characterisation of each type of micelles is presented in the next section (SANS measurements). For these WLM solutions, the ratio of surfactant/co-surfactant was kept constant at 5 : 1 (corresponding to the zero-shear viscosity peak) and ChEO₁₀ was varied from 2.5 to 10 wt% with 5.0 wt% of CMD, as for the shorter micelles.

The frequency sweeps for ChEO₁₀/C₁₂EO₃ alone are presented in Fig. 3. The systems behave as viscoelastic solids within the frequency range investigated with no visible cross-over point for G' and G'' . Upon addition of the polymer, both moduli G' and G'' increase by one order of magnitude, however the solutions also become less solid-like with a larger G''/G' ratio throughout most of the frequency ranges studied. With the 2.5% ChEO₁₀/C₁₂EO₃

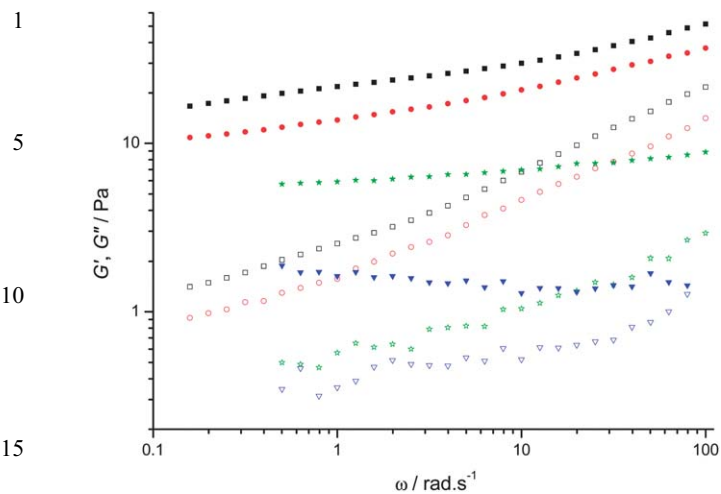


Fig. 3 Variation of the storage modulus G' (filled symbols) and loss modulus G'' (open symbols) as a function of oscillatory shear frequency in solutions of 10% ChEO₁₀/2.0% C₁₂EO₃ (■), 7.5% ChEO₁₀/1.5% C₁₂EO₃ (●), 5.0% ChEO₁₀/1.0% C₁₂EO₃ (★) and 2.5% ChEO₁₀/0.5% C₁₂EO₃ (▼).

system however, the addition of CMD induces a transition from a viscoelastic solid to a viscoelastic liquid, therefore showing the most significant contrast between ChEO₁₀/C₁₂EO₃ and ChEO₁₀/C₁₂EO₃/CMD systems. This drop in rheological response can be explained either by drastic structural changes, such as the breakup of micellar connectivity, or as dynamic changes giving rise to faster relaxation processes, such as branching.⁶⁸ As mentioned in the Introduction, both situations have been reported in WLM and polymer mixtures,^{34,37,40,43} and it is difficult to distinguish one case from another based solely on the rheological data. However, given the striking change in the frequency sweep data (Fig. 4), together with the absence of

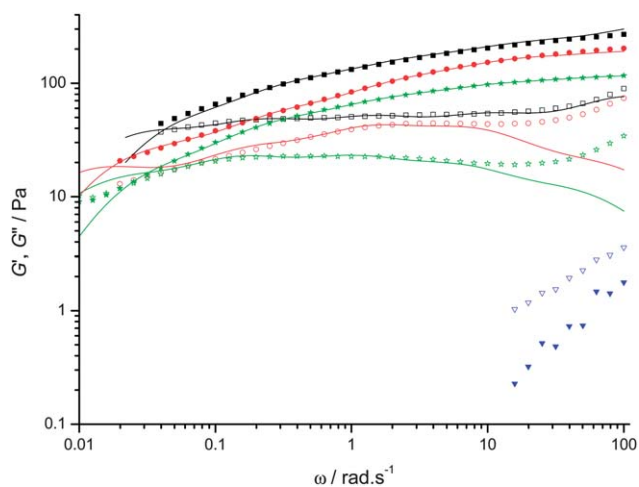


Fig. 4 Variation of the storage modulus G' (filled symbols) and loss modulus G'' (open symbols) as a function of oscillatory shear frequency in solutions of 5.0% CMD with 10% ChEO₁₀/2.0% C₁₂EO₃ (■), 7.5% ChEO₁₀/1.5% C₁₂EO₃ (●), 5.0% ChEO₁₀/1.0% C₁₂EO₃ (★) and 2.5% ChEO₁₀/0.5% C₁₂EO₃ (▼). Solid lines are fits to the Maxwell model with five elements.

a network formation at the same concentration of ChEO₁₀ short micelles (thus showing that this amount of short micelles is insufficient to provide junction points), we favour the hypothesis of a breakup of the WLM induced by CMD. SANS data presented in the next section give further support to this assumption. For all concentrations above 2.5%, the addition of CMD leads to a behaviour closer to the Maxwell model compared to the pure WLM, not too dissimilar to the one observed for the ChEO₁₀/CMD systems (Fig. 1), but with a larger spectrum of relaxation processes. Despite the obvious departure from the Maxwell behaviour, fits to a multi-elements Maxwell model were also performed here, purely as a mathematical tool, in order to extract and compare plateau modulus and relaxation time values and ‘quantify’ departure from the pure Maxwell model. Five elements were indeed needed to fit the frequency sweep curves (Table 1), which are presented in Fig. 4 as solid lines. The quality of the fits was much lower than for the ChEO₁₀/CMD system, even with the two additional elements. This clearly suggests a system with a wide range of relaxation times. This could possibly be explained by the presence of long, entangled micelles that are “sticked” by HMP, inducing several extra reptative relaxation modes of the entangled micelles by analogy with those predicted by the sticky reptation theory.^{69,70} However another feature of this system that distinguishes it from the previous one (short micelles/CMD) is the presence of the co-surfactant, which may also influence the junctions lifetime. As for the ChEO₁₀/CMD system, we attempted to construct a master curve from the normalised frequency sweeps (Fig. 5). Due to the lack of a well-defined cross-over, the normalisation was based on an arbitrarily chosen frequency. The master curve for this system however does not show such a good superposition as the one observed for the ChEO₁₀/CMD systems (Fig. 2). This suggests that the changes seen in the rheological behaviour between 5.0% and 10% WLM

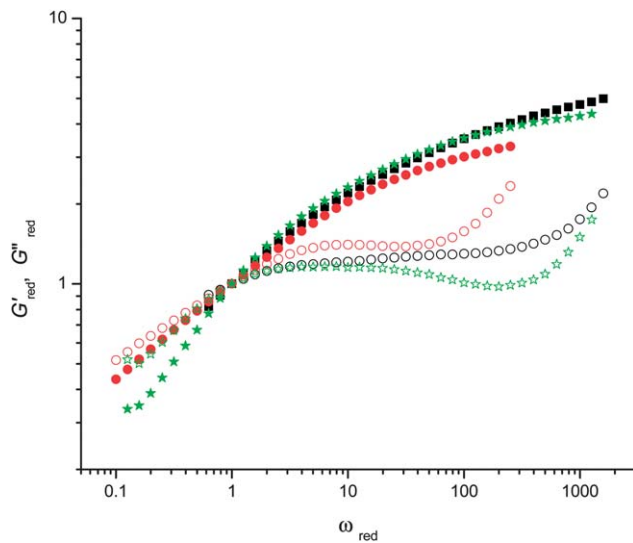


Fig. 5 A master curve of all three curves presented in Fig. 4 of solutions of CMD (5.0 wt%) with 10% ChEO₁₀/2.0% C₁₂EO₃ (■), 7.5% ChEO₁₀/1.5% C₁₂EO₃ (●) and 5.0% ChEO₁₀/1.0% C₁₂EO₃ (★). The master curve was obtained by dividing all frequencies by an arbitrarily chosen frequency, and equally dividing all the moduli by the respective moduli value at the chosen frequency.

are not simply due to concentration effects. Given the complexity of the mixtures, it is difficult to pinpoint the structural differences which lead this result, however some hypotheses can be proposed. In the 2.5% WLM system, the change from a viscoelastic solid to a viscoelastic liquid suggested that CMD reduces end-cap energy, leading to micellar break up, therefore suppressing, or strongly reducing, their entanglement. Under this assumption then, it would be expected that at lower polymer/surfactant ratio (*i.e.* increasing ChEO₁₀ from 2.5% to 10%), micelles long enough to entangle may still be present. Therefore at increasing concentration, not only the number of polymer/micelles associative junctions is affected, but also the length of the micelles present, thus the balance between relaxation due to junction association and dissociation on one hand and micelles reptation is modified. If we suppose instead that micelles have been shortened down enough to suppress reptation, the possibility of a distribution of lengths, leading to slightly different junction dynamics (compared to the short micelles) could explain the non-linear change with concentration.

In the ChEO₁₀/CMD solutions, while all relaxation times τ_r followed a clear linear dependence with surfactant concentration, the same is not observed for the ChEO₁₀/C₁₂EO₃/CMD systems, where τ_r at the intermediate ChEO₁₀ concentration (7.5%) is higher than at the highest ChEO₁₀ concentration (10%), and τ_r at 10% of the surfactant is slightly lower than that at 5.0% (Table 1). The relaxation times are also much longer than for the equivalent short micelles system (ChEO₁₀/CMD), by an order of magnitude (Table 1). In comparison also, the breathing or Rouse modes are less clearly visible, although a slight upturn of G'' at high frequency would suggest that they are still present. Unlike the relaxation time, there is a clear trend in the plateau modulus G_0 , which increases with concentration across the range studied.

Without yet making any assumption on the type of network formed, it is quite clear that junctions are created between the dextran cholesterol pendant groups and the hydrophobic tails of the surfactants, also cholesterol groups. The different viscoelastic behaviour between the two systems can be attributed to the different dynamics between short (rod-like) micelles and longer (broken wormlike) micelles: wormlike micelles have slow dynamics, implying longer hydrophobic junction lifetimes, thus longer relaxation times (Table 1). In addition, the density of connections is expected to be higher with the longer micellar aggregates than with the shorter near-spherical ones. The same long micelle could connect in several points along its length to either one or several polymer chains, thus giving rise to a combination of relaxation processes, both from the polymer and the longer micelles. These observations are reflected by the values of G_0 , which relates to the junction density.⁷¹ In both systems, G_0 increases with surfactant concentration and tends to reach similar values for both ChEO₁₀/C₁₂EO₃ and ChEO₁₀/C₁₂EO₃/CMD systems (G_0 values at 5.0% are quite different, while they are much closer at 7.5% and 10%, *cf.* Table 1). This reflects the different junction potential of both micellar aggregates: for the longer micelles, a larger number of connection sites is already established at the lowest concentration ($G_0 = 118$ Pa against 50.3 Pa for ChEO₁₀/CMD); as the concentration increases, the difference between G_0 for both systems is drastically reduced at the highest concentration ($G_0 = 382$ Pa against 341 Pa for ChEO₁₀/CMD).

The large deviation from the Maxwell behaviour for the ChEO₁₀/C₁₂EO₃/CMD when compared to the ChEO₁₀/CMD is not clear, but not surprising since the combination of long micelles with polymer is likely to give rise to a large spectrum of relaxation times. The nature of the distribution of the hydrophobic side chains along the polymeric backbone, probably irregular, may create local micro-domains with specific dynamics. For solutions of telechelic associative polymers and surfactants, a two-element Maxwell behaviour has been determined and was attributed to the existence of two relevant dynamics affecting the relaxation process: disruption of the hydrophobic junction and reptation of the micellar aggregates.⁴³ For telechelic associative polymers alone, one single relaxation time is expected, related to the dissociation of the hydrophobic junctions.⁷² Curiously enough, both surfactant/polymer systems show a behaviour closer to the expected Maxwellian behaviour than the wormlike micelles alone, strongly suggesting that the polymer itself directs the overall system dynamics through the hydrophobic junction dissociation. Also, as mentioned above, in the ChEO₁₀/C₁₂EO₃ systems at 2.5% ChEO₁₀ concentration, CMD induces remarkable changes in the rheological behaviour, which we suggest is due to the loss of network connectivity caused by the breakup of the micellar aggregates.

Flow behaviour. The ChEO₁₀/CMD and ChEO₁₀/C₁₂EO₃/CMD systems studied in this work present a typical shear-thinning behaviour (Fig. 6). For the ChEO₁₀/CMD systems, a Newtonian plateau is observed in the lower shear rate range, up to a critical shear rate value where shear-thinning starts. The zero-shear viscosities (η_0) determined from the Newtonian plateau are very high (Table 1): at 10% ChEO₁₀/CMD, the viscosity is 688 Pa s, an increase of over 5 orders of magnitude compared to the pure micellar solutions (*ca.* 2 mPa s) at the same concentration, showing a remarkable synergistic effect. The viscosity increases linearly with concentration, from 5.0 to 10%.

The zero-shear viscosity (η_0) and the plateau modulus (G_0) are given by, respectively, $\eta_0 = \sum_k G_k \tau_k \leq G_0 \tau_r$ ⁷³ and $G_0 = v_{\text{eff}} RT$,⁷¹ where R is the universal gas constant, T is the temperature and

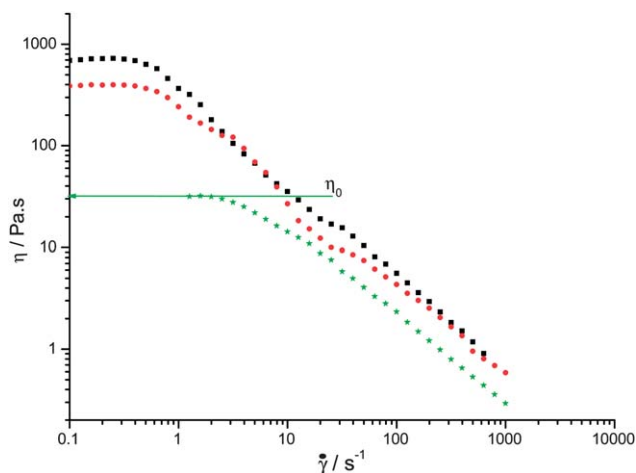


Fig. 6 Steady shear viscosity as a function of shear rate for a fixed concentration of CMD (5.0 wt%) and 10% (■), 7.5% (●) and 5.0% (★) ChEO₁₀.

ν_{eff} is the number of effective or active elastic chains, which is related to the junction density. Thus, the η_0 dependence on ChEO₁₀ concentration reflects the longer relaxation times and the higher junction density as the surfactant concentration increases. By using the relationship between G_0 and ν_{eff} it is possible to estimate a mesh size ξ , with $\xi \approx \sqrt[3]{1/\nu_{\text{eff}}}$. For the short micelles (with $G_0 = 50.9, 206$ and 342 Pa), we obtain $\xi \approx 432$ Å, 271 Å and 229 Å for 5.0, 7.5 and 10%, respectively. For the long micelles ($G_0 = 119, 353$ and 382 Pa), we obtain 326 Å, 227 Å and 201 Å, respectively, therefore (as expected) a decrease in mesh size with increasing micellar concentration and slightly denser networks with the long micelles, however more similar at the highest concentration studied (10%).

The flow curves for the ChEO₁₀/C₁₂EO₃/CMD system (Fig. 7) also show a typical shear-thinning behaviour. However, within the range of shear rates studied, no Newtonian plateau was observed. This may be a reflection of the larger increase of both relaxation time and entanglement density from ChEO₁₀/CMD to ChEO₁₀/C₁₂EO₃/CMD systems, increase larger enough to shift the Newtonian plateau to a lower shear region, outside the experimental window. The flow curves of the WLM without CMD are also quite similar (Fig. 7): the solutions shear-thin, without a measurable Newtonian plateau. The addition of polymer however increases the viscosity values by one order of magnitude, except at 2.5% of ChEO₁₀/C₁₂EO₃ where the presence of CMD causes a drastic viscosity reduction. This behaviour suggests that the CMD may disrupt the micellar aggregates, breaking them below entanglement level, and the amount of micelles and polymer in solution is below the percolation threshold⁴³ (as observed for 2.5% rod-like micelles). The small-angle neutron scattering results in the next section bring more support to this argument.

At low shear rates, the major contribution to viscosity comes from the junction points. As the shear rate increases, the macromolecules will orient under shear, possibly leading to

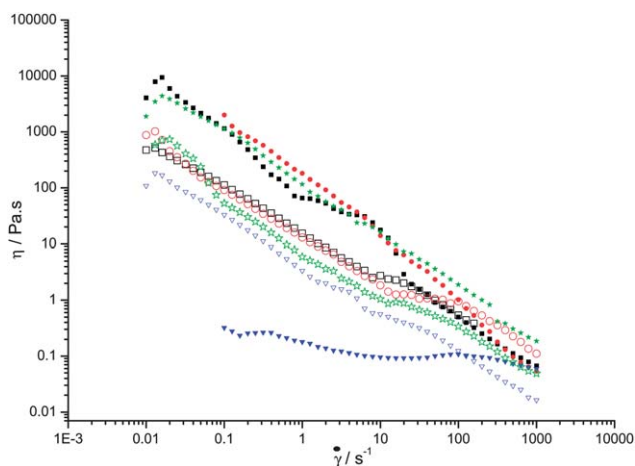


Fig. 7 Steady shear viscosity as a function of shear rate for a fixed concentration of CMD (5.0 wt%) and 10% ChEO₁₀/2.0% C₁₂EO₃ (■), 7.5% ChEO₁₀/1.5% C₁₂EO₃ (●), 5.0% ChEO₁₀/1.0% C₁₂EO₃ (★), 2.5% ChEO₁₀/0.50% C₁₂EO₃ (▼) and for the wormlike micelle solutions of 10% ChEO₁₀/2.0% C₁₂EO₃ (□), 7.5% ChEO₁₀/1.5% C₁₂EO₃ (○), 5.0% ChEO₁₀/1.0% C₁₂EO₃ (☆), 2.5% ChEO₁₀/0.50% C₁₂EO₃ (▽).

a decrease of the lifetime of junction points.⁷⁴ At high enough shear rates, the dissociation of the junction points may lead to a breakup of the supramolecular structures. Thus at very high shear rates the hydrodynamic contribution of the oriented remaining aggregates is likely to be the major component, instead of the network junctions. The viscosity values at high shear for the ChEO₁₀/CMD and ChEO₁₀/C₁₂EO₃/CMD systems fall within a small region (1.9 to 5.6 Pa s at a shear rate of 100 s⁻¹), with the exception of the 10% wormlike micelle system (ChEO₁₀/C₁₂EO₃), which shows a value lower by an order of magnitude (~ 0.5 Pa s at 100 s⁻¹). This suggests that, once the bridging between the supramolecular aggregates is broken, the remaining elements are of similar size and shape for both systems within the concentration range studied, except perhaps for 10% ChEO₁₀/C₁₂EO₃, where we may have smaller aggregates. The shoulder seen in the ChEO₁₀/C₁₂EO₃/CMD at 10% of ChEO₁₀ flow curves can be attributed to the formation of shear-induced structures⁷⁵ or shear-banding,⁷⁶ as this flow instability at intermediate shear rates is a signature pattern of these types of phenomena and has been observed both in wormlike micelles^{77,78} and associating polymer solutions.^{79,80}

The rheological data show the remarkable effect of introducing CMD into the micellar solutions, both for the short and the long micelles. The strong synergy observed denotes very favourable associations between the micellar aggregates and the polymer, which is not surprising since their hydrophobic moieties are identical. Bridging between the polymer and the micelles is expected, which could be of two types, as described in the Introduction: in one scenario, the polymer creates bridges between large entangled micellar aggregates,⁴¹ in another scenario, smaller spherical or rod-like micellar aggregates act as bridges between expanded polymer coils.¹⁴ We could envisage the first type of network for the wormlike micelles (ChEO₁₀/C₁₂EO₃) and the second for the pure ChEO₁₀ shorter micelles. However the rheological data alone do not provide a clear picture of how the interactions take place. The SANS measurements presented in the following section provide a structural characterisation of the systems which enables us to clarify the nature of the associative networks formed.

Structural study by small-angle neutron scattering measurements

Fig. 8 shows the SANS patterns from 1.0, 5.0 and 10% ChEO₁₀ solutions with and without the co-surfactant C₁₂EO₃. Structural information can be extracted from the dilute solutions (1.0%). At higher concentrations, the effect of the structure factor becomes clearly dominant as seen from the appearance of a marked interaction peak. The ChEO₁₀ micelles have been described as short rods or ellipsoids^{48,49} and we were able to obtain reasonable fits with polydisperse spheres or ellipsoids (with an axial ratio of ~ 2), but the best fits were obtained with short rods of 42.5 ± 2.5 Å radius and 180 ± 20 Å (Fig. 8a). The length is in good agreement with the previously reported value of 17 nm.⁴⁸ The line of the fit does not follow the experimental plot closely in the high q region (Fig. 8a), which is in part due to instrumental smearing of the data as well as the possibility of an elliptical cross-section of the rods. These data are analysed in more detail in another publication,⁵¹ however for our purpose here, assimilating the micelles to dense cylinders is a sufficient approximation, giving

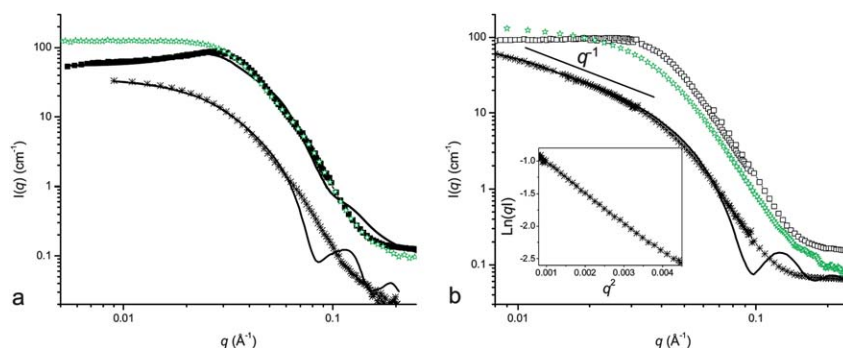


Fig. 8 (a) SANS experimental curves from ChEO₁₀ micellar solutions at 10% (■), 5.0% (★) and 1.0% (*) in D₂O. Solid lines are fits to the monodisperse rod model (for 1.0% ChEO₁₀) and to the hard-sphere model for the 10% ChEO₁₀ solutions. (b) Scattering intensity $I(q)$ from SANS experiments on wormlike micellar solutions at 10% ChEO₁₀/2.0% C₁₂EO₃ (□), 5.0% ChEO₁₀/1.0% C₁₂EO₃ (★) and 1.0% ChEO₁₀/0.20% C₁₂EO₃ (*) in D₂O. The solid line is a fit to a monodisperse rod model (1.0% ChEO₁₀/0.20% C₁₂EO₃). Inset: Kratky–Porod fit for 1.0% ChEO₁₀/0.20% C₁₂EO₃ scattering curve.

an aggregation number N_{agg} of *ca.* 800 by using the measured density of ChEO₁₀ of 1.065 g cm⁻³. Addition of C₁₂EO₃ promotes the one-dimensional growth of ChEO₁₀ micelles by intercalating between the large polyoxyethylene headgroups and thus decreasing the headgroup surface area. This leads to a clear change of the scattering pattern in the low q region, where the intensity follows a q^{-1} power law (1% ChEO₁₀/0.2% C₁₂EO₃), which is a clear signature of rods (Fig. 8b). The scattering at high q remains unchanged, showing that the cross-section is not affected by the growth. A Kratky–Porod representation of the data in $\ln(qI)$ vs. q^2 gives a linear plot over a wide q -range (Fig. 8b, inset), confirming the presence of elongated cylindrical structures, from which the radius of the cross-section and the mass per unit length can be extracted and were $R = 41 \pm 2 \text{ \AA}$ and $M_L = 3.0 \pm 0.5 \text{ kg mol}^{-1} \text{ \AA}^{-1}$. By considering a volume of cylinder of length 1 \AA and of the density of ChEO₁₀, we obtain a reasonably close value of 4 kg mol⁻¹ \AA^{-1} . Fitting the data to a monodisperse rod model confirms a cross-section of $R = 40 \pm 2.5 \text{ \AA}$ and contour length $L \approx 400 \text{ \AA}$ (the fit however is not sensitive to the value of L because it is outside the SANS measurement window, as seen by the absence of a plateau at low q). At a higher WLM concentration of 5.0% ChEO₁₀/1.0% C₁₂EO₃ (ratio unchanged), the q^{-1} behaviour is still detectable but interactions start to modify the pattern. In comparison, the scattering from the ‘short’ micelles at the same concentration (Fig. 8a) shows a flat plateau, indicative of a finite length. At higher concentrations (10%), the scattering in both systems is dominated by strong interactions. A reasonable fit is obtained for the short ChEO₁₀ micelles with a model of spheres of $42.5 \pm 2.5 \text{ \AA}$ interacting through a hard-sphere potential (Fig. 8a). Given that the rods are short (the length is about twice the cross-section diameter) and that interactions dominate the scattering pattern, it is not surprising that a simple hard-sphere model is sufficient to describe the data. At the same concentration in the mixed surfactants system (Fig. 8b) very long wormlike micelles are observed, with lengths well above 1 μm , as seen from the cryo-TEM image of 10% ChEO₁₀ with 2.0% C₁₂EO₃ (Fig. 9).

The addition of 5.0% modified dextran to the solutions of both short (ChEO₁₀) and long (ChEO₁₀/C₁₂EO₃) micelles is expected to give rise to a connected network, through cholesterol–cholesterol hydrophobic interactions between the dextran stickers and the micelles interiors. Given the different rheological

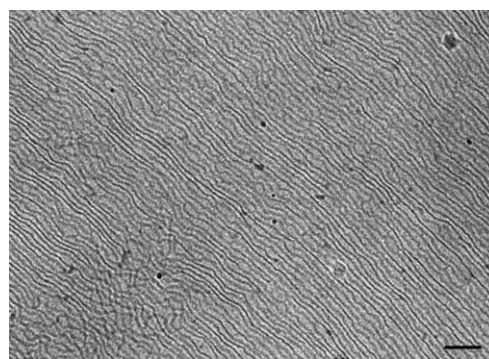


Fig. 9 Cryo-TEM image of wormlike micelles in 10% ChEO₁₀ and 2.0% C₁₂EO₃ aqueous solution.

behaviour obtained for these two systems (short and long micelles with CMD), quite different SANS patterns were expected. Quite surprisingly instead, the scattering curves from all samples at each concentration are seen to superimpose almost perfectly (Fig. 10) at each polymer/micelles ratio, suggesting a similar type of organisation. To show this more clearly, the scattering from 7.5% solutions of both short and long rods with CMD is presented in the inset of Fig. 10. The scattering of a filtered solution of polymer alone is also reproduced as a reference; it clearly shows no similarity with the patterns obtained from the mixed systems. The scattering of the polymer/micelles networks is dominated by strong interactions, with a clear structural peak. The peak position only shifts slightly with concentration, and is reminiscent of the peak observed in pure ChEO₁₀ micellar solutions at 10%, which is reproduced in Fig. 10 for comparison purposes; however, the interactions are stronger, as seen from the stronger depression of the intensity at low q . The data can be approximated to a system of interacting hard spheres (Fig. 10), with a radius of $42.5 \pm 2.5 \text{ \AA}$ and polydispersity $\sigma/\bar{R} = 0.2$, accounting for the possibility of more elongated structures (short rods) rather than spheres. Closer examination of the data shows that the structural peak shifts very slightly to lower q from 10% to 7.5% micelles, and in a more pronounced way from 7.5% to 5.0%, with a slight decrease of the intensity in the latter case. The narrow shape of the peaks suggests assimilating the scattering maxima q^* to a Bragg peak, with $d = 2\pi/q^*$, where d is an

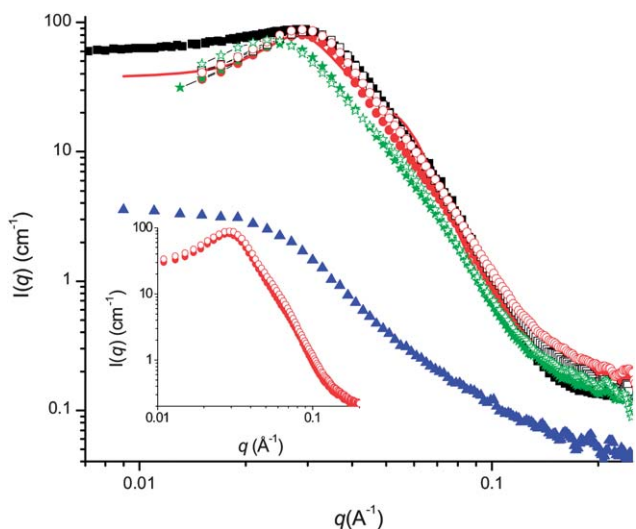
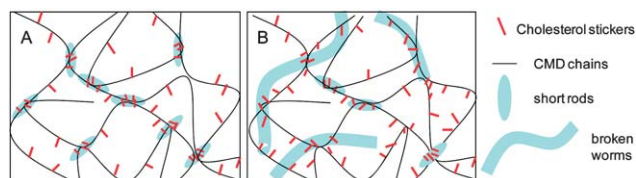


Fig. 10 SANS experimental curves from 5.0 wt% CMD (\blacktriangle), with short rod-like micelles (7.5% ChEO₁₀ (\bullet), 5.0% ChEO₁₀ (\star)) and wormlike micelle solutions (ChEO₁₀/C₁₂EO₃ at 10% ChEO₁₀ (\square), 7.5% ChEO₁₀/1.5% (\circ) and 5.0% ChEO₁₀/1.0% C₁₂EO₃ (\star)) in D₂O. For comparison 10% ChEO₁₀ micelles without CMD (\blacksquare) are also shown. The solid line is a fit to the hard-sphere model. Inset: scattering curves of 7.5% ChEO₁₀ (\bullet) and 7.5% ChEO₁₀/1.5% C₁₂EO₃ (\circ) with 5.0 wt% CMD.

estimate of the average size of the scattering density fluctuations, or network mesh size. A similar scattering pattern and interpretation have been reported for lysozyme gels.⁸¹ With this approach, we obtain mesh sizes of 273 Å, 233 Å and 217 Å for 5.0, 7.5 and 10% micelles, respectively (corresponding to $q = 0.023, 0.027$ and 0.029 \AA^{-1}), independently of the nature of the micelles. Although a very rough approximation, these mesh sizes are comparable to the ones obtained from the plateau modulus G_0 in the previous section and very close to the values calculated assuming that the micelles are simply arranged in a cubic lattice with $N_{\text{agg}} \approx 800$ (*cf.* above): 278 Å, 243 Å and 221 Å for 5.0, 7.5 and 10% micelles, respectively. It is of course expected that given the polydispersity of the micelles and polymer, heterogeneity of the cholesterol stickers distribution along the chains, the network is not perfectly uniform and regular as the value of a ‘mesh size’ would suggest. However, the picture emerging from the SANS data is overall the one of a network dominated by the scattering from a relatively ordered distribution of junctions of finite size, roughly the size of the individual ChEO₁₀ micelles. The most striking result is that the structure of the network seems to be identical in the case of the short (rods) and the long (wormlike) micelles. On the basis of this evidence, we tentatively propose that the network is dominated by the polymer chains which act as bridges between the micelles, as represented in Scheme 1, rather than the polymer cross-linking the wormlike micelles, as was found in other studies.^{41,45,69} Neutrons in this case are sensitive to the higher density regions formed by the junction zones (which we suggest are short rods) rather than the extended polymer chains, which scatter less. In this scenario, the association between CMD and the mixed-surfactant wormlike micelles leads to the breakup of the long (wormlike) micelles in order to shield more efficiently the pendent cholesterol groups from the solvent. In the case of the WLM, favourable interactions between the



Scheme 1 Schematic representation of the proposed networks formed by adding the modified dextran to solutions of (A) short ChEO₁₀ rods and (B) ChEO₁₀/C₁₂EO₃ wormlike micelles. In both cases, the elasticity of the network arises from polymer chains connected by micellar junctions through cholesterol–cholesterol interactions. In the case of ChEO₁₀/C₁₂EO₃ mixtures (B), the worms break down by interacting with the polymer, but it is expected that a distribution of lengths is still present, thus explaining the large spectrum of relaxation times in the rheological behaviour.

cholesterol groups from the ChEO₁₀ surfactant and the polymer could lead to the extraction of the co-surfactant (C₁₂EO₃) from the micelles, therefore favouring the rod-to-sphere transition. The hypothesis of a breakup of the wormlike micelles by CMD is strongly reinforced by the observation that at 2.5% WLMs, the solutions lost their solid-like behaviour when adding the polymer. Since, in the equivalent 2.5% ChEO₁₀ system, the samples remained liquid-like, this suggests that short rod-like micelles are present in both, and an insufficient number of them to provide a network connecting the CMD chains through their stickers. By assimilating the micelles to cylinders of 42.5 Å radius and 180 Å length, we would have at 2.5% concentration 2.3×10^{16} micelles per cm³ and 1.8×10^{18} cholesterol stickers. At 5.0% ChEO₁₀ and 4.6×10^{16} micelles per cm³, we have a sufficient number of junctions to form a network and at 10% (9.3×10^{16} micelles per cm³) still a large excess of CMD hydrophobes (1 micelle per ~ 20 stickers), suggesting that networks of higher strength could still be obtained at higher micellar concentrations.

Given however the different rheological response of the two systems, it is expected that some distribution of lengths still exists in the ChEO₁₀/C₁₂EO₃ micelles (Scheme 1B), accounting for the larger spectrum of relaxation times observed from the frequency sweep data (Fig. 4 and Table 1); in addition the presence of the co-surfactant may also influence the junctions lifetime. It is unlikely that slightly longer rods would be detected by SANS, since the scattering is dominated by strong interactions arising from the junction points; however the presence of long wormlike micelles would be expected to change the patterns to some extent. Therefore, we cannot exclude the presence of longer micelles in the ChEO₁₀/C₁₂EO₃/CMD systems, however the combination of the SANS data with the rheology experiments, which suggested a micellar breakup at 2.5% WLM (Fig. 4), as well as the occurrence of structural rearrangements with increasing ChEO₁₀/C₁₂EO₃ concentration (Fig. 5), all concur to give the picture that we propose in Scheme 1. We note that small structural differences that are undetected by SANS (or cryo-TEM) can lead to a very different rheological behaviour, as we have previously reported in a system of wormlike micelles with hydrophobic solutes.⁴⁰ Very subtle changes in self-assembly structures can lead to large changes in the rheological response by adding or removing relaxation processes. This is a point that we feel would deserve more discussion in the literature. This highlights again the necessity of using both techniques in combination to achieve

1 a better understanding of the systems: probing bulk *and* dynamic
aspects (rheology) as well nanoscopic organization (scattering).

Overall, the dynamics of the system are therefore dominated
by the association/dissociation processes of the polymer/micelles
5 junctions. By anchoring into the micellar cores, and possibly
partially wrapping around the micelles, the polymer creates
strong repulsions between the micelles, evidenced by the struc-
tural peak in the scattering pattern. The synergy between poly-
mer and micelles is such that micelles are able to uncoil the
10 polymer chains, which were collapsed and aggregated in solu-
tion, and act as connecting points between the long polymeric
chains, building-up a strong mixed network.

15 Conclusion

In this work we have reported that the addition of a cholesterol-
modified dextran (CMD) to short and long micelles of
polyoxyethylene cholesteryl ether (ChEO₁₀) leads to a strong
20 synergistic behaviour, inducing the formation of a strong
network. ChEO₁₀ forms short (rod-like) micelles in aqueous
solutions, which exhibit a weak Newtonian behaviour in the
concentration range studied. The addition of 5.0 wt% CMD to
these solutions (up to 10 wt% micelles) induces a strong visco-
elastic behaviour, with a significant increase of the zero-shear
viscosity, from *ca.* 2 mPa s to 688 Pa s. The frequency sweep
curves follow a near-Maxwellian behaviour (with 3 elements),
suggesting the formation of a solid-like network of mixed poly-
mer and micelles. At the lowest concentration studied of 2.5 wt%
30 micelles however, the behaviour remains liquid-like, suggesting
that the number of cross-links is insufficient to form a network. A
remarkable superposition of the frequency-sweep curves (when
normalised by the G' and G'' cross-over points) suggests that
changes in rheology with increasing micelle/polymer ratio can be
35 attributed solely to concentration effects, namely, the addition of
connection points.

The addition of small amounts of the lipophilic co-surfactant
triethylene glycol monododecyl ether (C₁₂EO₃) to ChEO₁₀
solutions (in a 1 : 5 ratio) leads to the uni-dimensional growth of
40 the short rod-like micelles into wormlike micelles (WLMs),
which show strong viscoelastic properties. The addition of
5.0 wt% CMD to the WLM solutions leads to a strong associa-
tive behaviour, characterised by the increase of G' and G'' by one
order of magnitude in addition to higher low-shear viscosity,
45 reaching *ca.* 10 000 Pa s. However, CMD/WLM solutions exhibit
a more liquid-like behaviour than the pure WLMs, as suggested
by the decrease of the G''/G' ratio. Interestingly at the lowest
WLM concentration studied (2.5 wt%), the samples become
more liquid-like when adding the polymer, suggesting a breakup
50 of the long micellar aggregates and an insufficient number of
crosslinks to form a network, as was observed at the same
concentration with the short micelles. The frequency sweep data
depart from the Maxwellian behaviour, pointing to a wide
spectrum of relaxation times. The frequency-sweep curves do not
55 superimpose as well as for the previous system, suggesting that
structural changes are occurring with increasing micelle/polymer
ratio, which are quite likely linked to the breakup of the worms.

Overall, the rheological data show the remarkable effect of
introducing CMD into the micellar solutions, both for the short

and the long micelles, due to cholesterol-cholesterol associations
between the micellar aggregates and the polymer.

Small-angle neutron scattering (SANS) measurements reveal
a very similar network for both systems: one dominated by the
5 scattering of micellar junctions which act as cross-links between
the polymeric chains, through cholesterol-cholesterol hydro-
phobic interactions. The combination of rheology results with
SANS therefore concur to suggest that the WLMs are broken-up
into shorter rods by the polymer, therefore giving a rheological
10 response not too dissimilar from the one in the network of
shorter micelles, with however a wider spectrum of relaxation
processes attributed to a distribution of lengths and possibly the
presence of the co-surfactant.

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